



Department of Energy

Washington, DC 20585

May 21, 2004

The Honorable John T. Conway
 Chairman
 Defense Nuclear Facilities Safety Board
 625 Indiana Avenue, NW, Suite 700
 Washington, DC 20004-2901

Dear Mr. Chairman:

This letter provides the U.S. Department of Energy (DOE) report (Enclosures 1 and 2) in response to the Defense Nuclear Facilities Safety Board (DNFSB) concerns raised in the DNFSB letter dated March 24, 2004. The DNFSB notes that Bechtel National, Inc. (BNI) is attempting to build a technical basis for addressing hydrogen hazards related to non-Newtonian high-level wastes with an experimental Research and Technology (R&T) program using surrogate materials. The DNFSB's concern is that the use of preliminary data so heavily based on experimental testing with surrogate materials, which has not undergone a thorough quality review, increases the chances of introducing errors into the design that may be irreversible. The DNFSB believes that decisions to proceed with the final mixing tank system design(s) and development of operating strategies to prevent hydrogen deflagrations/explosions are premature, given the degree of uncertainty that presently exists.

The DNFSB also reviewed the report of the DOE's Office of River Protection (ORP) on the adequacy of the "black cell" design concept. The DNFSB is concerned that open items in the ORP report, like the one on BNI's material selection basis, appear to be sufficiently significant to require resolution before proceeding with certain design activities.

With respect to the Board's concern that use of preliminary data based on surrogate materials experimental testing without a thorough quality review may introduce irreversible errors into the design:

Thorough quality reviews of the experimental testing have been completed and the surrogate materials used in testing have been endorsed by independent mixing experts. The design of the hydrogen mixing and control (HMC) system is based on conservative assumptions of waste behavior used in the testing program (described further in Enclosure 1).

On April 2, 2004, BNI completed its initial technical and quality verification of the R&T program test results for mixing system designs for vessels in the Pre-treatment facility containing non-Newtonian high-level wastes. The material in this report (Enclosure 3) has undergone quality reviews by the Battelle Pacific



Northwest Laboratory (PNL) and Savannah River Technology Center (SRTC) which carried out the testing. The results of the reviews are documented in Enclosure 3 of this letter. The enclosure provides the scaled prototypic test platform data, the mixing scale-up approach and basis, sparging zone-of-influence (ZOI) development, and demonstration that the mixing systems will both keep retained gas at low levels during normal operations and following a design basis event (DBE) for the March 2004 base design, thereby preventing hydrogen deflagrations or explosions. The quality review of the Enclosure 3 test report for the Pretreatment (PT) facility confirmed that the design and operating strategies specified in Enclosure 4 remain consistent with the test data.

A Peer Review meeting with an external panel comprised of experienced safety personnel from Savannah River, Oak Ridge, and other areas was held March 31, 2004, and April 1, 2004. The panel evaluated the safety strategy of the base design and provided recommendations for potential modifications. The panel observed (in their exit briefing) that the WTP hydrogen correlations are adequately conservative.

The acceptability of using surrogate materials to bound in-situ waste behavior was established through the analysis of actual wastes, tests of simulant materials, consultant input, and comparison with data of other high-level wastes. The simulant had higher weight percent solids than the expected waste, a high yield stress value, and very conservative consistency for physical modeling. An external panel of international mixing experts met on October 30, 2003, to November 1, 2003, and concurred with the WTP simulant selection. Additional details of the acceptability of the simulant are discussed in Enclosure 1, item 4.

Design Conservatism, Margins, and Flexibility

As a result of the experimental testing with surrogate materials, several mixing systems have been identified, which will accomplish the needed mixing and hydrogen control. These include combinations of pulse jet mixers (PJMs), recirculation systems, and spargers. PJMs will be used in all pretreatment inaccessible (black cell) tanks for mixing and suspension of solids off the tank floor. Spargers and/or recirculation pumps provide the mixing for the upper elevations of the tanks. The current pretreatment operational design for the Lag Storage and Blend vessels allows for mixing via *all* methods: PJMs, recirculation pumps, and spargers.

Conservatism has been included in both the design and safety strategy. Conservative approaches taken at each stage of the HMC testing program are outlined below. The safety margin in the mixing systems results from: 1) the conservative simulant selected for rheologic properties; 2) the scaled testing approach; 3) the methodology used to develop the sparging correlation; and 4) the techniques employed to evaluate gas retention and release behavior models. In order to put the conservatism of the design into perspective, scaling of test results

to date indicates that normal operation gas hold-up in the full scale vessels will be less than 1 percent by volume. Instantaneous gas releases were not observed during testing until more than 20 percent gas by volume accumulates in the tank. Even then, in the test of 7 Pascal simulant, where this was observed, the release occurred over a 20-second period. Because these hold-ups are so low, an instantaneous uncontrolled release does not appear to be credible.

As a further step to ensure the operational safety, hydrogen monitors will be provided in the vessel vent system (tank head space or vent pipe). While these systems will most likely not be safety class systems, the ability to monitor H₂ in actual plant operations is viewed as beneficial to confirm design assumptions.

In addition, safety class hardware to control the fill level of the tanks has been added to the proposed design, and the radionuclide inventory will be controlled by technical safety requirements (TSRs). By controlling dome volume and radionuclide inventory, the estimated time to LFL can be extended beyond the current 6- to 8-hour minimum. As an operational conservatism, the Pretreatment operation limits, including rheology, for each waste tank will be defined using the samples delivered by the tank farm contractor prior to WTP processing. This additional TSR will assure that the waste slurries remain well within the rheologic limits supported by the testing program.

A current review of mixing systems redundancy in the pretreatment lag storage and blend vessels indicates that mixing may be accomplished with solely PJM operation and sparger operation in *both* normal operations and post DBE. An initiative to eliminate the recirculation pumps in these tanks depends on current testing, which is evaluating the effectiveness of the spargers to rapidly establish a ZOI and to maintain a low gas hold-up with intermittent sparging. If this is not demonstrated, then the recirculation pumps will be retained.

The current non-Newtonian fluid mixing design requirements and operating strategies are provided in Attachment 2 for the Pretreatment facility. More complete design requirements and how they are achieved will be contained in a system description document under preparation.

Current designs and planning schedules allow for system refinement without compromises to safety. The ability to modify the design before the construction is "irreversible" ties to the placement of the ceiling over the black cells, which is projected to be February 2005 for Pretreatment.

With respect to the Board's concern that the Black Cell report open items appear sufficiently significant to require resolution before proceeding with certain final design activities:

BNI has developed the closure plan for each recommendation open item identified in the Black Cell Review report and has entered them into the BNI

Recommendation and Issue Tracking System for formal tracking and closure. ORP has reviewed and concurred with the closure plan activities, deliverables, and schedule. DOE expects that completion of the closure plan activities on the identified schedule will ensure that the recommendations and open items are resolved before proceeding with associated final design activities. The closure plan is statused and updated on a weekly basis and these updates are provided to the DNFSB staff. Enclosure 2 provides the April 19, 2004, update of the closure plan.

Regarding the Board's specific example of an inadequate basis for materials selection and wear rates, DOE has conducted a preliminary review of the technical basis for corrosion allowances, and determined that the likelihood of change to the material selection is minimal, once the basis for material selection is more rigorously defined.

This preliminary review found that the primary waste constituents of concern were the chlorides, fluorides, and sulfates. These constituents, along with the possible pH values expected in the tanks during operations were considered in the review. In this preliminary assessment, BNI concluded (and ORP concurred) that there was a wide margin before the maximum concentrations for the constituents would impact the current materials selected. This margin was greater than the expected variation in waste concentration of these constituents. Therefore, the risk of change is minimal.

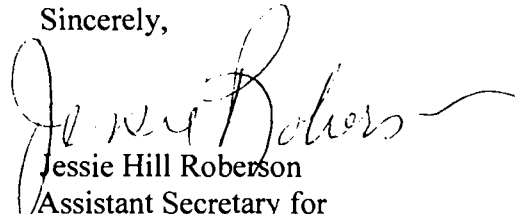
ORP will verify: (1) that waste feed hardness properties used by BNI in erosion evaluations are representative of tank farm waste information; (2) that BNI has evaluated the erosivity of the process waste streams; (3) that the corrosion evaluations have been adequately updated to account for erosion; and (4) ORP will review the need for modifications to the design required to accommodate changes in erosion allowances, if any. These actions are scheduled to be completed by July 30, 2004. Therefore, based on these reviews, both the erosion and corrosion attributes of the materials selected will be confirmed.

Based on the information provided in this letter and enclosures, ORP judges that the programmatic risk associated with continuation of the black cell vessel and piping design is acceptable.

Thank you for meeting with us on May 18. I realize that the Board has continuing concerns regarding the WTP and we have scheduled a follow-on meeting on June 2. As a result, we may need to provide refinements to this response.

If you have further questions, please call me at (202) 586-7709 or Patrice Bubar, Deputy Assistant Secretary for Integrated Safety Management and Operations Oversight, at (202) 586-5151.

Sincerely,



Jessie Hill Roberson
Assistant Secretary for
Environmental Management

4 Enclosures

cc:

M. Whitaker, DR/DOE
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SEPARATION

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**Report to Defense Nuclear Facilities Safety Board (DNFSB)
on Critical Information Requested
in DNFSB Letter dated March 24, 2004**

The following information identifies each of the DNFSB critical information requests and provides a U.S. Department of Energy (DOE) response.

DNFSB Information Request No. 1:

Identify the critical information needed to design mixing systems for non-Newtonian high-level wastes.

DOE Response:

The critical information for mixing design are the design basis rheology which establishes limits on waste receipt and revised hydrogen generation rate calculations. These are used in equipment sizing and for defining the operational modes of the mixing systems which establish normal and standby equipment sizes and controls system, and control logic requirements, and time required to generate flammable gas within these vessels. Also needed are limitations on equipment such as maximum vessel levels, which establish instrumentation and protective features, and mixing requirements that are needed to support adequate vessel heat transfer.

Other critical information needed to design mixing systems for non-Newtonian high-level wastes includes the mixing equipment specifics: the number of PJMs per tank, the PJM nozzle diameter, the PJM body diameter, the PJM nozzle velocity, PJM air demand, the number of sparge tubes, the air flow per sparge tube, the spacing and location of the sparge tubes, and the recirculation pumps used in the design. This information is developed from testing programs and analysis based on design basis rheology. The equipment information defines the support system requirements including the controls systems, instrumentation, air flow rates, line and tubing sizes, compressed air systems (including important to safety backup power and compressed air systems), anti-foaming capabilities, electrical power demands, and ventilation system capacities. The definition of these design features are discussed in the physical modeling report provided as Attachment 1 of this package. The report documents the data and scale-up methodology used to select the mixing systems for the non-Newtonian vessels in the Pretreatment (PT) facility. These designs are considered bounding in that they include the set of PJMs, recirculation pumps, and spargers. Although refinement of the operating strategies is likely, no further equipment will need to be added to these tanks to achieve the mixing and gas release requirements. These designs, also termed the March 2004 designs, are based on data developed in a testing program managed by the WTP Research and Technology (R&T) organization using resources at Battelle's Pacific Northwest Laboratory and the Savannah River Technology Center (SRTC).

Several mixing systems have been identified which will accomplish the needed mixing and hydrogen control. These include combinations of pulse jet mixers (PJMs), recirculation systems, and spargers. The current conceptual design for planning purposes is described in Attachment 4. Spargers are tubes that provide for a low volume of air to be bubbled through the liquid, thereby providing mixing via an air lift as the bubbles rise. From these considerations, designs have been defined that meet the mixing and gas removal requirements.

DNFSB Information Request No. 2:

Identify the quality and adequacy of test data being used to provide the information required by comment/issue number one.

DOE Response:

The test data developed by the two laboratories that underpin the selected designs have been reviewed by BNI in accordance with the approved Quality Assurance Program as well as undergone a quality review within the respective testing organizations. DOE considers the test data to be of high quality. A report of that test data is contained in Attachment 1. This report provides additional information on mixing system design, the quality of the test data, relevant mixing properties of the non-Newtonian high-level waste, and the use of surrogate test materials to bound actual waste behavior. The quality review of Attachment 1 for the PT facility confirmed that the design and operating strategies specified in Attachment 2 remain consistent with the test data. This report demonstrates that the tanks will be mixed and release gas generated by the wastes under normal and upset conditions.

DNFSB Information Request No. 3:

Discuss the relevant mixing properties of non-Newtonian high-level waste.

DOE Response:

The waste slurries delivered to the WTP are expected to be essentially Newtonian. The washing and leaching operations conducted within the Pretreatment Facility convert the waste to non-Newtonian slurries. Samples from waste tanks C-104, AZ-101, and AZ-102 have been pretreated in the laboratory and their flow properties measured. Based on these measurements, a design slurry rheology bases of 30 Pa Bingham yield stress and 30 cP consistency upper bound were established. The waste data, along with the behavior of the clay simulant are shown in Figure 3.2 of Attachment 1.

DNFSB Information Request No. 4:

Discuss the acceptability of using surrogate materials to bound in-situ waste behavior.

DOE Response:

The acceptability of using surrogate materials to bound in-situ waste behavior was established through the analysis of actual wastes, tests of simulant materials, and consultant input. For initial testing, Laponite, a synthetic clay, was used. Laponite is transparent and the yield stress can be adjusted via the composition. These properties were highly advantageous to evaluate mixing systems and mixing effectiveness. Laponite yield stress however is significantly reduced after mixing (shear thinning behavior) therefore a kaolinite/bentonite clay mixture was used to simulate high level waste in the actual tests which determined and demonstrated mixing system effectiveness. Clay mixtures are well known simulants for non-Newtonian materials. A clay mixture was selected for testing which had a 25-30 cP. This clay mixture had a weight percent

solids of 25-30%. The expected weight percent solids in the high level waste material is expected to be in the 15-20% weight percent range. The yield stress of 30 Pascals has been estimated to bound 70-80% of the material that will be fed to the WTP if these wastes are concentrated to 20 weight % solids in the Pretreatment facility. By measuring pretreated waste samples prior to processing, the Pretreatment operating parameters will be established for each waste tank to ensure that the waste remains less than or equal to 30 Pa. Rheology values will be included within the WTP Technical Safety Requirements document to ensure the extent of concentration within the PT facility is below the rheological design basis. It should also be noted that testing was done at broad range of yield stress levels (5 to 40 Pa clay simulants) to ensure adequate system performance. The mixed clay simulant was recommended by an international panel of mixing experts:

- Dr. Edward J. Lahoda – Westinghouse (Chair)
- Dr. Arthur W. Etchells – DuPont Consulting (retired DuPont)
- Dr. Alvin Nienow – University of Birmingham, England
- Dr. David Dickey – MixTech Consultants (retired Chemineer, Inc.)
- Dr. Richard Calabrese – University of Maryland
- Dr. Michael R. Poirier – Savannah River Technology Center

The mixing panel was convened in October, 2003 to review the PJM program and concluded that the clay simulant provides the best representation of waste slurry rheology. This simulant bounds the waste behavior for the following reasons:

- The mixed clay simulant used for the final mixing tests exhibits Bingham plastic behavior, or a constant consistency (viscosity) at increasing shear rates (better mixing). The properties of the actual pretreated wastes examined, however, are Herschel-Bulkley in nature. Herschel-Bulkley materials are shear thinning, i.e. the consistency is reduced at higher shear rates. Therefore, the clay simulant exhibits significantly higher viscous effects under representative mixing conditions than the actual waste.
- The design basis consistency for pretreated wastes is 30 cP. The consistency of the simulant should have been reduced to 5 to 8 cP to maintain similitude in the scaled models but was not lowered to provide additional conservatism. The clay simulant consistency was 20 to 30 cP. This means the viscous effects are much more pronounced in the models than at full scale. For example, jet velocities decay faster in the model than at full-scale.
- The yield stress of the simulant used for the scaled model mixing demonstrations generally exceeded the 30 Pa upper bound. Most of the PJM hybrid mixing tests were conducted with 33 to 37 Pa yield stress clay which would have created conservatively smaller mixed caverns than the 30 Pa waste will create.
- The waste density is expected to be in the 1.3 to 1.35 g/ml range at its high yield stress conditions. The clay density was 1.2 g/ml, which meant that the model PJM's were only developing ~90% of the momentum that the plant WTP design jets will deliver, again producing a conservatively smaller mixing cavern than the WTP design.

DNFSB Information Request No. 5:

Discuss the amount of excess capacity in the design to address experimental uncertainty.

DOE Response:

In order to put the conservatism of the design into perspective, scaling of test results to date indicates that normal operation gas hold-up in the full scale vessels will be less than 1% by volume. Instantaneous gas releases were not observed during testing until more than 20% gas by volume had accumulated in the tank. Even then, in the test of 7 Pascal simulant where this was observed, the release occurred over a 20 second period. Because the retained gas volume is so low, an instantaneous, uncontrolled release does not appear to be credible.

Although experimental uncertainty assessments are being performed where specific measurements are made, the PJM mixing program was planned and executed to provide greater assurance of significant design margin, using the measures described further below.

Conservative approaches taken at each stage of the PJM testing program are outlined below. The PJM safety margin results from the conservative simulant selected (see response to #4 above), the scaled testing approach, the methodology used to develop the sparging correlation, and the techniques employed to evaluate gas retention and release behavior of the models.

Approach to Scaling Test Stands

- Two of the three key non-dimensional parameters, the Strouhal number and the simulant yield stress, were preserved through the scaled designs. The third, the Reynolds number, was smaller in the model due to the high apparent slurry viscosity. This implies that the mixing behavior will be better at full scale.
- The highly turbulent mixing caverns developed by the PJM's appear to scale-up conservatively, i.e. the preliminary test data indicates larger mixing caverns at larger scale (data still undergoing quality review).

Sparging Correlation Development

- Spargers are effective in mixing the tank above the injection point due to the recirculation flow established by the air bubbles. Injection point is ~6 inches from the vessel floor. Therefore, any experimental uncertainties associated with the size of the PJM mixing cavern size does not reduce the overall mixing in the portion of the vessels mixed by spargers.
- The sparging tests were conducted in tanks that were approximately of half-WTP vessel depth. The test data indicates that the sparged volume is a weak function of submergence depth. When these tests were conducted, at half the immersed sparger depth, the sparged volume was fully developed. Sparger behavior will be scaled based on these observations, not the model-scaling factor.
- The full-scale WTP process vessels utilize only 67% of the measured sparger zone-of-influence (ZOI) diameter to conservatively apply the test data. Further, these conservative ZOI's are overlapped in the design so that the full surface of the mixing caverns are covered by a ZOI to carry the mixing to the entire tank.

- The layout of the spargers in the plant vessel designs does not credit synergism between adjacent spargers. Based on discussions with Dr. A. W. Eschels (of the expert mixing panel) the down-flow in the full-scale tanks is expected to be faster than the single sparger test data would suggest. To confirm this expectation, a multisparger test is planned to be conducted by the end of July.
- The sparger ZOI correlation was developed using data measured with 36 Pa simulant. This simulant rheology is 20% higher than the bounding 30 Pa waste. Thus, the ZOI's in the WTP will be larger than the ZOI predictions used to assure full mixing cavern coverage in the plant designs.

Gas Retention and Release Measurements

- The technique developed to rapidly generate in-situ bubbles in the simulated waste created conservatively small bubbles, which are more difficult to release. Therefore, the measured gas hold-up and release rates are lower and slower than the WTP will experience. This phenomenon occurs because the simulant gas generation rate was much more rapid than normal waste, preventing normal bubble "ripening" due to the physics of bubble formation.
- The WTP is evaluating anti-foaming agents to assure that gas release will not be impeded due to slow coalescence at the slurry surface. This will also assure that the gas bubbles will not be 'armored' by slurry particles, which could inhibit the gas release as has been observed in the Hanford waste tanks. This anti-foam technology is mature, and its specific application at WTP will be based on tests currently in progress at the Savannah River National Laboratory for WTP.
- The gas hold-up and release measurements conservatively do not include the effects of mass transfer by other means (e.g. stripping by the sparge air or from the slurry surface).
- Gas removal from both clay and a chemical waste simulant has been measured to enable estimation of the actual waste slurry gas hold-up behavior.

Work is continuing to optimize the design and it may be determined that equipment may be operated differently and/or that not all features are required. The need for applying active single failure criteria to PJMs for mixing the lower portions of the vessels during post-DBE conditions is being analyzed based on recommendations from the March 31 to April 1, 2004 external review panel. In no case however, is it envisioned that PJMs would not be a part of the mixing systems. DOE will keep the Board staff apprised of any developments in these areas.

DNFSB Information Request No. 6:

Discuss the impact of mixing designs on interfacing safety structures, systems, and components.

DOE Response:

The mixing system PJMs supplemented by spargers or recirculation pumps to achieve required mixing during normal operations and after a design basis event. These systems are designed to prevent hydrogen hazards based on conservative assumptions about hydrogen generation and release rates and recovery actions. The requirements are to maintain vessel head space flammable gas concentrations below the lower flammability limit (LFL) so there is appreciable margin against deflagration during normal operations, and not to exceed the lower flammability

limit following a DBE. These requirements impact the interfacing pulse jet ventilation, process vessel ventilation and compressed air systems. The hydrodynamic loads from pulse jet mixer operation and malfunctions are included in the vessel design criteria. A PJM System Description is in preparation that describes the functional criteria and safety design bases, how the test data is translated into the design, applicable standards, and detailed design requirements including interfaces with other facility systems, structures, and components.

Table 1 lists the primary interfacing systems and functions for the Pretreatment Facility and the related impacts and design actions associated with implementing the March 2004 base design.

DNFSB Information Request No. 7:

Discuss the ability to meet established design requirements and standards.

DOE Response:

Mixing system design requirements include achieving:

- off-bottom suspension to move solids through the system,
- blending of wastes and reagents,
- and release of evolved hydrogen gas.

As noted above, the hydrogen mitigation design requirement is to maintain vessel headspace levels at less than 25% of the LFL in the headspace for normal operation, and 100% of the LFL after a DBE.

The requirements above will be considered to have been achieved when the following conditions are met:

- no stagnant regions in the vessels
- low gas hold-up in the tanks
- predictable gas release during normal operation and following a DBE (to justify the operation cycles selected in the final design)

The results of the experimental tests, performed and anticipated, are intended to provide evidence that adequate mixing is achieved for a given configuration (Attachment 1). The tests include mixing tests and gas release tests. The March 2004 base mixing designs are derived from a successful testing configuration - number of PJMs, spargers, and recirculation pumps, their arrangement, and flow rates. This March 2004 base mixing design configuration is described in the design requirements outlined in Attachment 3. Additional design requirements will also be identified in the system description for support systems necessary for the mixing systems to accomplish their functions. These include the ability to keep spargers from blocking and capability to clear spargers and PJMs that might become blocked (even assuming continuous PJM and sparger operation), and addition of anti-foaming agents.

Table 1 - Summary of Principal Impacts of March 2004 Base Design In Pretreatment

Item	Impact/Status
Compressed air supply	Higher velocity PJMs and added spargers require more compressed air than previous design. <ul style="list-style-type: none"> • Analysis has shown that the existing plant service air compressor capacity is adequate • Higher peak demands of the redesigned PJMs will require larger air receiver vessels in PT • Supply headers in PT are not affected
Jet pump pair (JPP) design	Higher velocity PJMs require a larger capacity JPP <ul style="list-style-type: none"> • AEA has been released to develop "hybrid" JPP that boosts capacity • Design complete, confirmatory testing initiated • No impact to existing layouts
Added spargers	Air supply piping, sparger tubing, and valve racks are necessary <ul style="list-style-type: none"> • Layout studies have identified routing of piping and location of racks
Vessels	Vessel internals will change <ul style="list-style-type: none"> • Vessel sizing calculations updated to reflect more displacement by internals • Revision of vessel internals drawings for relocated PJMs , added spargers, and erosion wear plates are complete for all PT non-Newtonian vessels. • Vendors are released to start work and provide schedules and pricing for design changes
Vessel heat removal	Non-Newtonian fluid and changed mixing requires recalculation of heat-exchanger sizing <ul style="list-style-type: none"> • Complete for all PT vessels • Larger cooling jackets needed for Lag Storage and Blend vessels, added external heat exchangers necessary for ultrafiltration feed preparation (UFP) vessels • Added process cooling water demand identified and confirmed to be within existing capacity.
Erosion resistance	Higher velocity, relocated PJMs require more erosion allowance and at a different location <ul style="list-style-type: none"> • New allowance determined for PJM nozzles and impingement areas, calculation revision in progress • High velocity impingement zones identified using CFD analyses and included on revised vessel drawings
Pulse jet ventilation (PJV) system	Current capacity has been evaluated as adequate for increased PJM air flow. (Newtonian PJMs are assumed to operate no more than 25% of the time.) PJM operation frequency is still being finalized.
Process vessel vent system (PVV)	Has been redesigned to accommodate added flow from intermittent spargers. Higher potential liquid carryover being evaluated. Intermittent sparger frequency is still under development.
Added recirculation pumps	Evaluation confirmed adequate space exists and practical layout possible before decision to use pumps was made. Addition requires: <ul style="list-style-type: none"> • Revised hot cell layout and additional jumpers • New vessel nozzle connections and internal piping • Connection to power supply (adequate distribution capacity confirmed to supply 3 ~100 hp pumps) • Evaluation of adequate NPSH, preliminary calculation completed. • Additional process equipment platforms required
PJM controls	Controller logic not yet coded – no impacts; instrument and control racks under evaluation
C5 Ventilation system	In post-DBE operation, some purge may exhaust to black cells rather than be processed through PVV. Added load on C5 has been evaluated and can be accommodated.

A preliminary analysis suggests that the "idling" sparge flow rate that is specified to keep spargers from blocking may also be adequate for purging the headspace of any hydrogen (to be confirmed through the design process). Consequently, the normal vessel purge system will be disconnected from the non-Newtonian vessels and the purge function assigned to the spargers. Completion of the design process will include confirmation that measures to address indications of sparger plugging are included in the design and operating strategies.

The operation of the mixing equipment must be such that the hydrogen limits are maintained. Attachment 2 indicates the required operations. For normal operations, the mixing systems are operated continuously so that evolved hydrogen is continuously released to the headspace and purged, keeping headspace hydrogen levels well below the limits.

For post-DBE operation it is expected that intermittent operation of the mixing equipment can be shown to adequately manage hydrogen concentrations based on gas release testing performed to date. Intermittent operation will allow a much-reduced equipment requirement for ITS diesel-generators and compressors necessary to operate spargers and PJMs; ideally no more than one of the seven non-Newtonian vessels would need to be mixed at a time. As noted in Attachment 2, the mixing intervals are to be determined. Demonstration that these design requirements are satisfied post-DBE requires calculation of limiting hydrogen generation rates for each vessel (completed), calculation of the time to LFL (in progress), and demonstration by testing of the amount of time necessary to re-establish the sparging zones of influence that are needed to provided adequate mixing defined above (scheduled for May). It should be noted that in the limiting case, continuous operation of the spargers and PJMs provided by the March 2004 base design will satisfy post-DBE hydrogen management design requirements. Another potential post-DBE mixing alternative is continuous sparging with intermittent PJM operation. This is discussed in response to question 8 below.

Additional design requirements are set in the Integrated Safety Management System process and include accomplishment of the required mixing assuming a single failure of an active component, withstanding natural phenomena hazards such as earthquakes, and with required fire separation of redundant components.

The WTP Safety Requirements Document contains the standards applicable to the important-to-safety mixing design components and equipment include ASME Section VIII for vessels and ASME B31.3 for piping and tubing. These standards are implemented in design calculations, design documents, and procurement documents. Design specifications and design guides detail the analyses and features necessary to achieve compliance with the standards.

DNFSB Information Request No. 8:

Discuss any other options being pursued to supplement or replace pulse jet mixing.

DOE Response:

There are no options being pursued to replace pulse jet mixing, and no options being examined that involve other than PJMs, spargers or recirculation pumps to accomplish mixing. However, the project continues to evaluate means to simplify and improve the March, 2004 design (Attachment 2).

Two initiatives proceeded in parallel with the design work with the expectation that some of the design components could be deleted within the mixing systems, maintenance could be simplified, and improvements could be made to the control logic. These initiatives are: 1) employing an intermittent sparging of the Lag Storage and Blend process vessels to replace the recirculation pumps and 2) utilizing continuous sparging and intermittent PJM operation post-DBE. These initiatives proceeded on the basis of a Peer Review comprised of experienced safety personnel from Savannah River, Oak Ridge, and other locations, held on March 31, 2004 and April 1, 2004. This panel also observed that an intermittent mixing strategy appeared feasible.

The path forward to examine these initiatives is outlined below:

- Demonstration of intermittent sparging for normal operations requires showing that the time required for a sparger to establish the recirculation mixing pattern is much less than operational durations and that the gas retained in the vessels remains at low concentrations. The tests to determine the time to achieve steady state recirculation are in progress. The gas retention tests will be completed in June, 2004.
- Adoption of a post-DBE continuous sparging and intermittent PJM operating strategy is contingent on demonstrating that the WTP diesel generator capacity can supply the power needed for the sparger air supply, the rate that gas accumulates near the vessel floor where the spargers are least effective, and the ability of PJMs to release this retained gas when they are activated. The assessments are in progress and the tests are being scheduled.

The engineering details for implementation of these initiatives are under review.

SEPARATION

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Black Cell Review Recommendations and Open Items - CLOSURE PLAN

Report No.	RITS No.	Recommendation / Open Item	BNI Lead	ORP Lead (Notes 1 and 2)	Closure Plan	Deliverable	Closure Date	Minimum ORP Closure Verification
R-1	04-166	<p>BNI should prepare a procedure (or design guide) that completely defines the process conditions and materials selection process. This procedure should describe the process from the preparation of the material balance through the preparation of the corrosion evaluation report. BNI should reevaluate the process conditions and material selection process as part of the upcoming revision of the material balance (March 2004) to ensure:</p> <p>a. Documentation of the tracability of all chemical species from the WTP Contract requirements through the Corrosion Evaluation.</p> <p>b. Normal and bounding conditions are identified for the process chemistry.</p> <p>c. Normal and off-normal expected operational conditions are identified.</p> <p>d. A clear rationale for the material selection is documented.</p> <p>e. Wear allowance associated with both corrosion and erosion are separately identified.</p> <p>[LOI 1.1, 3.3, 3.4] [OI-10, 18, 20, 21, 22, 24]</p>	Duncan / Obenauer / Vail	Hamel [Holton, Miller, Brasel, Hamel]	<ol style="list-style-type: none"> 1. Prepare material selection design guide (2/6/04A) 2. Revise Design Guide on Preparation of Corrosion Evaluations (3/8/04A) 3. Prepare integrated mass balance calculation that includes bounding conditions (4/2/04A) 4. Issue draft corrosion evaluations using new design guides and meet with DOE to ensure that design guide meets needs (4/15/04A) 5. Update PCDSs 6. Update CEs and reconcile with vessels, piping.. procurement 7. Blue Ribbon Panel Review <p>Note: Preliminary assessment indicates no change in materials</p>	<ol style="list-style-type: none"> 1. Material Selection Design Guide 2. Updated Design Guide on Preparation of CE's 3. Integrated Mass Balance Calculation 4. Updated PCDSs 5. Updated CEs 6. Blue Ribbon Panel Report Out 7. RITS closure 	6/30/04	<ol style="list-style-type: none"> 1. Review and comment on preliminary BNI assessment of the ORP concerns and recommendations from the black cell review that concluded no immediate action was required to address those concerns other than develop the response plan contained in this table.(Complete) 2. Review and comment on Material Selection Guide and Design Guide on Preparation of Corrosion Evaluations 3. Review and comment on integrated mass balance calculation. 4. Review every corrosion evaluation prepared for black cell vessels and piping including the revised PCDSs. Confirm that the material of attached piping and components is consistent with the vessel material. This latter effort will be coordinated with the piping classification review, item R-5. 5. Review and comment on the Blue Ribbon Panel report. 6. Prepare ORP response and conclusion on closure of this item
R-2	04-169	<p>BNI should re-assess the technical basis for the erosion wear rates to determine if they are adequate and document this reassessment. BNI should determine if waste processing in WTP has the potential for increasing the erosion potential of the waste.</p> <p>[LOI 3.5] [OI-17, 19, 20, 22, 23]</p>	Duncan / Vail / Rangus	Hamel [Holton, Miller, Brasel]	<ol style="list-style-type: none"> 1. Assess adequacy of erosion wear rules <ol style="list-style-type: none"> 1a. Collect additional information from CHG regarding the basis for corrosion/erosion allowance (3/11/04A) 1b. Collect additional information from Tank Farms regarding the hardness of compounds present in solids (3/19/04A) 2. Evaluate production of erosive products in process stream 	<ol style="list-style-type: none"> 1. Respond in letter if no new information that impacts design is available incorporate into R1 products as appropriate 2. Revised calculation, if needed 3. RITS closure 	4/30/04	<ol style="list-style-type: none"> 1. ORP expects that BNI will review and revise, if needed, the calculation developing the erosion wear rules (i.e., the rules applied to ensure adequate wear allowances are specified) so that these rules are traceable to supporting documentation. ORP will review the calculation to ensure that the bases for the erosion wear rates are defensible, sufficient and complete. 2. Verify that the waste feed hardness properties used by BNI in erosion evaluations are representative of tank farm waste information provided to BNI. 3. Verify that BNI has evaluated appropriately the erosivity of the process waste streams. 4. Validate that the Corrosion Evaluations have been adequately updated to account for erosion. 5. Review the need for modifications to the design required to accommodate erosion. 6. Prepare conclusion and responses on closure of this item.

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R-3	04-170	BNI should develop design guidance on redundancy and spares and re-assess the current black cell design against that guidance to determine if additional redundancy or spares should be provided. Design features resulting in single or common mode failures of the process system should be addressed. This assessment should be documented and justified. [LOI 3.12, 3.13, 3.17]	Duncan / Hoffmann / Eichorn	Hamel [Holton, Miller, Brasel]	1. Put criteria for redundancy and spares in BOD. 2. Identify where only one flow path in process exists and communicate to DOE for further direction (4/15/04A) 3. Provide GAP analysis between design and criteria in letter 4. Address common mode failures in letter	1. BODCN 2. Letter to DOE 3. RITS closure	5/17/04	1. ORP expects that BNI will develop and apply criteria on redundancy and spares for the process and equipment design. ORP will verify that the updated guidance is clear, comprehensive and appropriate. 2. Verify that sufficient reviews have been performed to establish if additional redundancy or spares should be provided in the WTP facilities. 3. Evaluate information regarding single flow path and determine if additional redundancy or spares are desired. 4. Prepare conclusions and response for closure of this item.
R-4	04-172	BNI should evaluate the feasibility of modifying existing black cell openings such as HVAC or construction openings for future access to support unforeseen maintenance [LOI 3.8] [OI-12]	Braccia / Petrusha	Treadwell / Hamel [Treadwell]	Establish task team (complete) to prepare white paper that: 1. Identifies all existing access openings to black cells (4/20/04A) 2. Receives DOE criteria for size and location of openings (eg. three foot opening in current construction opening) (2/19/04A) 3. Evaluates feasibility of modifying them 4. Provides cost / schedule impact (ROM)	1. Report 2. Trend 3. RITS closure	5/31/04	1. Develop criteria for size and location of openings 2. Review and comment on BNI report and identify where BNI should develop Trends for modifications to provide these access openings. 3. Evaluate and provide recommendations on Trends.
R-5	04-171	BNI should establish design process "rules" for consistently and explicitly ensuring that black cell requirements are implemented that: a. Identify black cell boundaries on primary drawings and documents. b. Identify black cell requirements on physical fabrication and construction drawings, and collateral databases. c. Identify black cell requirement in procurement specifications and datasheets. d. Establish and document the requirements for black cell HVAC systems and components. [LOI 3.2, 7.2]	Duncan / Roth / Myatt / Hoffmann	Treadwell [Adams]	Establish task team of Mechanical and Plant Design (Complete) to: 1. Document Black Cell boundaries in BOD 2. Determine where there is a benefit for documenting Black Cell notation on design documents (4/15/04A) 3. Document rules in a design guide / checklist for clearly documenting black cell requirements on design deliverables and procurement documents 4. Assess pipe released for fabrication and determine schedule for documents that require update. 5. Address black cell HVAC requirements	1. BODCN 2. Design guide / checklist for black cell requirements 3. RITS closure	4/30/04	1. It is expected that once BNI has established guidelines for documenting black cell boundaries and documentation has been updated that BNI will perform a review to ensure that all components located within a black cell have been appropriately identified in the design output documents, e.g., piping lists, specifications, procurement documents, inspection requirements. 2. ORP will review and comment on the BNI guidelines for documenting black cell boundaries to ensure that they are comprehensive and will minimize the potential for failure to identify and correctly specify a black cell component. 3. ORP will review the results of the BNI review of conformance with the guidelines and confirmation that no component has been missed. 4. Prepare conclusions and response to close this item.

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OI-1	04-228	<p>BNI should continue to inform ORP on their progress in development of the AUT method. BNI should address the concerns identified in LOI 7.4.</p> <p>a. BNI intends to demonstrate that the proposed AUT method will have the capability to detect all the required weld imperfections at the detection levels specified for RT in Table 341.3.2 of B31.3 versus the less stringent criteria specified in Section 344.6.2 of B31.3. While this is not a specific B31.3 Code requirement, the Team believes this would provide increased confidence in the integrity of the welds for the 40 year plant design life. It also understood that the requirements of ASME Section V, Article 4 will be met. [LOI 7.4]</p> <p>b. BNI intends to consider the requirements added by the 2002 addendum to Section V, in Article 4, Subarticle T-421.1 for the examination of austenitic stainless steel welds and its applicability to this project. It is the Oversight Teams understanding that BNI will also evaluate complying with the non-mandatory Appendix E to Article 4, Section V, ASME for the Computerized</p>	Petrusha / Vail / Rangus	Treadwell [Adams]	<ol style="list-style-type: none"> 1. Prepare plan and schedule for program, procedures and qualification of AUT method and personnel. 2. Award AUT subcontract (3/16/04A) 3. Mobilize AUT subcontract at site 4. Confirm V&V of AUT software 5. Prepare AUT test plan 6. Perform AUT demonstration 7. Issue report on AUT demonstration 	<ol style="list-style-type: none"> 1. Qualification plan and schedule 2. Test procedures 3. Test report 4. RITS closure 	7/29/04	<ol style="list-style-type: none"> 1. Confirm that BNI has demonstrated that the proposed AUT method will have the capacity to detect the required weld imperfections that could lead to through wall leaks and meet the detection levels specified for RT in Table 341.3.2 of ASME B31.3 versus the less stringent criteria specified in Section 344.6.2 of ASME B31.3. Review the BNI procedures that implement AUT and observe the demonstration of the ability of AUT to consistently and reproducibly detect flaws that are rejectable using RT. 2. Verify that BNI is complying with the 2002 Addendum to ASME Code, Section V, Article 4, Sub-article T-421.1 for the examination of austenitic stainless steel welds and its applicability to WTP by reviewing the procedures(s) which implement AUT. 3. Evaluate "compliance" with the non-mandatory Appendix E to Article 4, Section V, ASME Code for the Computerized Imaging Techniques by confirming completion of AUT V&V. Confirm BNI has invoked the requirements of NQA-1 for software V&V. If BNI does not fully satisfy NQA-1 for software V&V, then confirm that an ABAR is being processed.
OI-1 Cont.	04-228 Cont.							<ol style="list-style-type: none"> 4. Review the BNI training plans to train, examine and certify field welding and quality control engineers for performance of AUT examinations and verify that the NQA-1 requirements for independence are being met. 5. Confirm that the AUT demonstration can detect the welding imperfections listed in Table A-110 of Section V of the ASME Code, e.g., burn through, cracks, excessive/ inadequate reinforcement, slag/tungsten inclusions, incomplete fusion, incomplete penetration, misalignment, porosity, root concavity and undercut. 6. BNI's stated intention is to examine 100% of the Black Cell pressure containing welds. Confirm that there is a plan for engineering review of any pressure containing welds that are not volumetrically examined and what controls are invoked approving such a decision.

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OI-2	04-174	BNI should document the operating conditions (and limitations) that were identified in the materials selection process for a vessel/component in the System Description for the respective system. The System Description should be placed under configuration control. [LOI 3.16]	Pisarcik	Hamel [Holton, Miller, Brasel]	<ol style="list-style-type: none"> 1. Revise System Description (SD) procedure to add operating conditions and limitations derived from material selection process. 2. Describe adequacy of existing SD change control. 3. Determine when SD should be placed under CM. 	<ol style="list-style-type: none"> 1. Revised SD procedure 2. RITS closure 	5/6/04	<ol style="list-style-type: none"> 1. Confirm that the modification of the System Description procedure captures the operational and other limitations identified in the Corrosion Evaluations. 2. Confirm that the operations organization uses the System Descriptions in a manner that ensures development of operating procedures will include these limitations. 3. Confirm that the System Descriptions are appropriately controlled under BNI change control. 4. Confirm that the System Descriptions have the appropriate priority in the document hierarchy to be up-to-date at testing and commissioning.
OI-3	04-229	BNI should brief ORP on the development of vacuum box leak testing and obtain ORP concurrence for the proposed usage. BNI should address the suggestions identified in LOI 7.5.	Petrusha / Vail / Rangus	Treadwell [Adams]	<ol style="list-style-type: none"> 1. Prepare plan and schedule for program, procedures and qualification of Vacuum Box method and personnel. 2. Obtain specimens with known leak rates (4/1/04A) 3. Fabricate vacuum box prototypes (3/16/04A) 4. Issue vacuum box leak test procedure 5. Prepare leak test plan 6. Perform leak test demonstration 7. Issue vacuum box test report 8. Issue ABAR and safety evaluation to PSC 9. Issue ABAR and safety evaluation to DOE 	<ol style="list-style-type: none"> 1. Qualification plan and schedule 2. Test procedure and report 3. ABAR and Safety Evaluation on Vacuum box test application on WTP 4. RITS closure 	6/15/04	<ol style="list-style-type: none"> 1. Review the ABAR and safety evaluation revising SRD Appendix C.26 and Appendix H to allow use of the vacuum box leak testing. If the information can be obtained, the ABAR should include a comparison discussion of the bases of the ASME B31.3 Code development of hydrostatic and pneumatic test pressure requirements and the bases for the Code specified vacuum box leak test sensitivity of 1E-3 atm-ml/sec. The ability to identify leaks with the vacuum box at the Code specified leak test sensitivity of 1E-3 atm-ml/sec and BNI's planned 7 psig vacuum and 20 second hold time should be compared with the ability of finding leaks at hydrostatic or pneumatic test pressures. The acceptability of utilizing the vacuum box testing when using AUT should be developed in the ABAR since ASME B31.3 allows use of partial AUT or RT in concert with hydrostatic or pneumatic testing. Determine the acceptability of the AUT combined with vacuum box leak testing based on the technical case in the ABAR, a review of the vacuum box leak test procedures and the demo test on known leak rate specimens.

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OI-4	04-196	BNI should evaluate the permissible configurations for black cell piping related to socket welds, branch connections, welded reinforcement pads and other welded attachments to piping. BNI should evaluate the required nondestructive examination for the permissible configurations. BNI should update Appendix A to shop fabrication and field piping specifications regarding inspection requirement for black cell piping as appropriate. [LOI 3.2, 7.2, 7.4]	Myatt / Vail	Treadwell [Adams]	1. Identify the permissible configurations for black cell piping 2. Determine appropriate NDE for these configurations 3. Revise Appendix A weld inspection table accordingly	1. Issue 24590-WTP-3PN-PS02-00018 2. RITS closure	4/2/2004 Closed	Review 24590-WTP-3PN-PS02-00018 and Appendix A, Weld Inspection Table, revisions addressing shop fabrication and field piping specifications for permissible socket weld, branch connections, welded reinforcement pads and other welded attachments to piping including the specified NDE for these configurations to confirm that it is consistent with the ASME B31.3 Code requirements.
OI-5	04-209	BNI should evaluate the air flow balance interaction of the Pretreatment Vessel Vent Process system with the vessel overflow system (PWD). [LOI 3.2]	R. Smith / Julyk	Shrader / Hamel [Ballweg]	Perform dynamic analysis	RITS closure	5/3/04	1. Verify that BNI has adequately addressed the issues associated with the Pretreatment Vessel Vent Process system with the vessel overflow system as identified in LOI 3.2.
OI-6	04-198	BNI should evaluate the overpressure / vacuum protection for the FRP system including system operational scenarios and equipment failure scenarios to determine the limiting design case. Describe how that was factored into establishing the vacuum design pressure for the FRP vessels, large diameter piping and other devices connected to the vessel vent header and system. [LOI 3.2]	Duncan / Hoffmann / Slater	Shrader / Hamel [Ballweg]	Check vessel design basis for absolute vacuum and review design provisions for vacuum conditions.	RITS closure	5/26/04	1. Verify that BNI has adequately addressed the issues associated with the FRP/Pretreatment Vessel Vent Process system with the vessel overflow system as identified in LOI 3.2. This appears to be the closure plan for Open Item 5.
OI-7	04-175	BNI should identify the design pressures basis for lines that may be used for unplugging fluidics components or other components than may become plugged. [LOI 3.2]	Duncan / Hoffmann	Shrader / Hamel [Ballweg]	Provide design pressure and unplugging recovery action	RITS closure	3/26/2004 Closed	1. Review the bases for the design pressures for these lines and confirm that the design pressures are consistent with the design code requirements.
OI-8	04-197	BNI should advise ORP on how independence will be maintained for weld acceptance as required by NQA-1. [LOI 7.4]	Petrusha / Esminger	Thomas [Treadwell]	Review existing QC program with ORP	RITS closure	4/2/2004 Closed	
OI-9	04-207	BNI should consider relocating the C5V volume dampers in the black cells to a place where they can be physically adjusted in the future or devising volume dampers that can be adjusted from the hot cell side. [LOI 3.2]	Duncan / Garcia	Orchard / Shrader [Ballweg]	Evaluate design change to move dampers out of black cells and make them adjustable, and provide rationale to DOE	RITS closure	5/28/04	1. ORP concern is that re-balancing of the HVAC flow will be difficult or not possible if required during plant operation due to the inaccessibility of these dampers. ORP will review and evaluate the BNI bases for the decision on the location and adjustability of the dampers in light of this concern. 2. Provide recommendation for concurrence of modification of the decision.

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OI-10	04-201	BNI should provide the engineering rationale for placing stainless steel pipe directly in contact with painted carbon steel support steel (w/o SS shims), using carbon steel bolts, carbon steel U-bolts, within black cells especially at lower levels which may be wetted by the cell wash/spray system or potentially rinsed with nitric acid solution for decontamination. [Reference requirement of 24590-WTP-GPG-ENG-005 Rev 1 item 10, page 42 of 163.] [LOI 3.2]	Myatt / Coutts / Rangus	Treadwell [Ballweg]	1. Identify coating requirements for black cell pipe support systems 2. Revise materials / coatings, if required	1. Specify black cell coating requirements in coatings table 2. Provide direction to engineers and designers 3. Revised design, if required 4. RITS closure	4/30/2004 Closed	1. Monitor resolution of these issues and any required modifications of the selected materials or coatings to ensure that the corrosion potential in these dissimilar material interfaces are adequately addressed. 2. Ensure that BNI coating requirements appropriately address these conditions in general.
OI-11	04-212	BNI should evaluate the approach for modular construction to ensure that significant deflections that will result from placing concrete and upper levels of steel are anticipated. [LOI 3.2]	Petrusha / Braccia / Myatt	Treadwell [Ballweg]	This open item is part of normal design process for future design of piping modules. 1. Develop design criteria and approach to address tolerances and deflections for piping modules in PTF. Consider later placement of concrete floors, removal of temporary supports, addition of other modules.	1. Design criteria and approach to address tolerances and deflections for PTF piping modules 2. RITS closure	5/17/2004 Closed	Confirm that all conditions that could affect the final configuration of the piping module, (e.g., later placement of concrete floors, removal of temporary supports, addition of other modules) are adequately considered in the design requirements documents.
OI-12	04-208	BNI should conduct a system engineering approach to assign functional requirements to the current Black Cell access openings (e.g. spray wand, camera viewing, shield plugs above separators). [LOI 3.8]	Braccia / Myatt	Treadwell / Hamel [Houghton, Ballweg, Barnes, Treadwell]	1. Identify current black cell access openings (see R-4) 2. Add access functional requirement in BODCN if directed by DOE	1. Trend 2. BODCN 3. RITS closure	6/15/04	-ORP will review and approve necessary changes to the BOD.
OI-13	04-200	BNI should evaluate the inclusion and formal documentation of thrust loading/fatigue in the design of the vessels. [LOI 5.2]	Duncan / Slater	Treadwell [Ballweg]	Document the process of inclusion of thrust loading/fatigue in vessel design	RITS closure	5/31/04	1. ORP expects that thrust loadings and fatigue are considerations in the vessel and piping design, required in procurement specifications and included in the documented design analyses for these components. 2. ORP will review the documentation for the black cell vessels and attached piping to confirm that these loads have been appropriately considered in the design.
OI-14	04-192	Due to the importance of items and/or materials destined for black cells, BNI should evaluate using independent testing laboratory services to verify quality requirements of materials and welding have been met. [LOI 5.5]	Myatt / Oldfather / M Watts	Thomas [Treadwell]	Review existing PMI program for piping with DOE	RITS closure	3/31/2004 Closed	ORP will review the PMI program and evaluate the adequacy for verifying the quality requirements of the materials.

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OI-15	04-193	BNI should evaluate the adequacy of the 24590-WTP-3PS-PS02-T-00003 Rev 1, Section 3.6.6, regarding pipe slope and determine if slope verification is adequately addressed for process lines requiring sloping. [LOI 3.2, 7.2]	Petrusha / Myatt	Treadwell [Adams]	Construction to revise procedure to specifically state that pipe slope will be verified.	1. Revised procedure 2. RITS closure	3/31/2004 Closed	Review revised procedure.
OI-16	04-202	BNI should identify what provisions have been incorporated into breakpot design and the design of other internals to make them sufficiently robust to assure that a forty year service life? Include an evaluation of fluid flashing, impingement or other transients have been considered in the design of these vessels and internals. [LOI 3.2]	Duncan / Hoffmann	Gilbert / Shrader [Ballweg]	1. Review breakpot suitability with flashing for 40 year life. 2. Describe code required aspect of the design of internals to support 40 year life.	RITS closure	4/30/2004 Closed	Confirm that the methods and results of the BNI evaluation support a 40 year design life for break pots and other vessel internals.
OI-17	04-194	BNI should evaluate the erosion wear allowance for vessels with PJMs accounts for the higher wear rate of stainless steel when the surface to particle angle is small, [e.g. <30 degrees] such as observed during site inspection of the perimeter PJMs in FRP-VSL-00002A/B/C/D. This should be evaluated for all other vessels with PJMs. (WSRC-TR-2001-00156, RPP-WTP Slurry Wear Evaluation Literature Search) [LOI 3.2]	Duncan / Vail	Hamel [Ballweg, Holton]	Re-iterate how this is accounted for in design.	RITS closure	3/31/2004 Closed	1. Verify that the methods and results of the BNI evaluation and design support a 40 year life for PJMs that are exposed to high velocities, with particular consideration of erosion wear in the nozzles that are exposed to particles at low impact angles. 2. Coordinate this review with that of OI-19.
OI-18	04-204	BNI should update the MSDS for FRP-VSL-0002A/B/C/D, the PT Waste Feed Receipt Vessels, 24590-PTF-N1D-FRP-00001 Rev 2, to have a minimum solids concentration consistent with the WTP contract value of 3.8 wt%. [LOI 1.1]	Duncan / Obenauer	Gilbert / Shrader / Hamel [Miller, Holton]	1. Update PCDS to align with revised contract and wear calculation (see R-1) 2. Preliminary review of information (3/8/04A)	1. Updated PCDS 2. RITS closure	5/7/04	1. As part of the closure of R-1, ORP will review all updated PCDS and Corrosion Evaluations for vessels and piping, where available, located in black cells. 2. ORP will confirm that the PCDS for PT Waste Feed Receipt Vessel has the appropriate waste concentration specified.
OI-19	04-205	BNI should present the results of the erosion testing (non-Newtonian fluid applications) for the PJM nozzle materials ORP for review and evaluation. The design impacts associated with application of the testing results should be identified. [LOI 3.3, 3.4]	Duncan / Hoffmann	Hamel [Holton, Miller, Brasel]	1. Conduct corrosion test for PJM inserts (2/26/04A) 2. Select material options (2/26/04A) 3. Conduct agitator blade erosion tests	1. Corrosion test report 2. Agitator blade erosion test letter report 3. RITS closure	5/14/04	1. Review the results of the corrosion tests and the bases for the selected materials. 2. Confirm that the results of the tests and other information support the selection of the materials for PJMs. This Open Item is just to provide test results. 3. Coordinate this review with that of OI-17.

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OI-20	04-233	BNI should assess the materials selection (corrosion evaluation-24590-PTF-N1D-UFP-0003) for the UFP feed Vessels (UFP-VSL-00002A/B) and other affected vessels based upon the proposed use of these vessels for oxidative leaching of the HLW sludge. These vessels will be used in process operations in which hard precipitates are produced (e.g. MnO ₂). This assessment should account the procurement and fabrication of the affected vessels. [LOI 3.3, 3.4]	Duncan / Vail / Obenauer	Hamel [Holton, Miller, Brasel]	Oxidative leaching is not part of the design basis nor required by contract. 1. Notify DOE of potential conservatism in erosion and material selection for hard precipitates. 2. Erosion precipitates addressed in R-2 3. Review preliminary R&T results and estimate impact on material selection of affected vessels	1. Letter to DOE 2. RITS closure	5/31/04	1. Evaluate potential conservatism in erosion and material selection for hard precipitates in UFP vessels.
OI-21	04-206	BNI should add the information presented in their response to LOI 3.4 on a corrosion assessment under acidic conditions for vessels as appropriate to the respective MSDS's and Corrosion Evaluations. [LOI 3.3, 3.4]	Duncan / Vail / Obenauer	Hamel [Holton, Miller, Brasel]	Update PCDS's and CE's for off-normal conditions (see R-1).	1. Updated PCDS's and CE's 2. RITS closure	5/17/04	1. As part of the closure of R-2, ORP will review all updated PCDS and Corrosion Evaluations for vessels and piping, where available, located in black cells. 2. ORP will confirm that acidic conditions have been appropriately considered in the updated PCDS and CEs. This review will identify and focus on vessels exposed to both acidic and alkaline conditions as part of normal and off-normal operations.
OI-22	04-199	BNI should reconcile the discrepancy between the SRD based requirements for corrosion/erosion wear allowance with the erosion "rules" contained in the wear allowance calculation (24590-WTP-M06-50-00004, Rev B). [LOI 3.5]	Duncan / Vail	Hamel / Miller [Miller]	Submit to PMT an ABAR to align SRD with wear allowance calculation	1. ABAR 2. RITS closure	4/15/2004 Closed	1. Resolution of Item R-2, above, may result in a change to the wear allowance calculation. ORP will ensure that BNI resolution of R-2 and the OI result in consistency in the requirements of the SRD and the design basis calculations.
OI-23	04-191	BNI should document the "rules" established for the erosion allowance (24590-WTP-M06-50-00004, Rev B) in a formal design guide. [LOI 3.5]	Duncan / Hoffmann	Hamel [Holton, Miller, Brasel]	Revise existing design guides on pipe sizing for erosion rules on maximum velocities	1. Revised design guides for erosion rules 2. RITS closure	3/24/2004 Closed	1. ORP expects that the revision of the existing design guides will be an outcome of the BNI resolution of item R-2, above. ORP will confirm that the updated wear "rules" are incorporated into the appropriate design guides.
OI-24	04-195	BNI should ensure that Black Cell Vessels have nozzle internals so that fluids entering the tank extend inside the vessel to assure that chemical additions drop freely into the vessel rather than dribble along the vessel wall. This reduces the risk that concentrated chemicals added to the vessel will attack the vessel locally. [LOI 3.2]	Duncan / Slater	Hamel [Ballweg]	1. Review all black cell vessels to assure this good design is implemented, as practicable. 2. Revise design and procurement documents accordingly for awarded procurements. All others as scheduled	1. Revised design and procurement documents, as required 2. RITS closure	3/31/2004 Closed	1. Review the results of the BNI assess of the status of vessel chemical addition penetration design to ensure that the nozzle internals have been specified to have sufficient penetration through the wall to prevent flow down the interior of the vessel wall. 2. ORP expects that this is a design feature that should apply to all vessels; not just those in the black cells. ORP will confirm that this is part of the general specification for WTP vessels, as applicable.

Black Cell Review Recommendations and Open Items - CLOSURE PLAN

Report No.	RITS No.	Recommendation / Open Item	BNI Lead	ORP Lead (Notes 1 and 2)	Closure Plan	Deliverable	Closure Date	Minimum ORP Closure Verification
OI-25	04-210	BNI to advise what provisions are being made to preclude seismic interaction of three vessel applications (TCP-1, TLP-9A&9B and CNP-4) were identified as SC-III items with SC-I items in the black cells. [LOI 3.2]	R. Smith / Julyk / Lowry	Treadwell [Ballweg]	Conduct a meeting w/ participants to get concurrence on path forward. 1. Conduct ISM / Hazard Topography meeting to evaluate the 3 Cat III vessels for appropriate seismic categorization and impact to Cat I designed equipment and systems in the black cell, as necessary. (4/8/04A) 2. Revise design and procurement documents, if required	1. ISM / Hazard Topography meeting minutes and action items, as necessary 2. RITS closure	4/22/2004 Closed	1. Review the results of the BNI assessment on the seismic classification of the TCP-1, TLP-A&B and CNP-4 vessels to ensure the seismic classification is appropriate.
OI-26	04-203	BNI should evaluate the method of installing, tightening and securing pipe support bolts, u-bolts and other mechanical fasteners to ensure they will remain secure for the entire service life of the facility. [LOI 3.2]	Myatt	Treadwell [Ballweg]	1. Review Black Cell connection requirements for piping systems 2. Provide written direction to engineers and designers, if required 3. Revise design, if required	1. Written direction, if required 2. Revised design, if required 3. RITS closure	4/30/2004 Closed	Review revised design guidance.
Executive Summary		It is suggested that BNI prepare a "Black Cell Management Strategy" that clearly delineates the Design, Operations, and Maintenance Standards for the WTP Black Cells	Duncan / Roth	Hamel [Hamel, Naft]	Add Black Cell requirements and criteria as a new section to the Basis of Design (BOD) rather than a separate stand-alone document	1. BODCN 2. RITS closure	5/17/04	Review BODCN.
		Note 1: The original members of the Black Cell Review Team will be consulted for verification of BNI closure actions depending on availability. Note 2: Names in brackets denote originators of recommendation/open item						
DOE-1	NA	ORP should evaluate the benefits of providing erosion hardening design features (e.g. PJM nozzle hardening and PJM jet wear plates) to the black cell vessels that are expected to be used for the storage of Newtonian fluids to provide future flexibility in waste processing. These tanks include the LAW and HLW receipt vessels. [LOI 3.3, 3.4]	NA	Hamel [Holton, Miller, Brasel]	Perform evaluation	Documented evaluation	4/30/04	NA

Black Cell Review Recommendations and Open Items - CLOSURE PLAN

Report No.	RITS No.	Recommendation / Open Item	BNI Lead	ORP Lead (Notes 1 and 2)	Closure Plan	Deliverable	Closure Date	Minimum ORP Closure Verification
DOE-2	NA	ORP should evaluate increasing the utilization of Government Acceptance Inspection on field welding performed on black cell piping. [LOI 7.3]	NA	Thomas [Treadwell]	Perform evaluation	Documented evaluation	8/1/04	NA
DOE-3	NA	ORP should add to their design oversight schedule an assessment review of BNI's vacuum testing program and AUT implementation. [LOI 7.2, 7.4]	NA	Treadwell [Hamel, Treadwell]	1. Develop draft AUT/Vacuum Test oversight plan (3/3/04A) 2. BNI review comments on plan and schedule input. 3. On-site test observation. 4. Prepare report.	Technical Report.	8/1/04	The important NDE issues for black cells were identified in detail in the W/TP Black Cell Design Adequacy Oversight Report. Elsewhere within this closure plan, DOE has identified the minimum NDE review attributes that are necessary and will be verified for closing the identified items including the AUT and Vacuum Box development plans, procedures and demonstration test.
DOE-4	NA	ORP should perform a follow-up design oversight that focuses on the resolution of any remaining recommendations and open items from this design oversight, design and construction processes for vessels and components in the WTP Black Cells, and the modular piping sections.	NA	Hamel [Hamel]	1. Add follow-up design oversight to ORP design oversight schedule (5/31/04) 2. Perform follow-up assessment (TBD)	Documented evaluation	TBD	NA

SEPARATION

PAGE

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This document will now be processed through the Hanford Waste Treatment and Immobilization Plant review procedure to produce an interim project report. The full test program will be documented in reports from our sub-contractors, which will be released this summer.

Steve Barnes
PJM Program Manager

SMB/jmp

Attachment:

“Hybrid Mixing System Test Data Supporting the Ultrafiltration Feed Process (UFP-VSL-00002A/2B) and HLW Lag Storage (HLP-VSL-00027A/B) and HLW Blend (HLP-VSL-00028) Vessel Design Efforts”

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Grabowski, S. w/a	MS4-A2	Wilson, J. w/a	MS12-B
Hoffmann, M. w/a	MS4-D2	Winkler, C. w/a	MS5-G

**Hybrid Mixing System Test Data
Supporting the Ultrafiltration Feed
Process (UFP-VSL-00002A/2B), HLW
Lag Storage (HLP-VSL-00027A/B)
and HLW Blend (HLP-VSL-00028)
Vessel Configurations**

PJM Task Team

March 2004

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Acronyms and Abbreviations

acfm	actual cubic feet per minute
APEL	Applied Process Engineering Laboratory
BHRG	BHR Group
BV	blend vessel
cfm	cubic feet per minute
CCN	correspondence control number
CFD	computational fluid dynamics
DACS	data acquisition and control software
DBT	design basis event
FEP	Feed Evaporation Process System
FMP	Fluid mixing Processes
FRP	Feed Evaporation Process System
gpm	gallons per minute
GR&R	gas retention and release
H/D	ratio of slurry height to vessel diameter
HLP	HLW Lag Storage and Feed Blending Process System
HLW	high-level waste
ID	inner diameter
ITS	important to safety
JPP	jet pump pair
LAW	low-activity waste
LRB	laboratory record book
LS	lag storage
OD	outer diameter
PJM	pulse jet mixer
PVC	polyvinyl chloride
PWD	Plant Wash and Disposal System
R&D	research and development
R&T	research and technology
RD	rheological additive
RDP	Spent Resin Collection and Dewatering Process System
RFD	reverse flow diverter
ROB	region of bubbles
scfm	standard cubic feet per minute
TEP	Technetium Eluant Recovery Process
TXP	Technetium Ion Exchange
UHP	ultrafiltration feed process
VFD	variable frequency drive
WTP	Waste Treatment Plant
ZOI	zone of influence

1.0 Introduction

1.1 Background

The Pulse Jet Mixer (PJM) Task Team (R&T, Engineering, R&D, and mixing consultants) developed an integrated strategy for scaled testing to validate PJM mixing in Waste Treatment Plant (WTP) vessels containing non-Newtonian fluids in June 2003. The scaled PJM mixing tests were to provide information on the operating parameters critical for the uniform movement (total mobilization) of these non-Newtonian slurries. In addition, the WTP project funded work to determine WTP-specific hydrogen generation rate source terms and gas transport characteristics in representative scaled prototypic mixing configurations during PJM operation. The gas transport testing included gas retention and release (GR&R) characteristics within non-Newtonian slurries during mixing operations to support design of the PJM mixing systems, to understand these characteristics within the selected mixing system, and to allow for development of normal operation and post-design basis event (DBE) mixing strategies.

Initial (physical) scaled testing confirmed in October 2003 that the baseline pulse jet designs in these vessels did not mix the non-Newtonian slurries to the extent necessary to meet WTP design requirements. Phase I of the PJM program developed an alternative "PJM-only" configuration that mixed the vessels containing non-Newtonian slurries in accordance with WTP design requirements toward the end of November 2003. Phase I scaled gas retention and release testing demonstrated that the WTP could provide safe gas control with these configurations in December 2003. In the same time frame, the hydrogen generation rate source testing was completed using actual waste samples from "expected worst case" tanks, and a better correlation to predict hydrogen generation for use by the WTP Project was developed. While the alternative PJM configuration was acceptable, implementation of the PJM-only mixing systems severely impacted the WTP facility designs due to increased numbers of PJMs, additional piping, and the significantly increased air consumption necessary to operate these systems.

To minimize the impact to the overall project cost and schedule, the PJM Task Team was directed to develop PJM hybrid mixing systems to reduce the WTP impact. Phase II of the PJM program investigated further alternative configurations to assess the effects of slurry rheology changes, reduced tank volume, PJM jet velocity and nozzle size, sparging, and recirculation pump operation. Phase II PJM hybrid mixing systems recently completed additional testing to confirm that the modified configurations mix non-Newtonian slurries in accordance with WTP design requirements. PJM hybrid mixing systems GR&R testing confirmed that the selected PJM configuration provides safe gas control in accordance with WTP design requirements.

This document describes the PJM hybrid mixing systems goals, mixing operation modes, test stands and experimental methods, selected configurations, and testing data supporting the ultrafiltration feed process (UFP) (UFP-VSL-00002A/2B), HLW lag storage (HLP-VSL-00027A/B) and HLW blend (HLP-VSL-00028) vessel configurations selected by the Pretreatment Facility Team and Central Engineering.

1.2 Design Goals for the Phase II PJM Hybrid Mixing Systems

In conjunction with Engineering, Pretreatment and HLW Vitrification Facilities, and Project Management personnel, the PJM Task Team conducted the hybrid mixing systems testing program with the following success criteria and constraints:

- Achieve complete mixing (i.e., no stagnant regions) with turbulent conditions in the majority of the slurry volume. Turbulent mixing conditions enhance heat transfer within the vessel. Turbulent mixing facilitates the suspension of waste particles.
- Use the baseline PJMs to mix in the bottom of the vessels; supplemental mixing would be used to mix the upper portion of the vessels.
- Limit the PJMs to the original baseline design, which includes a 4-inch exit nozzle; however, the standard jet pump pairs (JPPs), which use compressed air as the motive force to drive the pulse tubes, will be upgraded to allow for a nozzle velocity of 12 m/s. The number of PJMs is limited to six in the ultrafiltration feed process (UFP) vessel and eight in the lag storage (LS).
- Target the non-Newtonian slurry rheology in terms of yield stress from 5 to 30 Pa based on actual pretreated waste data from Tanks C-104 and AZ-102, respectively.
- Do not use the UFP recirculation pump above a vessel slurry height (H) over vessel diameter (D) ratio of 1.4.
- Mix the LS and blend vessels to the required H/D ratio of 0.74, with pretreated waste slurries having a yield stress of 30 Pa.
- Operate the air sparge systems within the constraints of the Pretreatment Facility vessel vent system; i.e., total additional air flow of 1520 scfm.
- Consider supplemental mixing technologies that are technically mature above emerging technologies. Eventually, PJMs, sparging, and steady jets (flow provided by recirculation pumps) were selected as the only options.
- Limit the full-scale recirculation pumps to 2200 gpm for each vessel.
- Provide robust mixing systems, i.e., provide for a 40-year operation life in a configuration that can be fabricated readily.
- Keep gas holdup (how much gas is retained at steady state in the mixed waste during normal, continuous PJM operation) as low as possible.
- Control gas release rate (how quickly gas is released upon PJM restart after a period of no mixing) after a post DBE or non-mixing period.
- Minimize air consumption requirements on both the supply and vessel ventilation systems.
- Minimize the number of vessel penetrations.
- Minimize the overall risk to the project.
- Minimize the overall cost and schedule impact to the project.

1.3 Operational Scenarios

1.3.1 Operational Processing Modes

Waste slurries with a sodium concentration of approximately 5 M is delivered to UFP-VSL-00002A/B for separation into solid (HLW) and liquid (LAW) fractions. The waste in the feed vessel is pumped through three bundles of cross-flow filters. The water and other soluble components of the waste permeate pass through the filter media and discharge into one of the permeate receipt vessels. The solids are recirculated into the feed vessels, where additional waste is received from the feed preparation vessels to replace the permeate and maintain a relatively constant volume (corresponding to an H/D of 1.4). While the solids are being concentrated, the filters will be back-pulsed periodically. Back-pulsing pushes permeate back through the filters into the concentrated slurry and dislodges solids that have built up on the filter surface, thus enhancing the overall permeate flux rate. The UFP vessels are equipped with PJMs, cooling jackets, high-pressure steam injectors, and chemical reagent feed lines. The cooling jackets are used to control the slurry temperature while filtering and to cool the waste after leaching. The filter pumps are large and add a significant amount of energy to the waste as heat. The high-pressure steam is used to heat and hold the waste at an elevated temperature during the leach process. The chemical reagents are used for leaching and filter cleaning.

Solids treatment begins after the solids are concentrated to approximately 20 wt% (dry basis) for Envelopes A, B, and D and 15 wt% for Envelope C. The first solids treatment step is to wash the solids with process condensate, using the same steps as solids filtering or concentration to remove soluble components. Process condensate is added to UFP-VSL-00002A/B to replace permeate that passes through the filters. After the Envelope A, B, and D solids are washed, they are leached (Envelope C solids are not leached) if warranted (corresponding to an H/D of 1.8). The first step in leaching is to add 19 molar sodium hydroxide until a calculated value of 3 molar free hydroxide is reached for the batch. The solution is then heated with high-pressure steam to 176°–194°F and allowed to digest for eight hours. After digesting, the slurry is cooled, then filtered until the solids concentration is increased back up to 20%. After the solids are reconcentrated they are washed again with process condensate to remove the residual sodium hydroxide and dissolved solids. The treated solids are then discharged to LS (HLP), and chemical cleaning of the filters, if required, begins.

Normally the LS vessels (HLP-VSL-00027A/B) receive treated solids from ultrafiltration; however, treated solids can be sent directly to the blend vessel (HLP-VSL-00028) if necessary. Backup blend vessel HLP-VSL-00027B can receive the same waste transfers as HLP-VSL-00028. Treated HLW solids, concentrated Cs, and Sr/TRU solids (if available) are blended together in HLP-VSL-00028, sampled, and routed to HLW vitrification.

1.3.2 Mixing Operation Modes

This section discusses the normal and post-DBE mixing operation modes. Normal mixing is that required for routine or normal plant operation. Post-DBE mixing refers to the mixing modes that will be available after a plant upset.

Normal Mixing: The hybrid mixing system will provide for complete mixing of the non-Newtonian slurry within the constraints of the Pretreatment Facility vessel ventilation system. Gas holdup in the waste slurry during normal operation will be low, i.e., gas release will be effective. Normal operations mixing can use a combination of PJMs, sparging, and recycle pump systems.

Post-DBE Mixing: After a DBE, sparging air can be diverted to the cell ventilation system and the incremental air flow constraint of 1520 scfm is lifted. Normal mixing uses equipment that is not rated for important-to-safety (ITS) use. Post DBE, the hybrid mixing system will use ITS-rated systems only and provide for complete mixing of the non-Newtonian slurry to ensure gas release from the slurry. Post-DBE mixing and gas release is limited to PJM and sparger operation.

1.4 Overview of the PJM-Hybrid Mixing Systems Design Approach

The hybrid mixing systems considered in this work involve the combined use of PJMs, steady mixing jets created by recirculation pumps, and air sparging. The mixing technologies were combined to take advantage of their respective strengths.

PJMs are used for mixing the lower region of the vessel contents and facilitating off-bottom suspension of solids. PJMs are ideally suited for these tasks because they discharge downward with nozzles near the vessel floor. The ideal PJM configuration for hybrid systems is one that creates a well-defined, highly turbulent cavern. The material in the upper region of the vessel is then transported to the turbulent cavern by the other systems where it is mixed (spargers and/or steady jets) as illustrated in Figure 1.1.

Having a high degree of turbulence is important to encourage both adequate mixing and gas removal, as well as to minimize scaling issues for prototypic test results that will be applied at full scale (scale-up is discussed in more detail in Appendix A). Additionally, having an obstruction-free interface between the mixed and unmixed regions simplifies the specification of spargers and jet nozzles.

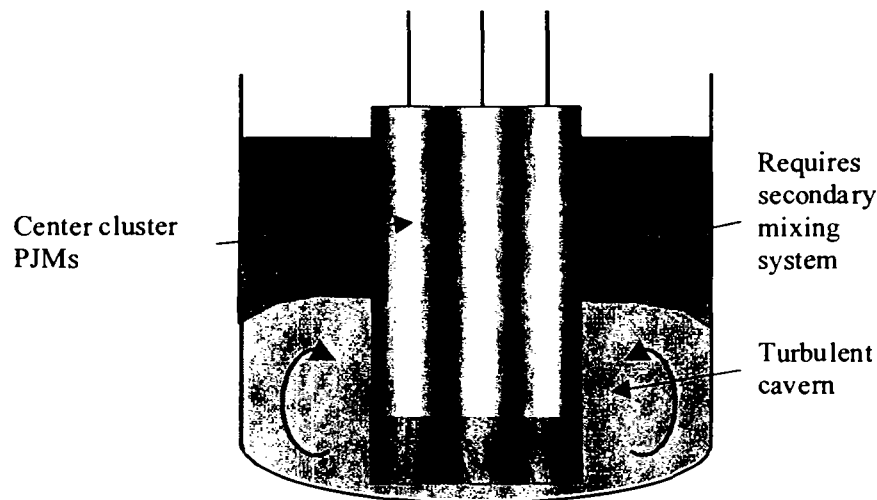


Figure 1.1. PJM-Hybrid Mixing Approach. Central cluster PJMs mix the lower region of the vessel and secondary systems mix the upper region.

A centralized cluster of PJMs with nozzles angled toward the tank wall was found to be the most effective at creating a distinct mixing cavern. Tests with a distributed array of PJMs were also conducted and found to provide good overall mixing (determined by the dye method); however, the uniformity of the cavern was found to be highly sensitive to nozzle impingement angle, and the quality of the turbulence was suspect.

Steady turbulent jets from recirculation pumps are known to be effective in mobilization and mixing applications. In general, mixing effectiveness is improved by increasing either the nozzle diameter or jet velocity. If the flow rate is fixed, the mixing performance is improved only by increasing the nozzle velocity, which implies a subsequent reduction in nozzle diameter.

Mixing performance can also be improved by increasing the number of mixing jets. Jets are a source of linear momentum and tend to be highly directional with relatively small spread angles (about 15 degrees for a free Newtonian jet). Once they impinge on solid surfaces, they tend to follow the contour of that surface. Further, cavern formation (or similar channeling) can occur for non-Newtonian slurries. Single jets can be used to mix entire vessels if the flow rates are high enough; however, a single jet will often break through the fluid surface and dissipate its energy before complete mobilization, particularly in a non-Newtonian slurry. Hence, by distributing the total available flow through multiple jets, more regions of the vessel can be affected and overall mixing can improve.

Ideally, the jet nozzles are located just below the PJM cavern interface, angled upward and aimed between the PJMs and the vessel wall. Material from the lower mixing zone is entrained and mixed into the upper region, a configuration well suited for operation at reduced operating volumes.

Air sparge tubes provide mixing an alternative mechanism. Rising air bubbles produce drag on surrounding fluid, creating an upward pumping effect. Once at the surface, fluid must recirculate downward. The net result is an upward bubble zone of mixing (in this document, this region is referred to as the region of bubbles [ROB]), surrounded by a larger, downward zone (in this document, this region is referred to as the zone of influence [ZOI]). Sparge ZOIs will interact in potentially beneficial ways if neighboring sparge points are spaced close enough. However, these interactions for non-Newtonian fluids are not fully understood and are not addressed in this document. Locating the outlet of the sparge tube near the bottom of the tank and well inside the PJM cavern should provide the capability to completely mix the tank contents.

1.5 Overview of the Scaled Testing Methodology

The scaled testing methodology involved conducting tests in a number of scaled vessels with representative non-Newtonian simulants. Five test stands were tested with PJMs; three were used to investigate the scaling laws and two were scaled versions of the full-scale tanks. Information on sparging was obtained with a single large-scale sparge tube. Scale-up and application of the mixing technologies are based on a mix of well-known theory and developments by the PJM mixing program.

As described in Section 3.1, the two primary simulants were Laponite and a mixture of kaolin/bentonite clay. Laponite is a thixotropic colloidal synthetic clay that forms a transparent gel when left unshered. This simulant was used for assessing the scale-up behavior of the PJMs and visualizing the flow behavior

in the scaled prototypes. The kaolin/bentonite clay mixture exhibits a Bingham plastic rheology that closely represents the rheology of actual waste slurries. This simulant was used to investigate the scale-up behavior of PJMs and GR&R characteristics. It was also used to assess the performance of the scaled prototypes.

The scale-up of the PJM mixing performance and the GR&R characteristics was investigated at three different scales with geometrically scaled test stands containing four PJMs. The largest test stand (described in Section 2.2) is the 12,000 gallon vessel in the Hanford 336 building which is similar in size to the actual concentrate receipt vessel. The intermediate sized test stand (described in Section 2.1) is located in the Applied Process Engineering Laboratory (APEL). It is approximately one-quarter scale (based on linear dimensions) relative to the large tank with a total volume of about 250 gallons. The small-scale tank, which is about one-half scale relative to the APEL test stand, is located at the Savannah River Technology Site.

The basis for scale-up of the mixing induced by PJMs and steady jets is based on modifications to turbulent jet theory to account for the non-Newtonian rheology and non-steady jets from the PJMs. Dimensional analysis (appendix A) was used to identify the important dimensionless parameters and guide the experimental design. The configuration for the sparging systems was based on the results of nearly full-scale tests with a single sparge tube (refer to Sections 2.4 and 3.3).

Scaled prototypes were used to evaluate the various mixing configurations. Both the LS and UFP vessels (both described in Section 2.3) had scale factors in the range of 4 to 5. Approximately 150 separate runs were conducted with these units containing various configurations of PJMs, recirculation pumps, and spargers. Only the mixing results from runs that have a direct bearing on the final configuration are reported in Section 3.2. GR&R results for the scaled prototypes are reported in Section 5. Development of the basis for scale-up of the GR&R results is ongoing and will be included in a future document.

2.0 Test Stands and Experimental Methods

This section contains a description of the test stands and the experimental methods. The APEL 4 PJM and the large-scale PJM test stands are described first and were used to demonstrate the scaling laws for mixing, gas release, and gas holdup. The scaled prototypes are described next and are geometrically scaled models of the full-scale UFP and LS tanks. The prototypes were used to evaluate various mixing configurations. Section 2.4 describes the equipment and methods used to obtain performance data for sparging in non-Newtonian slurries. Section 2.5 describes the methods used to assess GR&R behavior in the simulants. The final section describes the methods used to assess the extent of mixing.

The dimensional information presented in this section is divided into three categories based on (1) standard sizes; (2) measurements made prior to or during, or recreated after the testing; and (3) target values or ranges.

The first category pertains to the internal or external diameters of the stainless steel or PVC tubing/ piping materials used in construction of the pulse tubes, nozzles, recirculation lines, and sparger lines. Although the actual diameters vary from manufacturer to manufacturer, these values are generally within $\pm 5\%$. In the text, tables, and figures, unless otherwise noted, for the diameters of pulse tubes, nozzles, recirculation lines, and sparger lines, only the nominal values are listed.

The second category dimensional measurements are those that can be quantified and are presented as such in the discussion with the appropriate uncertainties.

The third category mainly corresponds to dimensional information that was impossible to measure directly, such as elevations of the vessel internals relative to the vessel bottom (e.g., distance of nozzles from bottom). Although significant effort was made to achieve the target values specified in the testing sequences, no direct as-built measurements were made because of space limitations within the tank (i.e., manned entry was not possible). In addition, this category also includes those measurements that were not recorded at the time of the testing and could not be recreated. The third category of measurements is indicated as approximate in the text, tables, and figures and should only be treated as such.

2.1 APEL 4 PJM Test Stand

The APEL 4PJM test stand (Figure 2.1) is a linearly scaled version of the 4PJM test setup in the 336 test facility. The configuration details, subject to the constraints presented at the beginning of Section 2, are discussed below. The diameter of the tank is 33.8 ± 0.5 inches, which corresponds to a scale factor of ~ 4.57 . This test stand consists of four PJMs constructed of 5-inch (5.29-inch ID) schedule 10 stainless steel pipe tapered to an approximately 60° angle cone truncated to a custom-built nozzle with a 0.88 ± 0.01 -inch ID. The length of the cylindrical section of the PJMs was 48 ± 1 inches. The height was intentionally set longer than the PJMs in the 336 test facility to enable testing at higher H/D ratios (up to H/D of 1.6 and a volume of approximately 180 gallons) than were possible in the large-scale test stands. The PJMs are situated around the center of the tank in a square along a pitch-circle diameter (PCD) of 21 ± 1 inches. The nozzles were approximately 2 inches above the tank floor directly under them.



Figure 2.1. APEL 4 PJM Test Stand

Unlike conventional PJMs, whose operation is regulated by JPPs driven by compressed air, the APEL 4 PJM test system used a series of solenoid valves and a combination of an air compressor and vacuum pump to simulate the drive and suction phases of PJM operation. These operations were controlled through a control logic program using DASYLab data acquisition and control software (DACs), which turns on and off the appropriate solenoid valves at specified time intervals. The duration of each phase, the applied pressure, and vacuum are all variables that can be varied independently to simulate the operation of the PJMs.

Each PJM was outfitted with a Drexelbook liquid-level capacitance sensor/transmitter and an Endress+Hauser ceramic pressure transducer, which enabled continuous measurement of the slurry level and pressure inside the PJM during operation. Additional sensors included in the test system are Type K thermocouples that measure the temperature of the tank contents and the ambient temperature.

During the GR&R tests, in addition to the above parameters, the liquid level in the tank and the H_2O_2 (used to generate in situ oxygen bubbles to study gas behavior) flow rate and density were also monitored and recorded digitally. The liquid level in the tank outside the PJMs was monitored continuously using ultrasonic level detectors. The H_2O_2 flow rate and density were monitored using a 0.25-inch Micro-Motion Coriolis mass flow meter. During each mixing test, several variables such as PJM liquid levels and pressures, tank and ambient temperature, and H_2O_2 flow rate and density were monitored continuously and recorded digitally on a computer.

2.2 336 PJM Test Stand Description

The large-scale PJM test stand installed in Battelle's 336 test facility has been described extensively in previous reports (e.g., Bontha et al. 2003); therefore, only a brief description is presented here.

The PJM system consisted of four pulse tubes each with a cylindrical section of ~10 ft length and ~2 ft internal diameter. Each tube has a dished head with a connection to a 2-inch pipe. The bottom end of each pulse tube was tapered at an approximately 60° angle cone truncated to a ~4 inch nozzle. The overall height of the pulse tube, which is shown in Figure 2.2, was approximately 12 ft. A schematic of the experimental system used to demonstrate the PJM system is shown in Figure 2.3. As-built dimensions are detailed in Bontha et al. (2003).

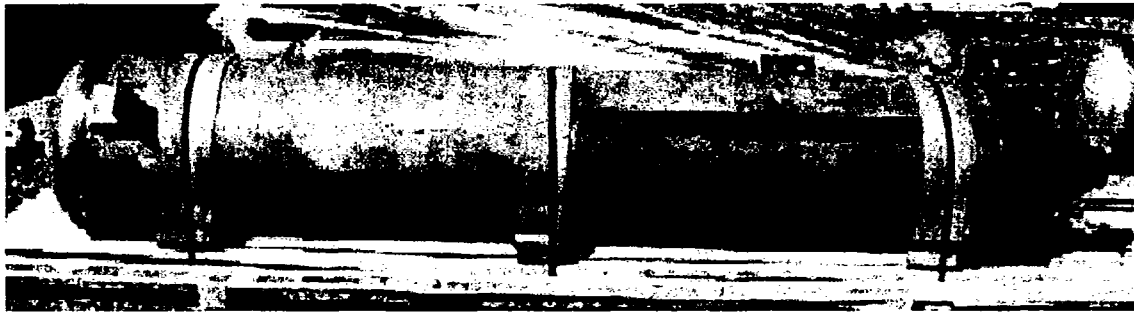


Figure 2.2. Photograph of the PJMs Used in the 336 Test Facility

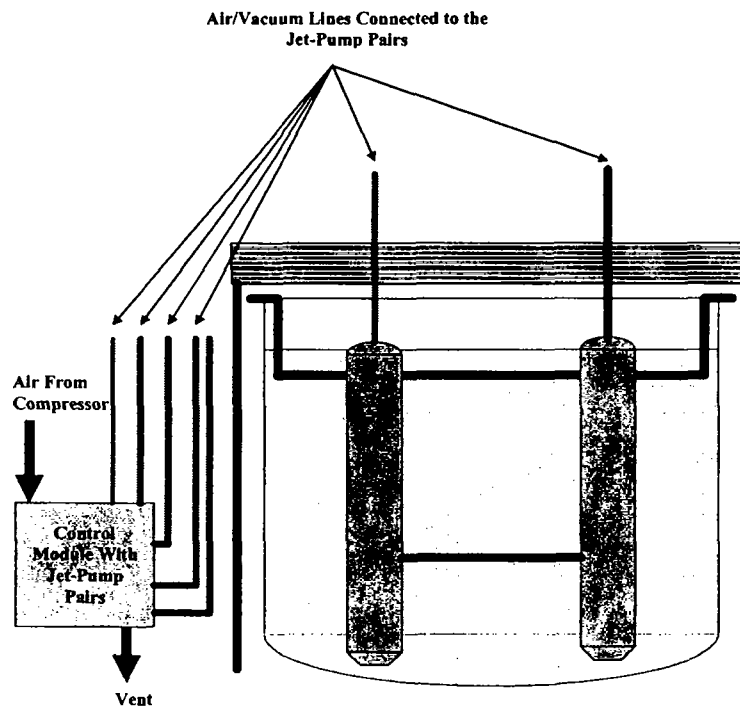


Figure 2.3. Schematic of Experimental System Used to Evaluate PJMs Using Non-Newtonian Simulants

The PJMs are inside a ~12.75-ft-ID x ~15-ft-tall supernate tank with a ~2:1 elliptical dish head. The nominal operating volume of the tank is about 10,000 gallons. The PJMs were held with brackets positioned on top of the tank. The brackets, which traverse the diameter of the tank and are welded to the sides, bear the weight of the tubes. The PJMs were positioned at the center of the four quadrants of the tank approximately 10 inches from the bottom of the tank.

During the operation of the PJMs, the pulse tubes were filled with the slurry by the application of a vacuum. The slurry was then expelled from the pulse tubes with compressed air. The suction and discharge of the slurry to and from the pulse tubes was regulated by JPPs in a control module on the ground level at the side of the tank. The JPPs were connected to the pulse tubes using 2-inch-OD wire-reinforced PVC tubing.

A compressor/accumulator(s) combination was used to regulate the air flow to the JPPs. The compressor chosen for the present study, which was based on the requirements for the air flow to the JPPs, was a Sullair compressor capable of delivering 1600 CFM at an operating pressure of 100 psig. The accumulators were an ASME standard 240 gal Brunner vertical air-receiver tank with pressure relief valves and a timed electronic drain valve. Both the compressor and the accumulators were located outside the 336 Building facility.

During the suction phase, liquid in the pulse tube piping can rise to a level of ~20 ft above the liquid level. To prevent suction of the liquid into the JPPs, the tubing connecting the pulse tubes to the JPPs was routed to the upper catwalk, ~40 ft above the top of the tank.

The sequence of operation and cycle frequency of the PJM and the RFD sampler was controlled by PRESCON™, an AEA Technology proprietary control system. Each PJM was outfitted with a Drexelbook liquid-level sensor/transmitter and a Ccomp pressure transducer that enabled continuous measurement of the slurry level and pressure inside the PJM during operation. Additional sensors included in the test system are Type K thermocouples for measuring the temperature of the tank contents and the ambient temperature. The data were digitally recorded on a computer using DASyLab DACS.

Simulant motion was detected either visually with the use of camera wells or with velocity probes (Figures 2.4 and 2.5). Video systems inserted into the camera wells were generally used to detect the mixed and unmixed regions in transparent simulants. A small video camera was moved up and down the camera well and the images recorded. The boundary of the mixed and unmixed regions was also recorded manually. Velocity probes were used to measure velocity although these results are not reported in this document.

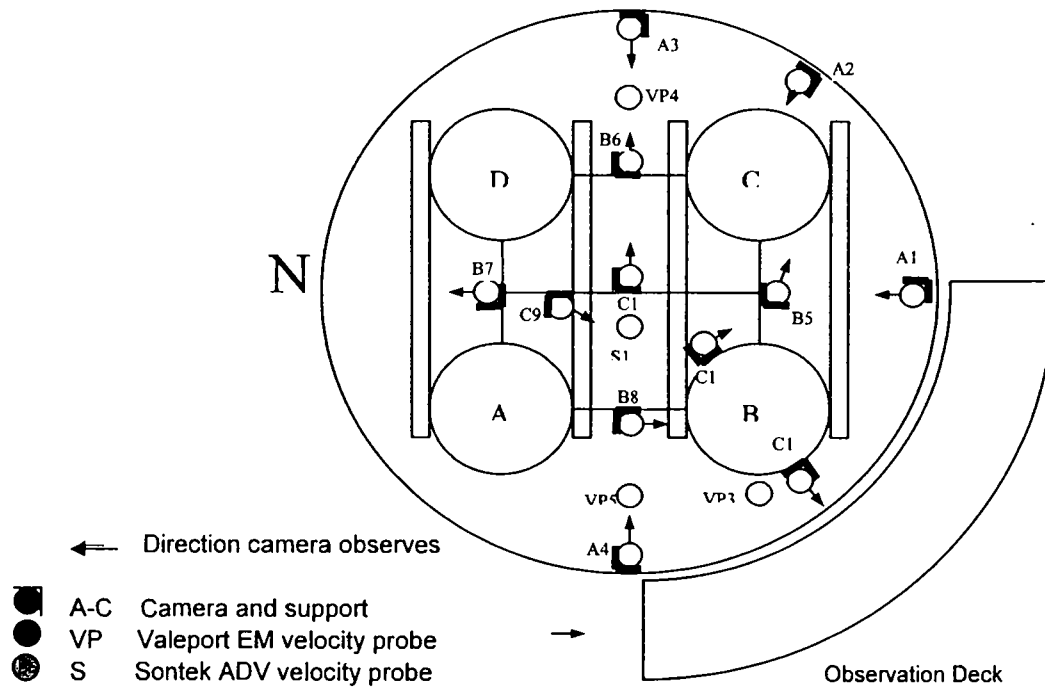


Figure 2.4. Plan View of the Instrument Locations for the 336 Building PJM 4 Test Stand

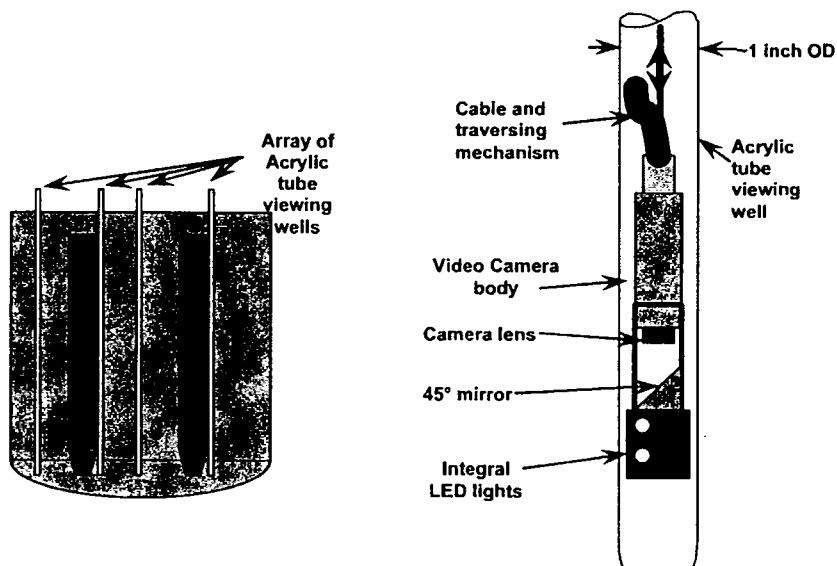


Figure 2.5. Video System for Detecting Mixed (cavern) Region in Large-Scale Testing

2.3 Scaled Prototypic Test Stands

2.3.1 UFP Prototype Vessel

The 168-inch-diameter, full-scale UFP tank was represented by a 34 ± 1 -inch-ID clear acrylic vessel. The geometric scale factor was ~ 4.94 . The scaled UFP prototypic test vessel was 91 ± 1 inches tall with a $\sim 2:1$ elliptical dish head made out of stainless steel. Mixing tests in this vessel were performed using different combinations of PJMs and spargers and a recirculation pump system. Top and plan views of the vessel and internals with nominal dimensions are shown in Figures 2.6 and 2.7. The various test sequences included in this document are presented in Table 2.1. The configuration details, subject to the constraints presented at the beginning of Section 2, are discussed below.

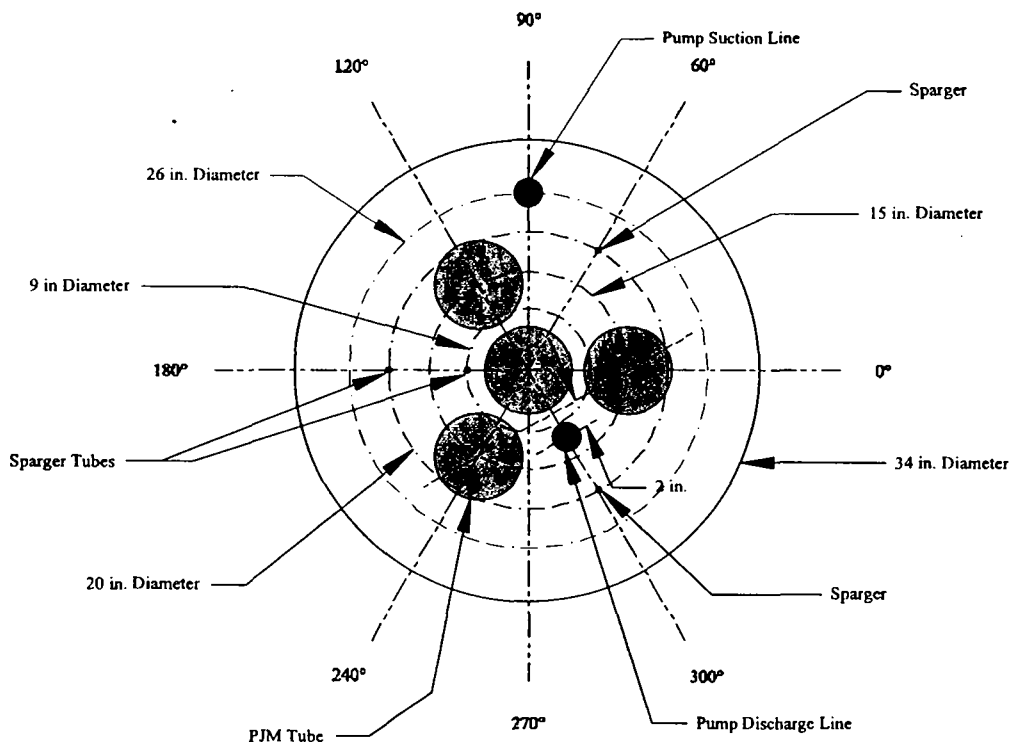


Figure 2.6. Top View of the UFP Prototypic Test Stand Showing Nominal Dimensions (measurement uncertainties are discussed in the text)

2.3.1.1 PJM Configurations

All of the PJMs for the UFP prototype were constructed from 6-inch-diameter (6.065-inch ID) schedule 40 stainless steel pipes with the end connected to an approximately 60° angle cone truncated to a 2-inch-diameter pipe fitting to which the nozzles were connected. The cylindrical section of the PJMs was 37 ± 1 inches tall; this corresponds to a PJM height scale factor of ~ 4.32 . The difference between the UFP tank dimension scale factor and the pulse tube dimension scale factor was due to the need to use

Table 2.1. UFP Test Sequences Presented in this Document and Corresponding PJM, Sparger, and Recirculation Pump Configurations^(a,b)

Seq No	Run	Test Type	Test Mode	PJM Configuration				Sparger Configuration			Recirc. Pump Discharge Configuration			
				PJM Arrangement	Nozzle Type	Noz. Dia. (in)	Elevation (in) ^(c)	No. of Spargers	Radial Pos.	Elevation (in) ^(c)	Noz. Dia (in)	PCD (in)	Elevation (in) ^(c)	Angle ^(d)
2	1	Mixing	PJM Only	Tri-Foil (3+1)	45° (3); Vertical (1)	0.824	2	-	-	-	-	-	-	-
2	2	Mixing	PJM Only	Tri-Foil (3+1)	45° (3); Vertical (1)	0.824	2	-	-	-	-	-	-	-
2	3	Mixing	PJM + Sparging	Tri-Foil (3+1)	45° (3); Vertical (1)	0.824	2	1 (near center)	Figure 2.6	-	-	-	-	-
2	4	Mixing	PJM + Sparging	Tri-Foil (3+1)	45° (3); Vertical (1)	0.824	2	3 (Outer)	Figure 2.6	-	-	-	-	-
3B	1	Mixing	PJMs Only	Tri-Foil (3+1)	45° (3); Vertical (1)	0.824	2	-	-	-	-	-	-	-
3B	2	Mixing	PJM + Pump	Tri-Foil (3+1)	45° (3); Vertical (1)	0.824	2	-	-	-	1.049	11	24	0°, Down
3B	3	Mixing	PJM + Pump	Tri-Foil (3+1)	45° (3); Vertical (1)	0.824	2	-	-	-	1.049	11	24	0°, Down
3B	4	Mixing	PJM + Pump + Sparging	Tri-Foil (3+1)	45° (3); Vertical (1)	0.824	2	1 (near center)	Figure 2.6	4	1.049	11	24	0°, Down
5	1	GR&R	PJM + Sparging	Tri-Foil (3+1)	45° (3); Vertical (1)	0.824	2	1 (near center)	Figure 2.6	4	-	-	-	-
5	2	GR&R	PJM + Sparging	Tri-Foil (3+1)	45° (3); Vertical (1)	0.824	2	1 (near center)	Figure 2.6	4	-	-	-	-
5	3	GR&R	PJM + Pump	Tri-Foil (3+1)	45° (3); Vertical (1)	0.824	2	-	-	-	1.049	11	24	0°, Down
5	4	GR&R	PJM + Sparging	Tri-Foil (3+1)	45° (3); Vertical (1)	0.824	2	1 (near center)	Figure 2.6	4	-	-	-	-
6	1	GR&R	PJM + Sparging	Tri-Foil (3+1)	45° (3); Vertical (1)	0.824	2	1 (near center)	Figure 2.6	4	-	-	-	-
6	2	GR&R	PJM + Sparging	Tri-Foil (3+1)	45° (3); Vertical (1)	0.824	2	1 (near center)	Figure 2.6	4	-	-	-	-
6	3	GR&R	PJM + Sparging	Tri-Foil (3+1)	45° (3); Vertical (1)	0.824	2	1 (near center)	Figure 2.6	4	-	-	-	-
6	4	GR&R	PJM + Sparging	Tri-Foil (3+1)	45° (3); Vertical (1)	0.824	2	1 (near center)	Figure 2.6	4	-	-	-	-
7 ^(e)	1-4	Solids Lift	PJMs only	Cluster (5+1)	45° (5); Vertical (1)	0.824	2	-	-	-	-	-	-	-

(a) Test results discussed in Section 3.

(b) Configuration spatial and dimensional distances values in table do not reflect the type of measurement or accuracy. See text for details.

(c) Approximate distance from the bottom of the tank under the nozzle

(d) Angle from vertical.

(e) Configuration selected.

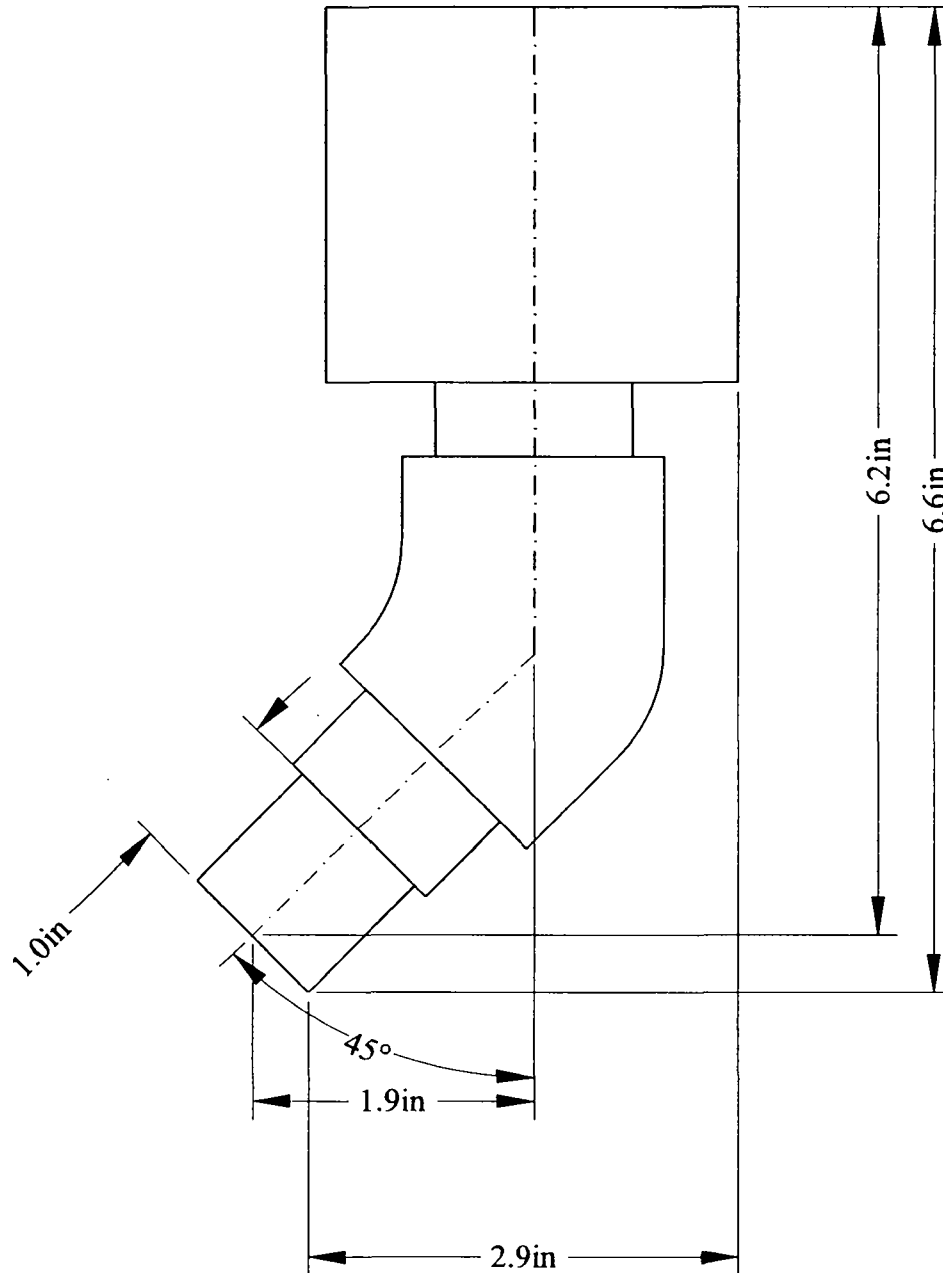


Figure 2.8. Schematic of the 45° Nozzle Used in the UFP Test Stand Showing Nominal Dimensions (measurement uncertainties are discussed in the text)

and sparger hybrid configurations without obstruction of the spargers by additional PJMs. Because the purpose of PJMs was to generate a mixed region at the bottom of the vessel with the spargers extending the mixed region to the vessel surface, four PJMs were suitable for testing. The actual vessel will have six PJMs, which will provide a larger mixed region. The final test sequence (#7 in Table 2.1) used six PJMs.

The center PJM nozzle was constructed from 0.75-inch (0.824-inch ID) schedule 40 stainless steel pipe and was pointed straight down toward the center of the tank bottom and raised approximately 2 inches off the bottom. The perimeter PJM nozzles were constructed from 0.75-inch (0.824-inch ID), schedule 40 stainless steel pipe angled 45° (using a standard 45° elbow fitting) radially outward from the tank center and raised approximately 2 inches off the tank floor. Figure 2.8 is a schematic of the 45° nozzle; all dimensions listed are within ± 0.5 inches. All tests with the PJMs were conducted using a target stroke of 33.5 to 35.5 inches (85 to 90 cm) and a target average nozzle velocity of 8 ± 0.8 or 12 ± 1 m/s.

2.3.1.2 Sparger Configuration

Tests using spargers were performed using an array of four (one center and three perimeter) spargers. The center sparger was approximately midway between adjacent perimeter PJMs at a radial position of approximately 4.5 inches from the tank centerline. The perimeter spargers were placed at approximately midway between adjacent perimeter PJMs at a pitch diameter of 20 ± 1 inches. All of the sparger tubes were made from 0.5-inch-OD (0.37 inch ID) stainless steel tubing, and the lower ends of the sparger tubes were approximately 4 inches above the bottom of the tank as measured from the tank floor (or approximately 2 inches above the tip of the nozzle). Tests with sparging were carried out either with the center sparger operating at a target flow rate of 3 acfm or the perimeter spargers operating at a target flow rate of 1 acfm each.

2.3.1.3 Recirculation System Configuration

The pump recirculation system consisted of two centrifugal pumps placed in parallel and connected in series with a diaphragm pump that served to eliminate cavitation and prime the centrifugal pumps. The recirculation pump system was operated at a target flow rate of 90 ± 5 gpm (which corresponds to ~ 2200 gpm at full scale), and the discharge line nozzle was sized such that the linear velocity exiting the nozzle was ~ 30 ft/sec.

For the test sequences presented in this document, the recirculation configuration consisted of a single discharge line of 2-inch (2.067 inch ID) schedule 40 stainless steel pipe with a 1-inch (1.049-inch ID) schedule 40 stainless steel nozzle pointing down. It was located approximately midway between two of the perimeter PJMs at a radial position of approximately 5.5 inches from the tank centerline and an elevation of approximately 24 inches from the bottom center of the tank floor. The pump suction line consisted of a 2-inch (2.067-inch ID) schedule 40 PVC pipe located at a radial position of approximately 4 inches from the tank wall on the opposite side of the tank from the discharge line and at an elevation of approximately 4 inches as measured from the center of the intake to the tank floor beneath it.

2.3.2 Lag Storage Prototypic Vessel

The 300-inch-diameter, full-scale LS tank was represented by a 70 ± 1 -inch ID clear acrylic vessel. The scale factor was ~ 4.29 . The scaled LS prototype acrylic vessel was 91 ± 1 inches tall with a 100:6 elliptical dish head made of stainless steel. Mixing tests in this vessel were performed using different combinations of PJMs and spargers and the recirculation pump system. Top and plan views of the vessel and internals are shown in Figures 2.9 and 2.10. The various test sequences included in this document are presented in Table 2.2. The configuration details, subject to the constraints presented at the beginning of Section 2, are discussed below.

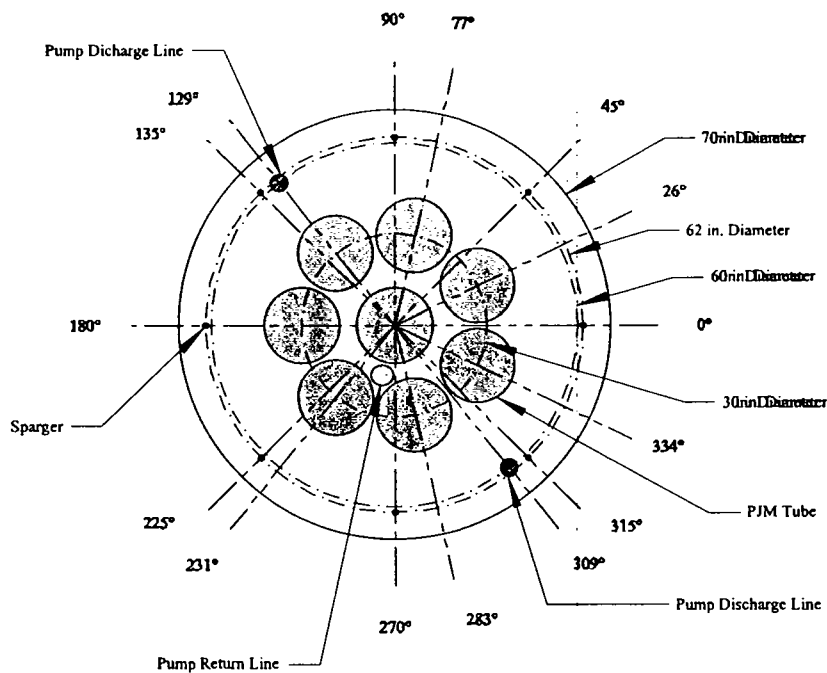


Figure 2.9. Top View of the LS Prototype Test Stand Showing Nominal Dimensions (measurement uncertainties are discussed in the text)

2.3.2.1 PJM Configurations

All the PJMs for the LS prototype were constructed from 12-inch-diameter (12-inch ID) schedule 40 stainless steel pipe with an approximately 60° angle cone truncated to a 2-inch-diameter pipe fitting to which the nozzles were connected. The height of the cylindrical section of the PJMs was 31 ± 1 inches, corresponding to a PJM height scale factor of ~ 4.93 . The difference between the LS tank and pulse tube dimension scale factors is due to the need to use standard pipe sizes for procurement expediency. However, the volume expelled from the PJMs was consistent with the LS scale factor of ~ 4.29 .

For the LS test sequences presented in this document, the PJM array consisted of eight PJMs, with one near the center of the tank and the other seven nearly equally spaced around the center PJM on a pitch diameter of 30 ± 1 inches. This was referred to as the “cluster” configuration.

For all but one test sequence (# 20) presented in this document, the center PJM nozzle was constructed from 1-inch (1.049-inch ID) schedule 40 stainless steel pipe pointed straight down toward the center of the tank bottom and raised approximately 2 inches off the bottom. For sequence 20, a 1-inch (0.957-inch ID) schedule 80 stainless steel pipe was used for the center nozzle. For all test sequences presented in this document, the perimeter PJM nozzles were constructed from 1-inch (0.957 inch ID), schedule 80 PVC pipe, angled 45° (using a standard 45° elbow fitting) radially outward from the tank center and raised approximately 2 inches off the tank floor. Figure 2.11 is a schematic of the 45° angled nozzle; all dimensions in this figure are within ± 0.5 inches. All tests with the PJMs were conducted using a target stroke of 29.5 to 31.5 inches (75–80 cm) and a target average nozzle velocity of 8 ± 0.75 or 12 ± 1 m/s.

Table 2.2. LS Test Sequences Presented in this Document and the Corresponding PJM, Sparger, and Recirculation Pump Configurations^(a, b)

Seq No.	Run	Test Type	Test Mode	PJM Configuration				Sparger Configuration				Recirc. Pump Discharge Config.				
				PJM Arrangement	Nozzle Type	Noz. Dia. (in)	Elevation (in) ^(c)	No.	OD (in)	PCD (in)	Elevation (in) ^(c)	No. of Nozzles	Nozzle dia (in.)	PCD (in)	Elevation (in) ^(c)	Angle ^(d)
4	1	Mixing	PJMs Only	Cluster (7+1)	45° (7) Vertical (1)	(7) 0.957 (1) 1.049	2	-	-	-	-	-	-	-	-	-
4	2	Mixing	PJMs Only	Cluster (7+1)	45° (7) Vertical (1)	(7) 0.957 (1) 1.049	2	-	-	-	-	-	-	-	-	-
4	3	Mixing	PJMs + Spargers	Cluster (7+1)	45° (7) Vertical (1)	(7) 0.957 (1) 1.049	2	4	0.5	61	5	-	-	-	-	-
4	4	Mixing	PJMs + Spargers	Cluster (7+1)	45° (7) Vertical (1)	(7) 0.957 (1) 1.049	2	8	0.5	61	5	-	-	-	-	-
7	1	Mixing	PJMs Only	Cluster (7+1)	45° (7) Vertical (1)	(7) 0.957 (1) 1.049	2	-	-	-	-	-	-	-	-	-
7	2	Mixing	PJMs Only	Cluster (7+1)	45° (7) Vertical (1)	(7) 0.957 (1) 1.049	2	-	-	-	-	-	-	-	-	-
7	3	Mixing	PJMs + Pump	Cluster (7+1)	45° (7) Vertical (1)	(7) 0.957 (1) 1.049	2	-	-	-	-	4	0.622	60	29	30° Up
7	4	Mixing	PJMs + Pump	Cluster (7+1)	45° (7) Vertical (1)	(7) 0.957 (1) 1.049	2	-	-	-	-	4	0.622	60	29	30° Up
7	5	Mixing	PJMs + Pump + Sparging	Cluster (7+1)	45° (7) Vertical (1)	(7) 0.957 (1) 1.049	2	4	0.5	61	5	4	0.622	60	29	30° Up
7	6	Mixing	PJMs + Pump + Sparging	Cluster (7+1)	45° (7) Vertical (1)	(7) 0.957 (1) 1.049	2	8	0.5	61	5	4	0.622	60	29	30° Up
11	1	Mixing	PJMs + Pump	Cluster (7+1)	45° (7) Vertical (1)	(7) 0.957 (1) 1.049	2	-	-	-	-	2	0.91	60	16	30° Up
11	2	Mixing	PJMs + Pump	Cluster (7+1)	45° (7) Vertical (1)	(7) 0.957 (1) 1.049	2	-	-	-	-	2	0.91	60	16	30° Up
15	1	GR&R	PJMs + Spargers	Cluster (7+1)	45° (7) Vertical (1)	(7) 0.957 (1) 1.049	2	4	0.5	61	5	-	-	-	-	-
15	2	GR&R	PJMs + Spargers	Cluster (7+1)	45° (7) Vertical (1)	(7) 0.957 (1) 1.049	2	4	0.5	61	5	-	-	-	-	-
15A	3	GR&R	PJMs + Pump	Cluster (7+1)	45° (7) Vertical (1)	(7) 0.957 (1) 1.049	2	-	-	-	-	4	0.622	60	16	30° Up
15A	4	GR&R	PJMs + Spargers	Cluster (7+1)	45° (7) Vertical (1)	(7) 0.957 (1) 1.049	2	4	0.5	61	5	-	-	-	-	-
16	1 - 6	Solids Lift	PJMs Only	Cluster (7+1)	45° (7) Vertical (1)	(7) 0.957 (1) 1.049	2	-	-	-	-	-	-	-	-	-
20	1	Mixing	PJMs	Cluster (7+1)	45° (7) Vertical (1)	(7) 0.957 (1) 0.957	2	-	-	-	-	-	-	-	-	-
20 ^(d)	2	Mixing	PJMs + Pump	Cluster (7+1)	45° (7) Vertical (1)	(7) 0.957 (1) 0.957	2	-	-	-	-	2	0.803	60	14	25° Up

(a) Test results discussed in Section 3.
(b) Configuration spatial and dimensional distances values in table do not reflect the type of measurement or accuracy. See text for details.
(c) Approximate distance from the bottom of the tank under the nozzle
(d) Angle from vertical.
(e) Configuration selected.

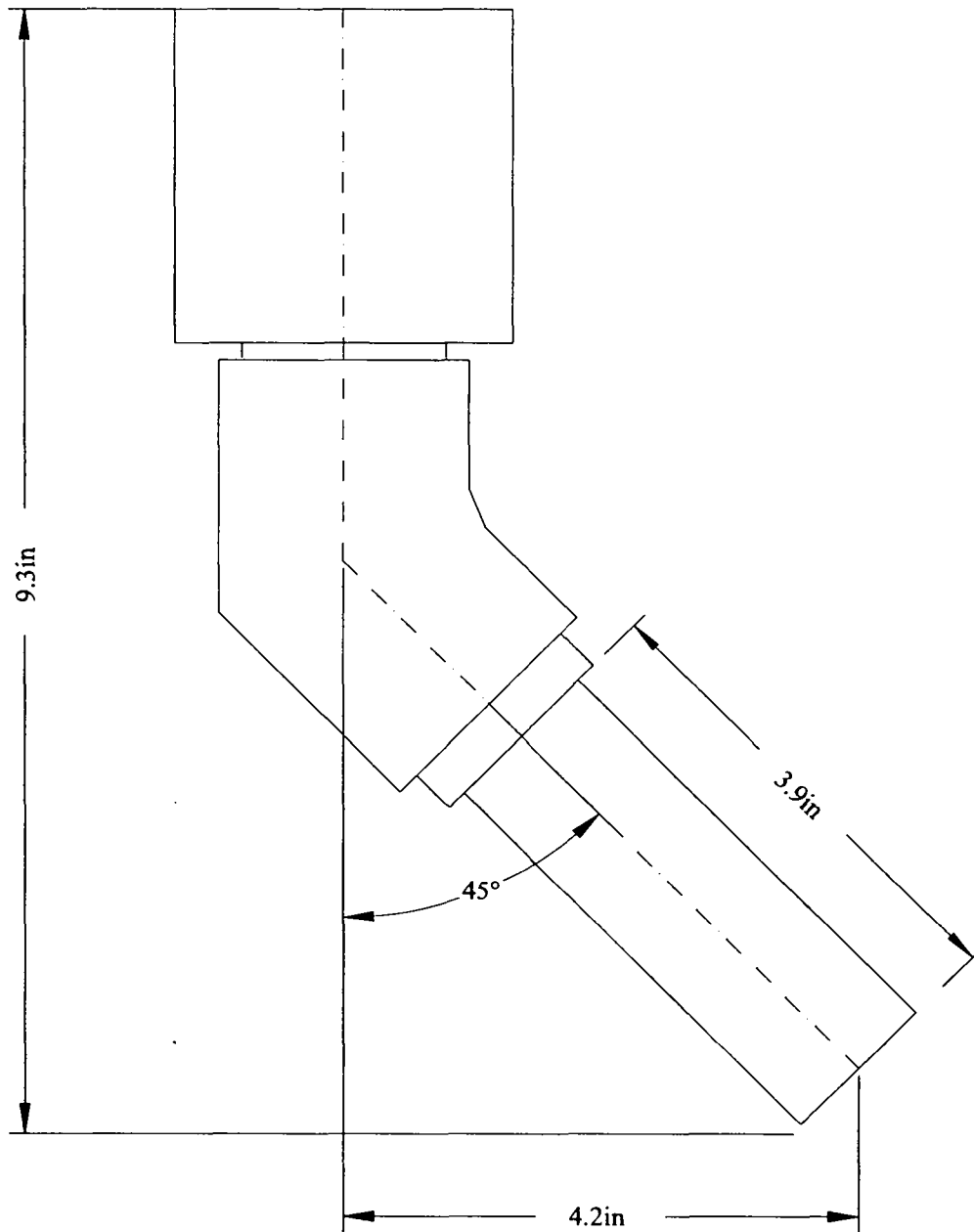


Figure 2.11. Schematic of the 45° Nozzle in the LS Prototype Test Stand Showing Nominal Dimensions (measurement uncertainties are discussed in the text)

2.3.2.3 Recirculation System Configuration

The pump recirculation system consisted of two centrifugal pumps placed in parallel and connected in series with a diaphragm pump that served to eliminate cavitation and prime the centrifugal pumps. The recirculation pump system was operated at a target flow rate of 120 ± 5 gpm (which corresponds to ~ 2200 gpm at full-scale). The nozzles at the discharge were generally sized such that the nominal linear velocity was ~ 30 ft/sec, although during Sequence 20 tests with two discharge nozzles, the nozzle diameter was selected to provide a linear velocity of ~ 40 ft/sec.

For the test sequences presented in this document, the pump suction consisted of 3-inch (2.900-inch ID) schedule 80 PVC pipe. The end of the suction line had several 1.5-inch holes drilled along its side to provide additional area for simulant flow. For all test sequences except Sequence 11, the suction line was in the space between the center and two adjacent perimeter PJMs. For Sequence 11, the suction line was roughly at a distance midway between the two discharge lines at a radial position of about 30 inches from the tank centerline. Except for Sequence 20, the elevation of the suction line varied from 4 to 12 inches above the tank floor during the testing to minimize cavitation due to its proximity to the spargers or to minimize its influence during dye injection near the bottom of the tank. In Sequence 20, the suction line was 10 ± 1 inches off the tank floor, as measured directly under the nozzle and shown in Figure 2.11.

In this document, four sequences of tests conducted with the recirculation pump are presented. This includes two tests with two discharge nozzles (sequences 11 and 20) and two tests with four discharge nozzles (sequences 7 and 15A). A schematic of the recirculation nozzle used in Sequence 20 is shown in Figure 2.12; all dimensions listed in this figure are within ± 0.5 inches.

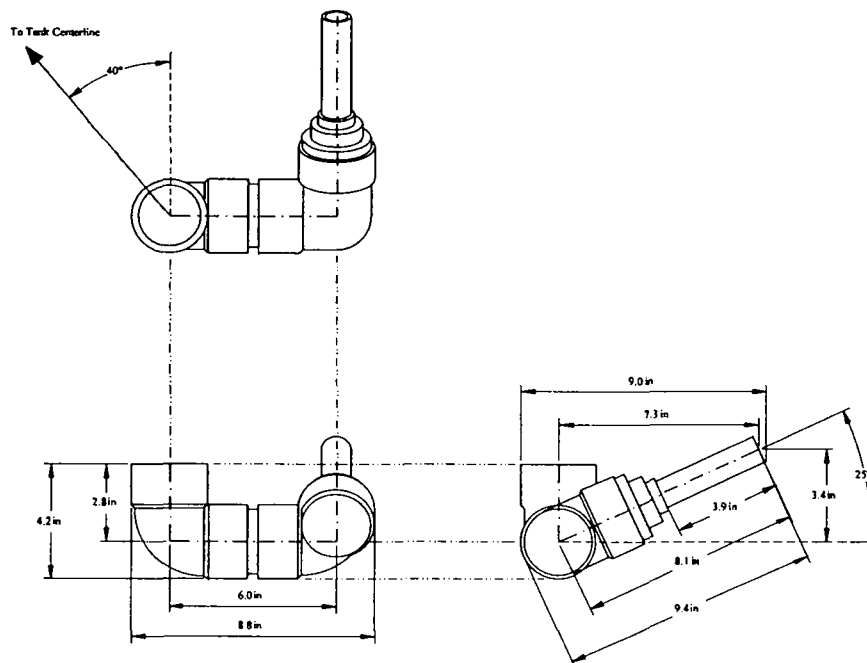


Figure 2.12. Recirculation Pump Discharge Nozzles Used in LS Sequence 20 (measurement uncertainties are discussed in the text)

All discharge lines were constructed out of 2-inch (2.067-inch ID) schedule 40 stainless steel pipe. The first test with two discharge nozzles (Sequence 11) consisted of 0.75-inch schedule 40 stainless steel pipes that were bored out to a 0.91 ± 0.01 -inch ID. The second test with two discharge nozzles (Sequence 20) consisted of 0.75-inch schedule 80 stainless steel pipes that were bored out to a 0.80 ± 0.01 -inch ID. The nozzles for the two test sequences with four discharge lines (Sequences 7 and 15) consisted of 0.5-inch (0.622-inch ID) schedule 40 stainless steel pipe.

For Sequences 11 and 20, the nozzles were diagonally opposite each other in the tank at a pitch diameter of 60 ± 1 inches. For Sequence 11, the nozzles were pointed upward at an angle of approximately 30° and raised to an elevation of approximately 16 inches. The nozzle angles formed by the line from the center of the tank to the center of the discharge line and the line passing through the discharge nozzle were approximately 40° pointing inward. For Sequence 20, the nozzles were pointed up at an angle of $25 \pm 2.5^\circ$ and raised to a 14 ± 1 -inch elevation. The nozzle angles formed by the line from the center of the tank to the center of the discharge line and the line passing through the discharge nozzle were $40^\circ \pm 5^\circ$ pointing inward and are shown schematically in Figure 2.13.

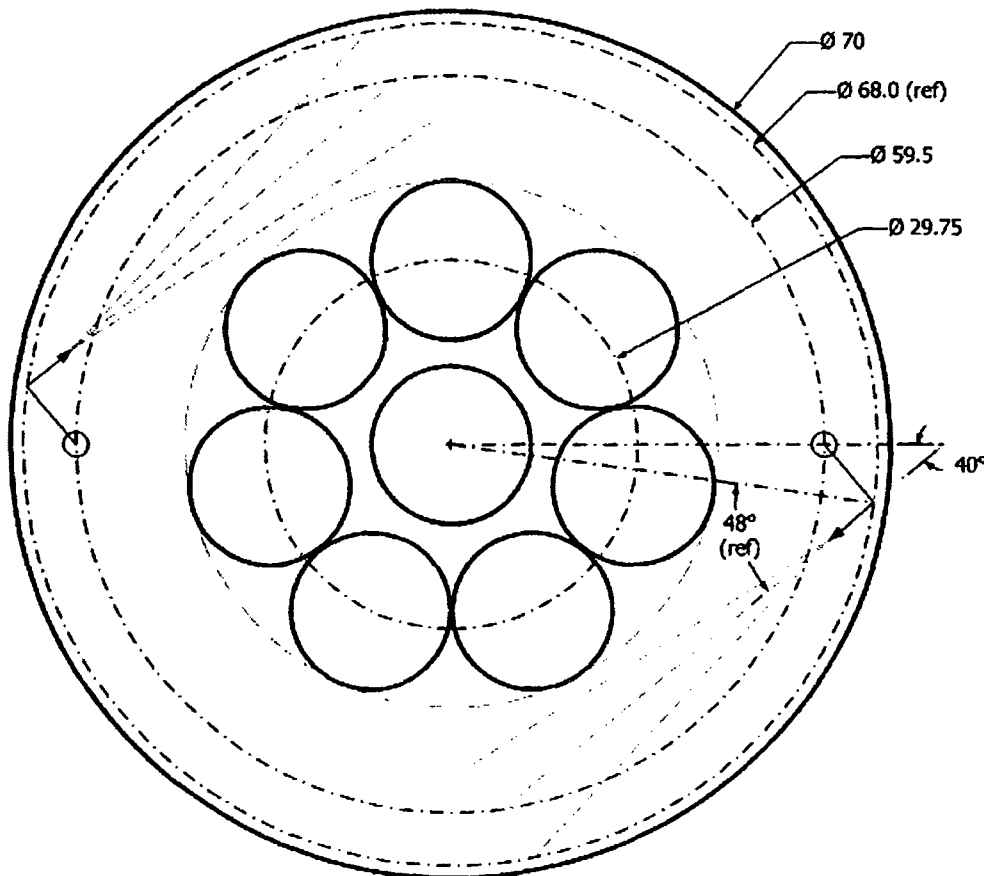


Figure 2.13. Top View of the Recirculation Nozzles in the Tank for LS Sequence 20 (measurement uncertainties are discussed in the text)

For Sequences 7 and 15, the nozzles were located along the four corners of a nearly square rectangle at a pitch diameter of 60 ± 1 inches. For Sequence 7, the nozzles were pointed upward at an angle of approximately 30° and raised to an elevation of approximately 29 inches relative to the bottom center of the tank. The nozzles were pointed approximately tangential to the tank wall. For Sequence 15, the nozzles were pointed up at an angle of approximately 30° and raised to an elevation of approximately 16 inches relative to the bottom center of the tank. The nozzles were approximately tangential to the tank wall.

2.3.3 System Operation and Data Acquisition

Unlike conventional PJMs, whose operation is regulated by JPPs driven by compressed air, the prototype test systems used a series of solenoid valves and a combination of an air compressor and a vacuum pump to simulate the drive and suction phases of PJM operation. These operations were controlled through a control logic program using DASyLab that turns the appropriate solenoid valves on and off at specified time intervals. The duration of each phase, the applied pressure, and the vacuum are all variables that can be independently varied to simulate the operation of the PJMs. The PJMs were operated at a specific average nozzle velocity (\bar{u}_{disch}), which is defined as

$$\bar{u}_{disch} = \frac{\Delta H}{\Delta t} * AR \quad (2.1)$$

where ΔH is the length of the PJM stroke, Δt is the time for achieving the stroke, and AR is the area ratio of the PJM to the nozzle. This equation is the same as Equation A.7 in Appendix A.

In addition to the PJM operation, the recirculation pump flow rates were controlled using a variable frequency drives (VFDs) on the centrifugal pumps and the air pressure to the diaphragm pump. Finally, the sparger air flow rates were controlled using rotameters.

During each mixing test, several variables such as PJM liquid levels and pressures, tank and ambient temperatures, recirculation pump flow rate, and density were monitored continuously and recorded digitally on a computer. The liquid/slurry level inside each of the PJMs was measured using Drexelbook capacitance level probes and transmitters. The functionality of the level probes was checked prior to the start of a sequence of tests which typically ran from 4 to 8 hours. Compressor and vacuum supply pressures and the pressure inside each PJM were monitored using Endress+Hauser ceramic pressure transducers. The tank and ambient temperatures were measured using Type K thermocouples. The flow rate and density of the slurry from the recirculation pump was measured using a 3-inch MicroMotion Coriolis mass flow meter. In addition to these variables, which were digitally monitored, the sparger air flow rates and pressures were recorded manually on the run data log sheets or in the project laboratory record books (LRBs).

During the GR&R tests, in addition to the above parameters the liquid level in the tank and the H_2O_2 flow rate/density were also monitored and recorded digitally. The liquid level in the tank outside the PJMs was monitored continuously using ultrasonic level detectors. In addition, the H_2O_2 flow rate/density was monitored using a 0.25-inch MicroMotion Coriolis mass flow meter.

2.3.4 Mixing Effectiveness Determination

The primary measurement in the scaled prototypic test platforms is the size and extent of the mobilization cavern resulting from PJM operations and PJMs combined with recirculation (i.e., steady jet) and/or sparging. This was achieved using a chemical tracer method discussed in detail in Section 2.6 and Appendix B. This section deals only with the method in which the tracer was injected into the tank and how the samples were collected.

The required amount of tracer (typically Brilliant Blue dye in an amount equal to ~ 5 g per 100 gal of clay simulant in the tank) was mixed with ~2 liters of the same clay simulant that was used in the testing. The concentrated tracer/clay mixture was injected prior to the start of a sequence of tests at lowest nozzle velocity of that test sequence. The concentrated tracer slurry was injected into the center PJM during the vacuum phase of the PJM cycle over a period of approximately 10 minutes. Once tracer injection was completed, the tracer injection line was purged with clean clay to ensure complete transfer of the tracer into the PJM. Once the line was purged, simulant samples from the tank were collected over a period of at least 45 minutes of PJM operation. Samples were withdrawn at various times from five different sample lines installed in the PJMs and the tank. Three of these samples were drawn from three perimeter PJMs and the remaining two samples were drawn from the annulus between the PJM and tank wall at elevations representing the lower and upper halves of the tank, respectively. After completion of the specified run conditions, the tank was completely homogenized and final homogenized samples collected. Comparison of the tracer concentration in the various samples with the final homogenized samples provides the percent mixed as a function of time and run conditions. Complete and successful mixing is defined as 100% as indicated by the chemical tracer method.

2.3.5 Solids Suspension Under Turbulent Conditions

Under some conditions the rheology will be low and solids may settle to the bottom of the tank. PJMs are well designed to pick up such solids because they direct a turbulent jet against the bottom of the tank. Solids suspension in mechanically stirred tanks is characterized by the "just suspended" criteria developed by Zwietering (1958; Atiemo-Obeng 2003), where no solids remain on the bottom of the tank for more than a few seconds. The BHRG-FMP consortium has shown that for steady downward-pointing jets an equation of functionality similar to that of Zwietering can be developed.^(a) The same form and functionalities would be expected to apply for multiple pulsed jets.

$$V_{js} = K * (\Delta\rho)^{0.43} (dp)^{0.2} X^{0.14} \quad (2.2)$$

where

- V_{js} = minimum velocity to suspend solids
- $\Delta\rho$ = density difference between solids and liquid
- dp = maximum particle size
- X = wt% of solids.

To determine the solids suspension characteristics of several of the pulse jet mixed tanks in the WTP area, tests were run similar to those done by Zwietering and FMP. A small concentration of 4-mm glass beads

(a) Personal communication with FMP on "Jet Solid Suspension Design Guide." FMP report 064.

was placed in the bottom of the tank and the PJM velocity increased in increments until the solids were observed to lift off the bottom. Many workers have shown that visual and instrumentation methods for determining the just-suspended velocity give very similar results (Brown et al. 2003).

The concentrations used in the Zwietering terms were 0.4 to 0.5 wt%. The Zwietering and FMP correlations show that the minimum velocity to pick up solids is a weak function of solids fraction and particle size and mainly depends on the density difference. Thus, using dense glass (2500 kg/m^3) and large particles gives a good estimate of the exact velocity required and makes observation easier.

2.3.6 Visual Observations During (Dye) Tracer Tests

Visual observations of the tank surface and walls supplemented the understanding of the test results. General observations were made to characterize flow conditions on the tank surface, including easily observed upwelling of material due to PJM discharge, recirculation pump operation, or air sparging. Because in all experiments the chemical tracer was Brilliant Blue dye, observations of the slurry surface were made to verify that dye did not prematurely break through the surface during tracer injection. The surface was also monitored during the run to determine whether dye broke through the surface due to upwelling of new slurry onto the surface. A video camera recorded the simulant surface image during each test. The tank walls were monitored during tracer dye injection to verify that the perimeter PJMs were discharging dyed slurry. After dye injection, the tank walls were monitored for evidence of dyed slurry spreading upward and/or laterally along the wall. Dry erase markers were used to map dyed areas on the tank wall and for sketching a cylindrical projection map of the dyed areas on the acrylic tank wall. The markings on the tank wall were also recorded with a video recorder. Mapping tracer locations along the tank walls supplemented interpretation of tracer on the slurry surface for breakthrough due to cavern growth, flow due to the spargers or pump recirculation, and interpretation of tracer sampling results. In some runs, direct evidence was observed of turbulence due to air spargers or PJM discharges, which were seen as a rippling effect extending up the tank wall at specific locations. This supplemented the tracer observations of cavern height at the tank wall. Observations of dark particulates entrained in the slurry at the tank wall were also made to follow flow lines during some of the recirculating pump operations, particularly of flow toward the pump return line.

2.3.7 Specific Observations During Gas Retention and Release Tests

During GR&R tests the tank level changes were monitored to determine the gas fraction in the simulant. The retained gas volume was estimated by measuring the simulant level changes referenced to the simulant level with no retained gas (see Section 2.5 for details). While the liquid levels were recorded digitally by the DACS, observers visually recorded slurry levels along the tank walls at three locations using tape measures attached to the tank wall and/or suspended from the tank rim. In addition, a video recording was made of one of these stations. These observations supplemented data collected from ultrasonic level indicators and were used to interpret gas release and holdup. In addition, periodic general observations were made as appropriate to the run to aid in interpretation of the tests.

2.4 Sparger Testing

To assess the performance of spargers in WTP actual waste rheology bounding non-Newtonian slurries near full scale, an experimental apparatus consisting of a large-scale tank with a single sparge point was used. The tank used had an approximately 10-ft ID conical bottom and a height of about 12 ft. This vessel was filled with a kaolin:bentonite simulant with Bingham plastic rheological parameters of 25 cP consistency and 35 Pa yield stress. A 3/8-inch schedule 40 pipe was used as a sparge point. The tube was immersed in the simulant to a known depth, and air was forced through the pipe. A rotameter coupled with a pressure gauge and thermocouple was used to determine the actual volumetric air flow through the pipe (Figure 2.14).

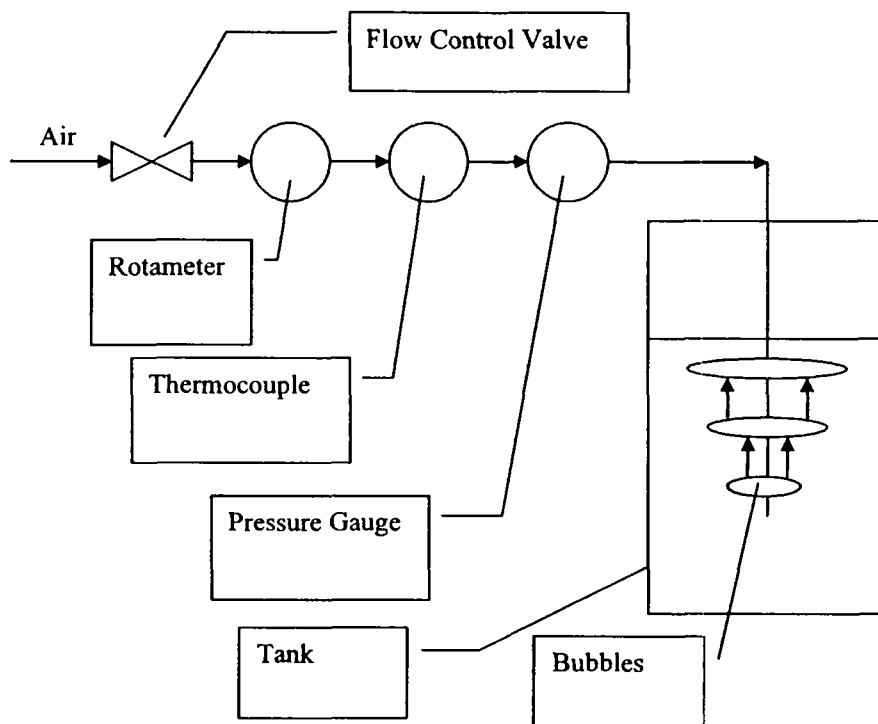


Figure 2.14. Diagram of Sparging Experimental Setup

The rotameter, pressure gauge, and thermocouple readings were converted to the actual volumetric air flow in the slurry at the sparger submergence depth. Isothermal expansion at the sparge tube orifice was assumed. The following equation is used for this calculation:

$$Q_{sparge} = \frac{Q_{rot}}{\rho_{sim} g h_{sparge} + P_0} \left(\frac{P_{std} P_{rot} T_{rot}}{T_{std}} \right)^{\frac{1}{2}} \quad (2.3)$$

where

Q_{sparge}	is actual volumetric air flow in the slurry at the submergence depth of sparge tube (ft ³ /min)
Q_{rot}	is the calibrated volumetric flow rate read from the rotameter at standard conditions (ft ³ /min)
P_{std}	is the absolute pressure at which the rotameter is calibrated (1 atm)
T_{std}	is the absolute temperature at which the rotameter is calibrated (530 R)
P_{rot}	is the absolute pressure read at the rotameter
T_{rot}	is the absolute temperature read at the rotameter
ρ_s	is the bulk density of the simulant tested (1.2 g/mL)
g	is the gravitational constant (9.81 m/s ²)
h_s	is submerged depth of the sparge tube
P_0	is atmospheric pressure (1 atm)

Measurements on the surface of the simulant were made to determine the areas of the tank affected by the upward motion of the bubbles. Two major areas were measured. The first area is referred to as the region of bubbles (ROB). This area is the region surrounding the sparge tube that contains the plume of bubbles rising from the sparge point. Surrounding the ROB is an area of induced radial and downward flow. This area is referred to as the zone of influence (ZOI). Both of these regions are reported as diameters. These concepts are portrayed graphically in Figure 2.15.

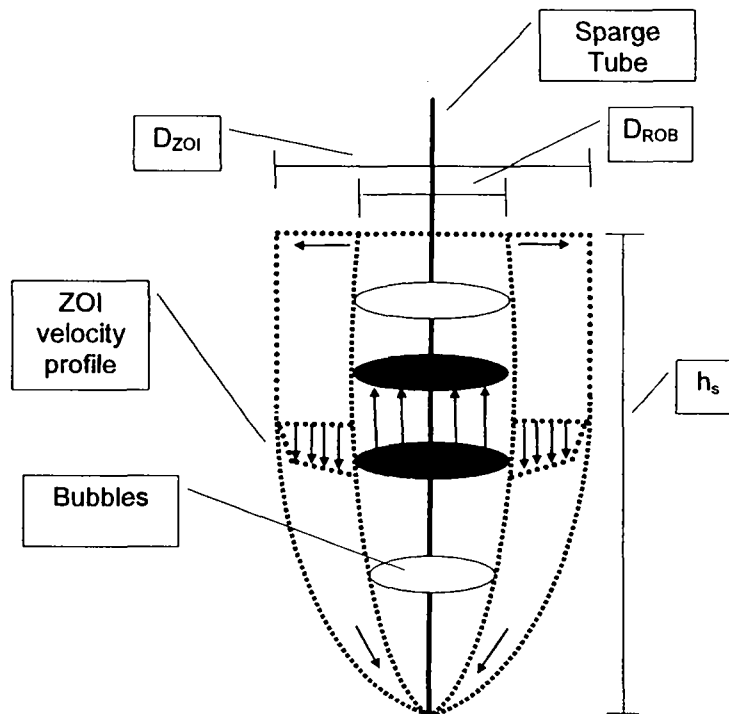


Figure 2.15. Diagram of Sparging Experiment Concepts

The radii of the ZOI and ROB were measured with the sparge tube inserted to several submergence depths and operated with a range of air flow rates. The ROB was measured through the use of a laser reference system coupled with video imaging software. This technique involves placing two laser points

a known distance apart on the surface of the simulant. As testing proceeds, the surface of the simulant is video recorded. The frames from the video images are then analyzed to determine the number of pixels between various features in the frame. By knowing the number of pixels and the actual distance between the laser reference points, the actual distance between two features on the frame were determined.

After the steady-state flow due to sparging was achieved,^(a) the video images of the surface were collected. The technique described above is used to measure the diameter of the bubble plume on the surface of the simulant. Since the bubble plume is dynamic and asymmetric, several images at different times are analyzed to determine the ROB. These values are then averaged to represent a single datum. The ZOI was measured by placing buoyant flow followers on the surface near the sparge tube. The flow followers then move radially outward due to the induced secondary flow. The flow followers stop at the point where the radial flow stops and axial flow downward dominates. A measuring tape was used to determine the distance from the center sparge tube to the flow followers.

2.5 Gas Retention and Release

GR&R tests were completed using ≥ 30 Pa yield stress (Bingham plastic model) kaolin:bentonite clay simulant in the UFP and LS prototype vessels in configurations similar to the selected designs. The basis for the simulant selection is presented in Section 4.1. The vessel configurations and nominal operating conditions and the gas retention and release experimental methods are summarized here.

Of the GR&R test sequences run in the prototype vessels, one sequence in LS (Sequence 15) and two in UFP (Sequences 5 and 6) were closely matched to the final designs. The configurations and nominal or target operating conditions used in the gas retention and release tests are as follows:

- LS Sequence 15 – 0.74 H/D initial fill; 8-PJM cluster with 7-45° + 1-vertical nozzles at ~12 m/s; four-nozzle recirculation at ~120 gpm (gas holdup tests only) and four (or eight) air sparge tubes flowing at ~3 acfm each (gas release tests only).
- UFP Sequence 5 – 1.4 H/D initial fill; four-PJM tri-foil configuration with three 45° plus one vertical nozzles at ~12 m/s; single-nozzle recirculation (gas holdup test only) and one center (and three peripheral) sparge tube flowing at ~3 acfm (gas release tests).
- UFP Sequence 6 – 1.8 H/D initial fill; four-PJM tri-foil configuration with three 45° plus one vertical nozzles at ~12 m/s; one center (and three peripheral) sparge tube flowing at ~3 acfm (gas holdup and gas release tests).

Extra sparge tubes (indicated in parentheses above) were used in some tests to release additional gas after the specified release tests with fewer sparge tubes were completed. Unless otherwise specified, the results described in Section 5 were obtained with the nominal operating conditions including fewer sparge tubes.

The decomposition of nominal 30 wt% hydrogen peroxide (H₂O₂) solution was used to generate gas in situ in the kaolin:bentonite clay simulant. The H₂O₂ solution was injected with a peristaltic pump through a single tube into the well-mixed cavern area adjacent to pulse tube nozzles while the PJMs were oper-

(a) Steady state was determined visually by observing that the ZOI flow followers had stopped moving radially from the sparge tube.

ating under prototypic conditions. The amount of H_2O_2 introduced was quantified by weight. In the initial preparation for gas release tests, a specified amount of H_2O_2 was introduced over a short period of time (e.g., 10 to 20 min); after some period of additional mixing (e.g., 10 to 30 min), the system was shut down to allow the H_2O_2 to decompose and gas bubbles to be retained in the quiescent simulant. The accumulated gas was typically released by operating the PJMs and spargers the following day.

In gas holdup tests, H_2O_2 solution was added to simulant at a fixed rate over an extended period of time (e.g., 2 to 3 hours) to continuously generate O_2 gas while the simulant was mixed in the PJM vessel using specified "normal" operating conditions. Injection continued until a new steady-state level was achieved in the test vessel. The rate of H_2O_2 injection was determined by recording the weight of a solution feed container as a function of time. (A MicroMotion flow meter and a data acquisition system were also used to measure and record the solution flow rate.) The mixing system was shut down shortly after the completion of gas holdup tests, resulting in simulant volume growth as residual H_2O_2 decomposed. (Further analysis of the growth profile following shutdown will provide additional information on the apparent gas generation rate.) After a short period of gas retention (30 min or less), a gas release test was typically conducted.

In GR&R experiments, retained gas volume fractions in the prototype vessels were assessed by changes in surface level, which were independently correlated to tank volume. Several methods, including instrumental techniques and visual/camera observations, were used to track changes in surface level over time. Ultrasonic level sensors (Gems Corporation model UCL-200) were deployed in each of the prototype vessels (two in LS and one in UFP), and signals were output to a data acquisition system where they were recorded at 10 Hz. These sensors sample an area of the surface, which increases with the sensor-to-surface separation distance; the sensors were typically placed 0.5 to 1 m from the surface. In both prototypes, the sensors were located in the annular region between the PJMs and the tank wall. Gas volume fractions presented here were determined from the ultrasonic level sensor data. In the case of LS, the results are the average for two sensors located over nearly opposite sides of the tank. Typically, a single volume was determined for each pulse cycle using the minimum level obtained when PJMs were drawn full (suction phase).

A detailed analysis of the uncertainty in reported gas volume fractions and estimated gas generation rates has yet to be completed. The accuracy of reported gas volume fractions is a function of the accuracy of surface level-volume correlations and the accuracy of level measurement techniques. In turn, level measurements are limited by instrument resolution and calibration accuracy and are subject to variability due to surface movement and irregularity during mixing operations. Data fluctuations about central values (e.g., standard deviation in gas holdup tests) give an indication of the variability. These data will be reported in the final DBE strategy.

2.6 Tracer Techniques

Mixing performance in the PJM test vessels was assessed through the use of tracer chemicals. A tracer was injected during the initial stages of the PJM test. Samples of the simulant were taken from several locations during each mixing test to determine the changes in dye concentration as a function of time and operating parameters. At the end of a test cycle, the test vessel was homogenized and a final sample collected. A summary of the technique used is shown in Figure 2.16.

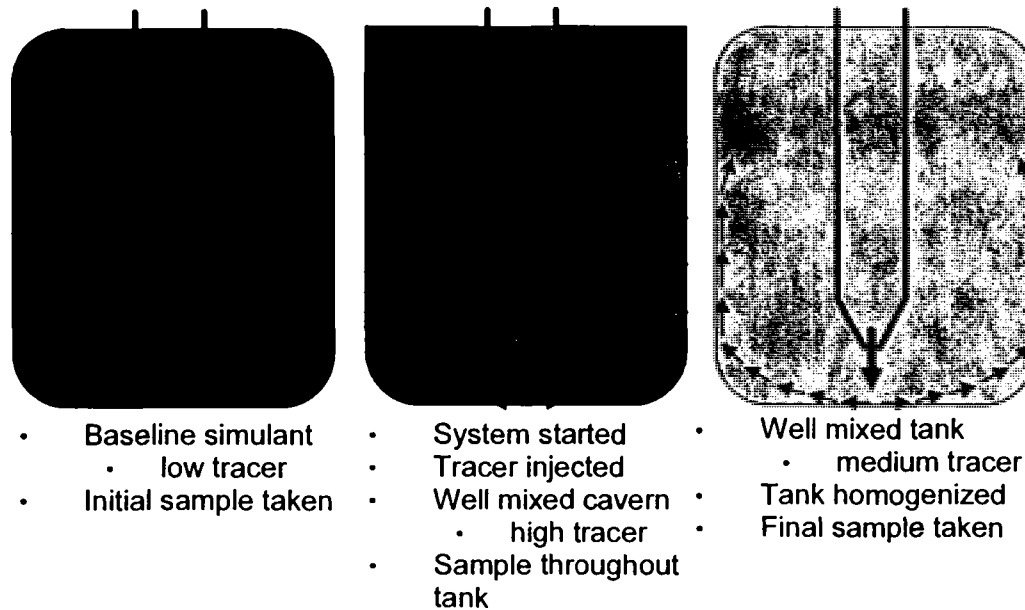


Figure 2.16. Summary of Tracer Dye Technique Steps

The chemicals used were food dye color No. 1, (Brilliant Blue FCF) and sodium chloride (NaCl). Initially, a sample of simulant was drawn from the test vessel to baseline the tracer levels. Next, a stock solution of these materials was prepared by dissolution in water. This stock solution was then blended with a sample of the test simulant to achieve rheological properties close to the actual test simulant. This solution was introduced into the center PJM tube during operation by opening a valve on a sample injection line during the PJM suction phase. During the drive phase, the valve was closed and the injected dye was driven from the PJM tube. Use of this procedure allowed for the gradual introduction of the tracer dye into the system over several drive/suction cycles and minimized the potential for a large amount of concentrated tracer to enter a stagnant region of the tank. This was observed when the concentrated tracer had significantly different physical properties from the bulk simulant. Such physical properties include density, entrained air due to surface tension, and rheological parameters.

After the dye was injected, the experimental clock started and samples were drawn from five locations in each test vessel. Locations 1, 2, and 3 were samples taken directly from three separate pulse tubes. These samples represent the contents of the well-mixed cavern. Sample locations 4 and 5 were placed between the pulse tubes and the tank wall. Location 4 was at a low elevation and location 5 was at a high elevation. Schematic diagrams of the tracer sampling locations are shown in Figures 2.17 and 2.18 for the LS vessel and UFP vessel, respectively.

Multiple run conditions were typically achieved for each tracer injection. The tracer test started with the lowest mixing energy condition to form the initial well mixed cavern. Additional systems (e.g., recirculation pumps or sparging tubes) or increased pulse tube velocities were then used as subsequent run conditions to form larger mixing caverns.

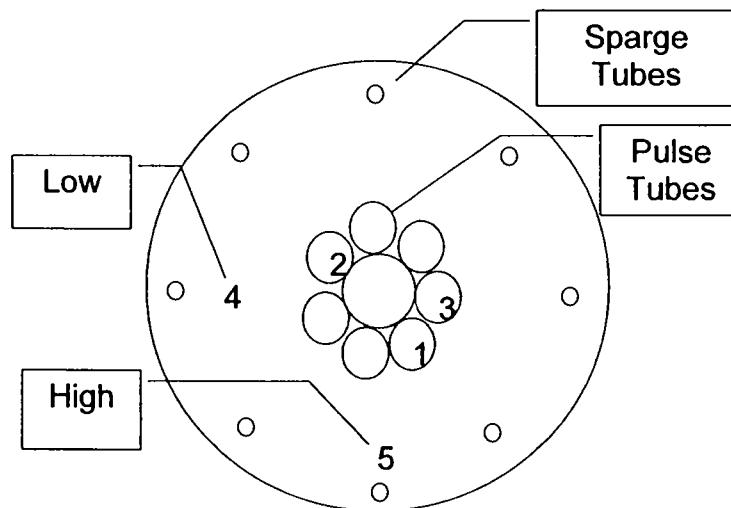


Figure 2.17. Schematic of LS Vessel Tracer Sampling Locations

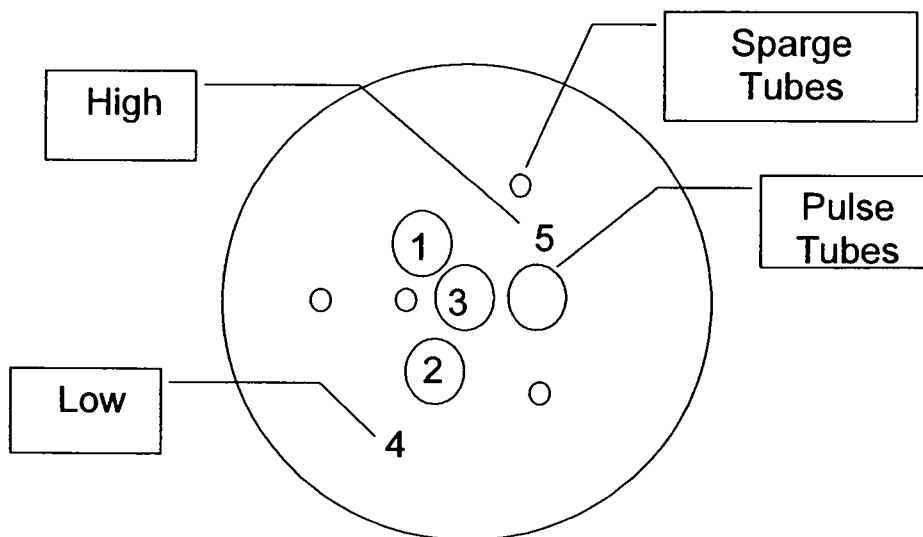


Figure 2.18. Schematic of Ultrafiltration Process Vessel Tracer Sampling Locations

During the initial run condition, samples were drawn from sample locations 1, 4, and 5 taken approximately every 10 minutes after completion of dye injection. After 50 minutes of operation, samples were drawn from all sample locations and the next run experimental condition employed. During subsequent run conditions, samples from locations 1, 4, and 5 were taken every 15 minutes. After 45-90 minutes of operation, samples were drawn from all sample locations, and the next run experimental condition was employed.

This procedure was used to quantify the transient behavior of the mixed regions within the tank. The first run condition was examined in more detail because the anticipated amount of energy required to reach steady state for the first run is greater than subsequent runs where a significant mixed region already exists.

Samples were drawn using a vacuum system. In this system, a vacuum was placed on the sample lines in the tank. The simulant was drawn through the lines and collected in stoppered beakers using a trap. When sampling, the lines were initially purged of simulant into a separate beaker. This step loaded the sample line with simulant from the sample location at the appropriate sample time. A clean beaker was then attached and the newly loaded simulant was collected. The simulant was then transferred into sample containers for tracer analysis. A sample extraction typically took 2 to 5 minutes to complete.

Tracer analysis consists of two measurements, one for the dye and one for the NaCl. The concentration of dye was measured using a UV-VIS spectrometer. This instrument requires a transparent sample. To overcome this limitation, the opaque kaolin:bentonite simulant was centrifuged, and the analysis was performed on the centrifuged liquid portion of the sample. The spectrometer measures the optical absorbance of the sample at multiple wavelengths of light. When the dye is present in the system a peak absorbance is observed at approximately 630 nm. According to Beer's law, the magnitude of this absorbance peak is directly proportional to the concentration of dye in the system.

For the NaCl tracer, a chloride ion selective electrode was used to measure the concentration of chloride present in the samples. This instrument measures the potential difference across an electrode that is surrounded by a membrane that allows chloride ions to pass from the sample material into the electrode cell. Unlike the spectrometer method, this measurement was performed directly on the simulant with no required preparation steps.

The equation used to calculate the fraction mixed is shown below:

$$X_j = \frac{C_f - C_0}{C_j - C_0} \quad (2.4)$$

where

- X_j is the fraction mixed of the j-th tank sample
- C_f is the tracer concentration of the final homogenized simulant
- C_0 is the tracer concentration of the initial baseline simulant
- C_j is the tracer concentration of the j-th tank sample

When the aqueous phase tracer does not absorb onto the solid phase, the liquid phase concentration can be measured with the techniques above, and Equation 2.4 can be used to directly calculate the fraction of the tank mixed. The chloride ion did not appear to absorb onto the simulant particles, and this equation is used for the NaCl tracer. Because the spectrometer measures absorbance, which is proportional to concentration, Equation 2.4 can be rewritten for the dye tracer as follows:

$$X_j = \frac{A_f - A_0}{A_j - A_0} \quad (2.5)$$

where

- X_j is the fraction mixed of the j-th tank sample
- A_f is the optical absorbance of the final homogenized simulant
- A_o is the optical absorbance of the initial baseline simulant
- A_j is the optical absorbance of the j-th tank sample

Unfortunately, the dye tracer absorbs onto the clay particles in significant quantity. In this situation Equation 2.4 still applies, but the concentrations used in the equation must account for both the liquid and solid phases. This is accomplished using the following equation:

$$C = Y_l C_l + Y_s C_s \quad (2.6)$$

where

- C is the tracer concentration
- C_l is the tracer concentration of the liquid phase
- C_s is the tracer concentration of the solid phase
- Y_l is the liquid phase mass fraction
- Y_s is the solid phase mass fraction

The distribution of tracer between the liquid and solid phases is typically described using a distribution coefficient:

$$C_s = K_d C_l \quad (2.7)$$

where K_d is the distribution coefficient.

To complicate matters further, the distribution coefficient is also a function of liquid phase dye concentration. When Equations 2.6 and 2.7 are substituted in Equation 2.4, the following equation results:

$$X_j = \frac{Y_l (A_f - A_o) + Y_s (K_{df} A_f - K_{do} A_o)}{Y_l (A_j - A_o) + Y_s (K_{dj} A_j - K_{do} A_o)} \quad (2.8)$$

where

- K_{df} is the distribution coefficient at the homogenized tank tracer concentration
- K_{do} is the distribution coefficient at the initial baseline tracer concentration
- K_{dj} is the distribution coefficient at the j-th tank sample tracer concentration

When K_d is null or constant, Equation 2.8 reduces to Equation 2.5. Over the small dye concentration ranges observed in the prototype testing, the assumption of a constant distribution coefficient is valid, and Equation 2.5 can be used. Note that as A_j approaches A_f , K_{dj} approaches K_{df} , and the error associated in using Equation 2.5 approaches zero. In addition, the distribution coefficient function varies from batch to batch of simulant, and other factors such as temperature and contact time will also affect the distribution

coefficient function. Lastly, the solids loading of the simulant was often varied for rheological purposes. For these reasons, Equation 2.5 is used to estimate the fraction mixed using the dye tracer. The error associated with this assumption is predicted using estimated values for the liquid and solid mass fractions and the distribution coefficient. Appendix B contains further details on these parameters.

3.0 Supporting Data for Mixing System Vessel Configurations

3.1 Rheology

3.1.1 Bounding Conditions

For all seven WTP vessels that will contain non-Newtonian fluids, it was assumed that the HLW pretreated sludge bounding physical and rheological properties would hold (CCNs 069099, 065607, and 082255).

Normal Plant Operation Rheological Bound: Data from actual radioactive and simulant waste rheograms combined with general engineering design techniques were used to define a set of bounding physical and rheological properties that agree well with actual data (POLOSKI 2004). The non-Newtonian HLW pretreated sludge rheological properties were fit using a linear Bingham plastic model. The bounding conditions were used to develop the waste simulants used in the PJM program. Figure 3.1 is a plot of actual pretreated waste rheograms and the upper bounding rheological properties curve. The linear Bingham plastic model fit parameters are yield stress (y-axis intercept) of 30 Pa and consistency (slope) of 30 cP. Table 3.1 contains a summary of expected physical and rheological properties.

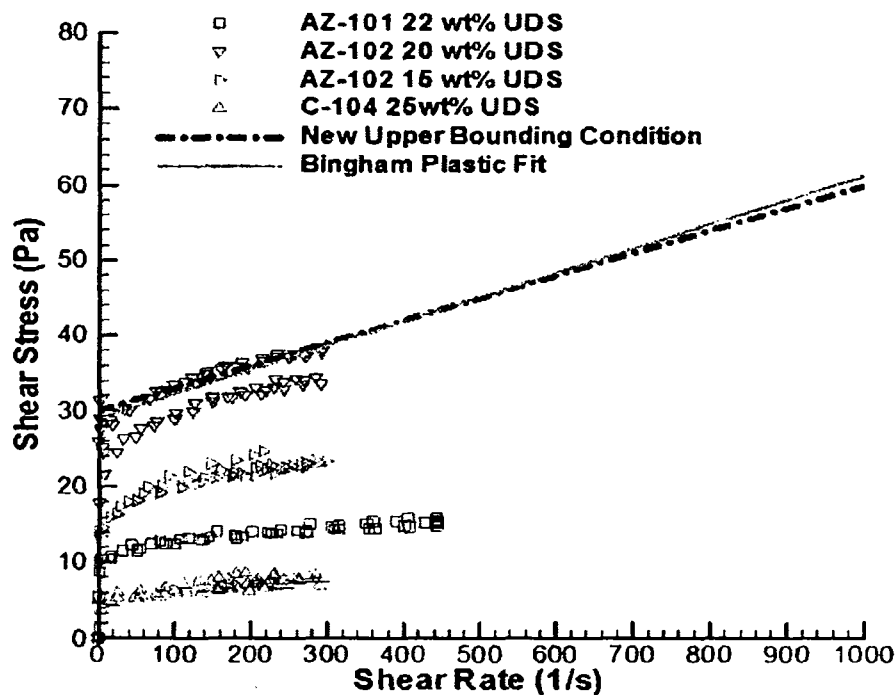


Figure 3.1. Rheogram of Actual HLW Pretreated Sludge Samples with Upper Bound Rheological Curve

Table 3.1. Physical and Rheological Properties that Help Define Simulants for Rating or Qualifying Fluidic Mixing Systems

Property	HLW Pretreated Sludge
pH	$\approx 12^{(a)}$ -14
Particle size distribution (D_{50}) ^(b)	2 μm
Particle size distribution (D_{95}) ^(c)	20 μm
Bulk density	1.1-1.6
Supernatant liquid density	≈ 1.0
Vol% settled solids	10%-90%
Wt% total dried solids	5%-25%
Wt% total oxide	7%-15% ^(d)
Shear stress versus shear rate (ambient and 40°C)	Bingham Plastic
(a) Expected pH after washing leaching in 0.01 M NaOH.	
(b) 50% of particles are smaller than the indicated value.	
(c) 95% of particles are smaller than the indicated value.	
(d) Based on simulant data.	

Because the rheological window is based on only four samples from three tanks, it is possible that slurries from other tanks could exceed the rheological boundary. It has been estimated that 20 to 30% of HLW tanks may have rheological properties higher (yield stress and consistency higher than 30 Pa and 30 cP, respectively) than those documented in the three active tank samples analyzed to date (CCN 082255). This uncertainty will be addressed by laboratory testing prior to receipt of the waste at the WTP to define the extent to which the slurry may be concentrated and stay below the rheological boundary.

Plant Upset Operation Rheological Bound: It is important to note that measured maximum shear strength values (an actual physical property that must be overcome in order for these fluids to flow) for actual HLW pretreated sludge samples when allowed to stand in an unmixed condition, that is, post-DBE, they reach values greater than 30 Pa. For this reason, a bounding yield strength value of 70 Pa should be used (CCN 065607). In addition, the "gel" time (the time required for the actual waste to reach its maximum shear strength value) of actual waste samples will need to be taken into account along with the maximum shear strength values for plant operation considerations.

3.1.2 Simulants

One transparent simulant and one opaque simulant were used in the PJM program. The transparent simulant was Laponite RD (Southwestern Clay Products), a thixotropic colloidal synthetic clay that forms stable gel networks when unsheared. Due to the thixotropic nature of Laponite, the flow behavior of the simulant is dynamic, and it was allowed to gel and reach a target shear strength. Speers et al. (1987) demonstrated that the shear strength of clay drilling muds increases over time following first-order rate kinetics. Laponite shear strength behavior was observed to agree with the Speers et al. (1987) correlation for drilling muds. At this point the PJM system was started and a mixing cavern formed as defined by the gel's shear strength. After constant shearing, a steady-state flow behavior was approached. Unfortunately, this flow behavior was lower than the bounding rheology of WTP waste streams. This is illustrated in Figure 3.2, where actual HLW pretreated sludge rheograms are compared with PJM simulants. The bounding rheological parameters of the HLW pretreated sludge (Poloski 2004) are defined as Bingham plastic consistency of 30 cP and yield stress of 30 Pa.

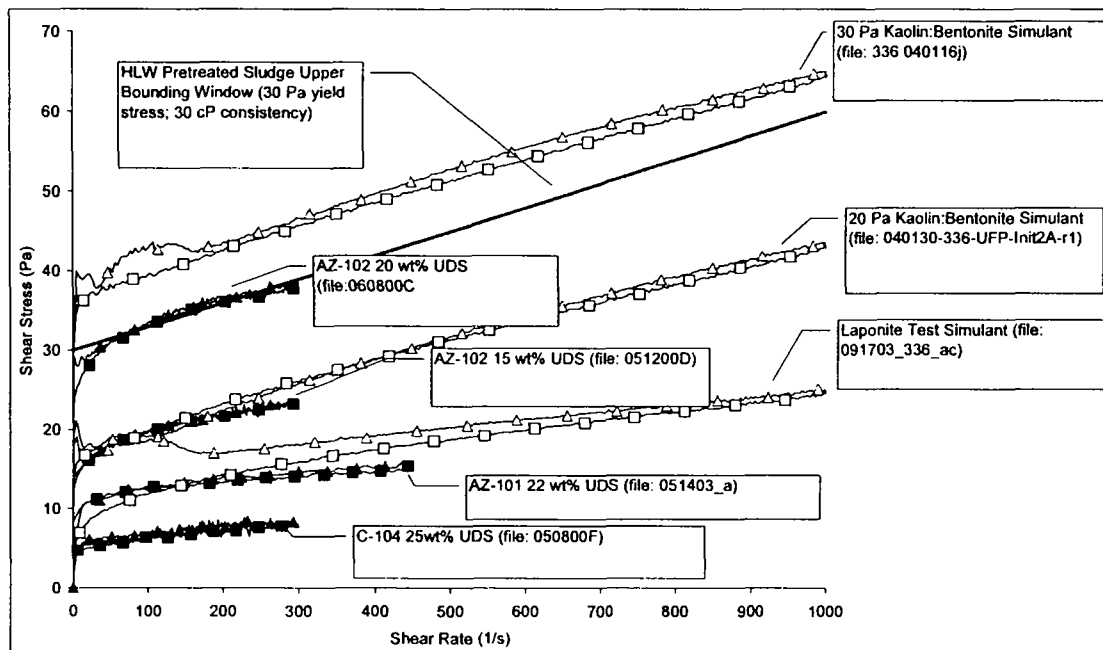


Figure 3.2. Flow Behavior Comparison of PJM Simulants and Actual HLW Pretreated Sludge

In addition to not possessing the target rheological parameters desired for PJM testing, the Laponite composition also does not match other target values given in Table 3.1. The Laponite recipe calls for 1-2 wt% Laponite RD in water where the actual waste is in the 15 to 25 wt% undissolved solids range. And the Laponite simulant consists of particles on the order of tens of nanometers, whereas the actual waste consists of particles in the tens of microns range. These differences may result in varying turbulent flow behavior in the PJM mixing cavern. For these reasons, a more representative particulate slurry was developed to enhance confidence in the PJM testing results. Unfortunately, this simulant is opaque.

The particulate simulant developed consists of a mixture of kaolin clay (EPK Feldspar Pulverized) and bentonite clay (WYO-Ben Big Horn CH-200) in water. To meet the WTP bounding parameters of Bingham plastic consistency of 30 cP and yield stress 30 Pa, a recipe was developed using these two clays. The recipe calls for a composite of 80% kaolin and 20% bentonite mixed with water to a loading of approximately 27 wt%. Water is then added to the simulant to adjust the rheological parameters to other target values. Figure 3.2 compares these simulants with actual waste at various solids loadings to target 30+ and 20 Pa yield stress. A summary of the measured rheological parameters for significant prototype tests and sparging tests is shown in Tables 3.2, 3.3, and 3.4. In addition, the bentonite/kaolin simulant shear strength behavior was observed to agree with Speers et al. (1987) correlation for drilling muds.

Table 3.2. Rheological Model Fits for LS Prototype PJM Simulants at Ambient Temperature

Model/Model Parameter	LS Test 4	LS Test 7	LS Test 11	LS Test 15	LS Test 20
File Name	336 040116j	040123-336- ls-T7-R1	336 04129b	040216 apel- 0002f	040309 apel- r1-0001f
Bingham Plastic:					
τ_O^B - Bingham yield stress (Pa)	37	36	37	36	36
k - Bingham consistency coefficient (cP)	28	27	26	27	24
R - correlation coefficient	1.0	1.0	1.0	9.6 ^(a)	5.8
Herschel-Bulkley:					
τ_O^H - yield stress (Pa)	35	34	35	34	34
k - Herschel-Bulkley consistency coeff. (Pa·s ^{-b})	0.092	0.090	0.082	0.12	0.063
b - Herschel-Bulkley power law exponent	0.83	0.83	0.84	0.79	0.86
R - correlation coefficient	1.0	1.0	1.0	1.3 ^(a)	1.6 ^(a)
(a) Standard error.					

Table 3.3. Rheological Model Fits for UFP Prototype PJM Simulants at Ambient Temperature

Model/Model Parameter	UFP Test 2	UFP Test 3B	UFP Test 5	UFP Test 6
File Name	040130-336- ufp-T2-init-r1	040213 apel- 0003f	040213 apel- 0006f	040213 apel- 0012f
Bingham Plastic:				
τ_O^B - Bingham yield stress (Pa)	34	33	36	37
k c - Bingham consistency coefficient (cP)	27	18	19	20
Rc - correlation coefficient	1.0	6.4 ^(a)	4.8 ^(a)	5.2 ^(a)
Herschel-Bulkley:				
τ_O^H - yield stress (Pa)	33	32	35	36
k - Herschel-Bulkley consistency coefficient (Pa·s ^{-b})	0.086	0.059	0.046	0.053
b - Herschel-Bulkley power law exponent	0.84	0.83	0.88	0.87
R - correlation coefficient	1.0	2.1 ^(a)	1.8 ^(a)	1.7 ^(a)
(a) Standard error.				

Table 3.4. Rheological Model Fits for LS Prototype PJM Simulants at Ambient Temperature

Model/Model Parameter	336 simulant 2-20-04	336 simulant after peroxide addition	336 simulant during gas retention test	336 simulant after gas retention test
File Name	040224 apel- 0007f	040227 apel- 0002f	040301 apel- 0009f	040308 apel- 0001f
Bingham Plastic:				
τ_O^B - Bingham yield stress (Pa)	37	33	33	32
k - Bingham consistency coefficient (cP)	22	21	22	23
R - correlation coefficient	7.7 ^(a)	7.8 ^(a)	7.2 ^(a)	9.1 ^(a)
Herschel-Bulkley:				
τ_O^H - yield stress (Pa)	35	32	32	30
k - Herschel-Bulkley consistency coefficient (Pa·s ^{-b})	0.088	0.082	0.073	0.098
b - Herschel-Bulkley power law exponent	0.81	0.81	0.83	0.80
R - correlation coefficient	1.9 ^(a)	1.8 ^(a)	1.7 ^(a)	2.1 ^(a)
(a) Standard error.				

3.2 Prototype Results

The various mixing tests performed and the percent mixed for both the UFP and LS prototype test stands are summarized in Tables 3.5 and 3.6, respectively. The PJMs, spargers, and recirculation pump configurations for the various sequences listed in the tables are presented in Tables 2.1 and 2.2 (Section 2.3). All tests were performed with a kaolin/bentonite clay simulant, the yield stress of which was determined from thoroughly mixed samples (mixed by PJM overblow and sparging) collected prior to and at the completion of a sequence of runs. The yield stress of the kaolin/bentonite clay simulant is the average of the results for these samples. The H/D_T is the ratio of the simulant fill height to tank diameter.

The nozzle velocities listed in Tables 3.5 and 3.6 were calculated based on the average velocity (\bar{u}_{disch}) as defined by Equation A.7 in Appendix A. The average nozzle velocities are based on averages of all the PJMs (four or six for UFP and eight for LS) taken over typically 25 representative cycles of PJM operation during a run. The cycle times listed in Tables 3.5 and 3.6 for the two test stands were set based on scaling approximately equal to the inverse of the geometric scale factor, that is, 4.94 and 4.24 for the UFP and LS prototype test stands, respectively.

For tests that used a recirculation pump, the pump flow rates were scaled approximately by the square of the geometric scale factor, that is, 4.94^2 (=24.4) and 4.29^2 (18.0) for the UFP and LS prototype test stands, respectively. The recirculation flow rates listed in Tables 3.5 and 3.6 are based on the average of the flow rates measured over the duration of a run. In calculating recirculation pump averages, startup transients were ignored.

For tests that involved sparging, no scaling was applied in setting the operating air flow rates and acfm through the sparger tubes are based on the readout of the rotameters included in-line with each sparger. No corrections were applied to the sparger air flow rates, and a post-calibration of the flow meters indicated the sparger flow rates were within $\pm 15\%$.

Table 3.5. Test Conditions and Fraction Mixed Results for Tests Performed in UFP Test Stand

Seq	Run	Test Mode	H/D	Yield Stress (Pa)	Noz. Vel. (m/s)	Cycle Time (sec)	Sparger Flow Rate (acfm)	Pump Flow Rate (gpm)	Fraction Mixed	Error ^(a) (\pm)
2	1	PJM Only	1.8	35	9.0	27	-	-	0.53	0.093
2	2	PJM Only	1.8	35	12.3	27	-	-	0.64	0.074
2	3	PJM + Sparging	1.8	35	12.3	27	3	-	1.1	0.013
2	4	PJM + Sparging	1.8	35	12.4	27	1	-	0.96	0.0088
3B	1	PJMs Only	1.4	37	9.3	27	-	-	0.65	0.12
3B	2	PJM + Pump	1.4	37	9.3	27	-	90	0.98	0.0074
3B	3	PJM + Pump	1.4	37	14.1	27	-	87	1.0	0.0019
3B	4	PJM + Pump + Sparging	1.4	37	14.1	27	3	95	1.0	0.0038

(a) Estimated error due to assumption of linear isotherm for dye absorption. Experimental error not included.

Table 3.6. Test Conditions and Fraction Mixed Results for Tests Performed in LS Test Stand

Seq	Run	Test Mode	H/D	Yield Stress (Pa)	Noz. Vel. (m/s)	Cycle Time (sec)	Sparger Flow Rate (cfm)	Pump Flow Rate (gpm)	Fraction Mixed	Error ^(a) (±)
4	1	PJMs Only	0.74	38	7.8	45	-	-	0.54	0.15
4	2	PJMs Only	0.74	38	11.3	45	-	-	0.65	0.13
4	3	PJMs + Spargers	0.74	38	11.1	45	3	-	0.87	0.052
4	4	PJMs + Spargers	0.74	38	11.4	45	3	-	0.97	0.014
7	1	PJMs Only	1	36	4.6	55	-	-	0.24	0.11
7	2	PJMs Only	1	36	7	45	-	-	0.42	0.085
7	3	PJMs + Pump	1	36	7	45	-	121	0.55	0.06
7	4	PJMs + Pump	1	36	10.3	45	-	119	1.1	0.01
7	5	PJMs + Pump + Sparging	1	36	10.4	45	3	122	1.1	0.0058
7	6	PJMs + Pump + Sparging	1	36	10.5	45	3	121	0.93	0.0067
11	1	PJMs + Pump	0.74	37	8.2	45	-	121	0.66	0.033
11	2	PJMs + Pump	0.74	37	11.9	45	-	115	0.95	0.0055
20	1	PJMs	0.74	35	12.3	45	-	121	0.96	0.0097
20	2	PJMs + Pump	0.74	35	12.2	45	-	122	1.0	0.00069

(a) Estimated error due to assumption of linear isotherm for dye absorption. Experimental error not included.

The fraction mixed data presented in Tables 3.5 and 3.6 are based on the measurements obtained from the dye/tracer injected into the simulant prior to the start of a test sequence, and the approach is discussed in Section 2.6 and Appendix B. The error in the fraction mixed values is due to a linear isotherm assumption for dye absorption (see Appendix B for details). This error goes to zero as the fraction mixed goes to 100%. Experimental variability due to sampling and analysis is still present. The percent mixed versus yield Reynolds number for the various tests conducted with the UFP prototype test stand are shown in Figure 3.3. Similar results for the LS prototype test stand are shown in Figure 3.4.

It can be seen from the data in Figure 3.3 that with PJMs only an increase in the yield Reynolds number results in an increase in the percent mixed. It can also be in Figure 3.3 that PJMs alone are not sufficient to completely mix the tank. The addition of sparging and/or recirculation generally results in complete mixing. Similar observations can be made for the LS prototype test stand.

The test conditions and results of the various solids lift tests performed in both UFP and LS prototype test stands are shown in Tables 3.7 and 3.8. The PJM configurations for UFP and LS prototype test stands are presented in Tables 2.1 and 2.2 and discussed in Section 2.3. For all the solids lift tests, a slurry of 4-mm glass beads (specific gravity 2.5) in water was used. The concentration of the glass beads was ~0.2 vol%. In Tables 3.7 and 3.8, the nozzle velocities were determined based on the averages described above.

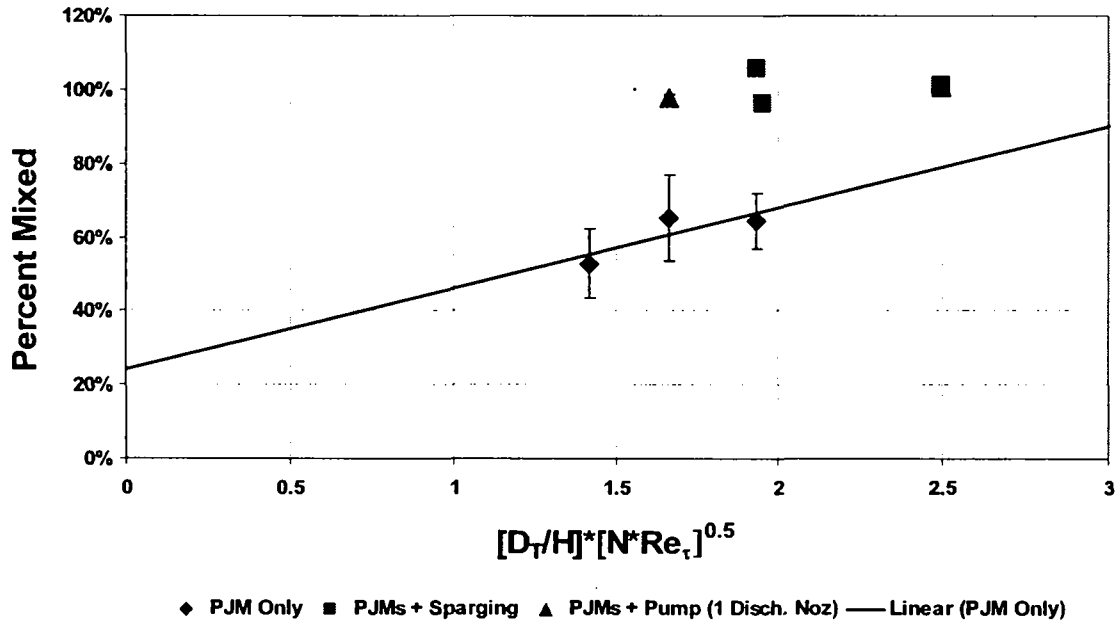


Figure 3.3. Percent Mixed Versus Yield Reynolds Number for UFP Prototype Test Stand During Various Operating Conditions

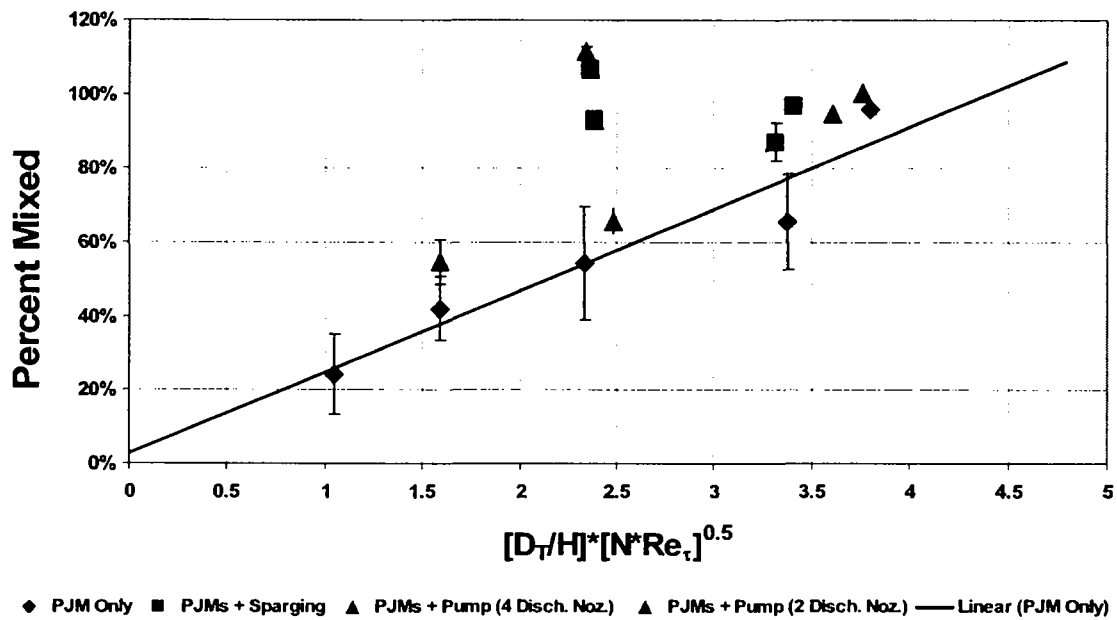


Figure 3.4. Percent Mixed Versus Yield Reynolds Number for LS Prototype Test Stand During Various Operating Conditions

Table 3.7. Test Conditions and Results of Solids Lift Tests Performed in UFP Prototype Test Stand

Seq	Run	Test Mode	H/D	Noz. Vel. (m/s)	Cycle Time (sec)	Solids Lift (Yes/No)
7	1	PJMs only	1.8	4.7	27	No
7	2	PJMs only	1.8	6.1	27	No
7	3	PJMs only	1.8	6.5	27	Yes
7	4	PJMs only	1.8	6.9	27	Yes

Table 3.8. Test Conditions and Results of Solids Lift Tests Performed in LS Prototype Test Stand

Seq. No.	Run	Test Mode	H/D	Noz. Vel. (m/s)	Cycle Time (sec)	Solids Lift (Yes/No)
16	1	PJMs only	0.74	7.8	45	No
16	2	PJMs only	0.74	8.6	45	Yes
16	3	PJMs only	0.74	9.2	45	Yes
16	4	PJMs only	0.74	7.0	45	No
16	5	PJMs only	0.74	7.6	45	No
16	6	PJMs only	0.74	8.0	45	Yes

During the solids lift tests, visual observations were made to assess whether at any moment during the drive phase all the solid glass beads were lifted off the floor. These observations are indicated by Y or N (yes or no) in the last column of Tables 3.7 and 3.8.

The data for the bead lift tests in the UFP test stand using the "cluster" configuration and 45° nozzles indicates that the minimum velocity needed to lift the beads from the floor was between 6.1 and 6.5 m/s. For the LS prototype with cluster PJM configuration and 45° nozzles, the minimum velocity to lift the beads from the floor was found to be between 7.8 and 8 m/s. All the values are below the minimum jet velocity of 8 m/s being considered for the PJMs.

These velocity values can be extended to other concentrations and particle sizes through the functionalities given in Section 2.3.5 and Equation 2.2. FMP found that the effect of scale was given by the function V_{js} proportional to $T^{0.3}$ for constant ratio of tank to jet diameter. This scale-up effect is small because of the large scale of the test tanks.

The above data only refers to whether solids are lifted off the tank bottom. How well they are distributed vertically in the tanks depends on different factors with different functionalities. In WTP this has been studied with computational fluid dynamics (CFD) and has shown with slow settling particles that the solids are fairly well distributed. However CFD cannot currently determine whether the solids are lifted off the bottom; this requires the experimental verification discussed above.

3.3 Sparging

The mobilization performance of a single sparge tube in a rheologically bounding WTP simulant was investigated as described in Section 2.4. The rheological properties of the simulant are described in

Section 3.1.2. ZOI and ROB diameter results were plotted against the actual volumetric air flow in the slurry at the end of the submerged sparge tube. These results are shown in Figure 3.5. Examination of the data reveals that the ROB and ZOI diameters are a weak function of submergence depth. This indicates that these regions have a nearly cylindrical submerged vertical profile. The full-scale PJM sparger systems will be submerged deeper than measured in these experiments. Because the ZOI and ROB diameters will increase slightly with submergence depth, this assumption is conservative from a design perspective. A correlation of the ROB and ZOI diameters to actual volumetric flow rate adequately describes the size of these regions (see Equations 3.1 and 3.2).

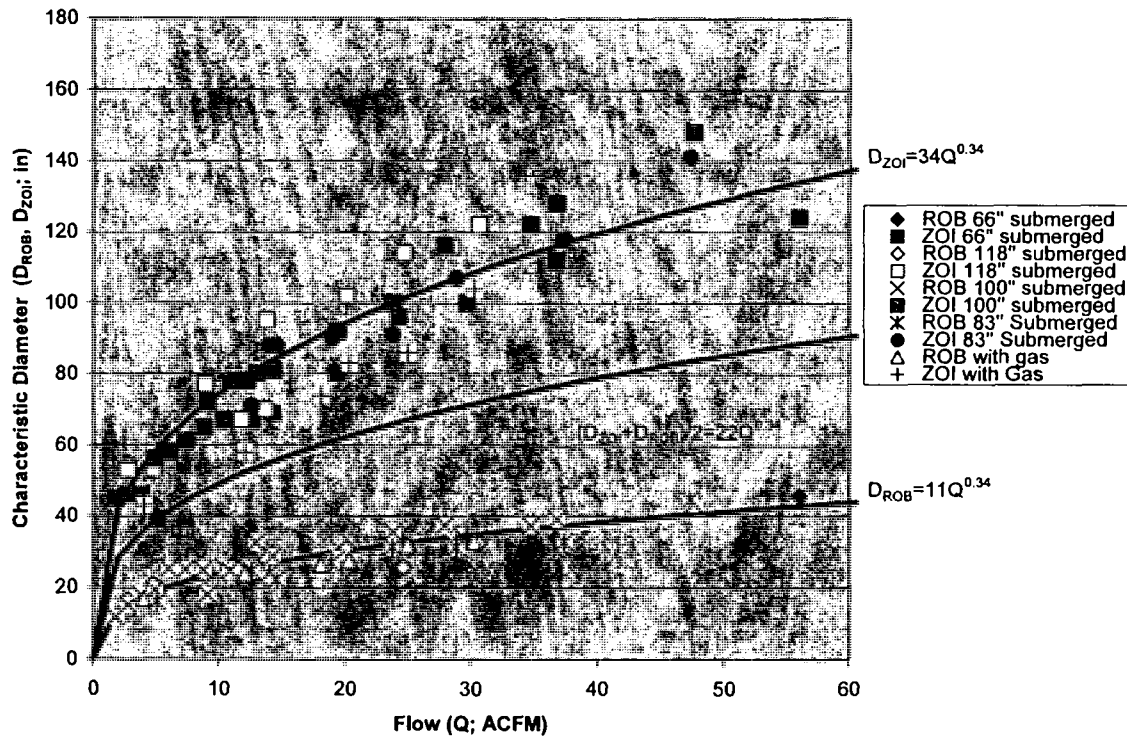


Figure 3.5. ZOI and ROB Sparger Diameters at Various Air Flow Rates

$$D_{ROB} = 11Q_{air}^{0.34} \tag{3.1}$$

$$D_{ZOI} = 34Q_{air}^{0.34} \tag{3.2}$$

where

- D_{ROB} is the ROB diameter (in)
- D_{ZOI} is the ZOI diameter (in)
- Q_{air} is the actual volumetric flow rate of the air in slurry at the end of the sparge tube (ft³/min)

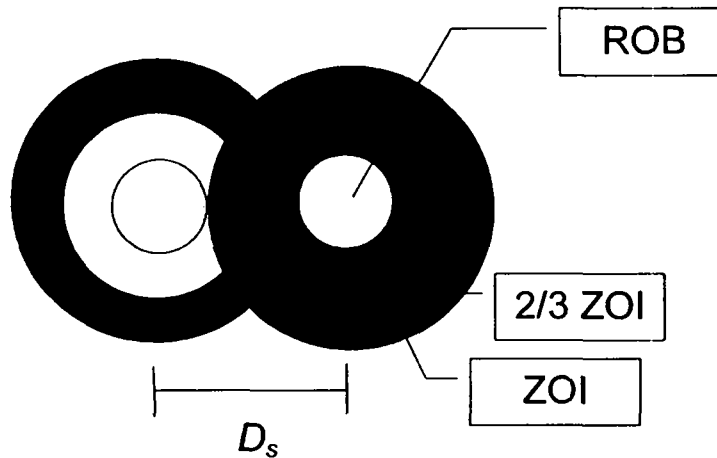


Figure 3.6. Adjacent ZOI and ROB Interaction Options

From these correlations one can see that $D_{ZOI} \approx 3 \cdot D_{ROB}$. If one designs a sparging system such that the ZOI from one sparger tube meets the ROB from an adjacent sparger (Figure 3.6), the sparger spacing shown in Equation 3.3 can be specified.

$$\begin{aligned}
 D_s &= \frac{D_{ROB} + D_{ZOI}}{2} \\
 D_{ZOI} &= 3D_{ROB} \\
 D_s &= \frac{2}{3}D_{ZOI}
 \end{aligned}
 \tag{3.3}$$

where

D_s is the sparger spacing
 D_{ROB} is the ROB diameter
 D_{ZOI} is the ZOI diameter

Another single-tube sparging test was performed by mixing hydrogen peroxide with the simulant. The simulant was then allowed to sit undisturbed for a time as the decomposition of hydrogen peroxide proceeded to load the simulant with gas. The sparging experiment was then performed on simulant loaded with gas that had developed a shear strength due to remaining undisturbed for several hours. These results are shown in Figure 3.5 and indicate a significant decrease in ZOI diameter during these tests.

The ROB diameter appears unaffected by the presence of gas. Potential factors that influence the measured ZOI diameter were the presence of gas in the system at startup and increasing shear strength due to gelation of the slurry as it sat undisturbed. Nonetheless, this test illustrates that actual sparger performance in the WTP will be affected by letting the waste remain undisturbed for periods of time, allowing for increasing rheological parameters and gas holdup. Startup procedures to recover from these scenarios should be considered to ensure successful operation of the WTP.

4.0 Description of Selected Pretreatment Facility Designs

The PJM cluster configuration concept, that is, one central pulse tube with the remaining pulse tubes clustered around the central tube, was chosen for both the UFP and LS vessels. This configuration provides a mixed turbulent cavern in the bottom of the vessel that suspends waste particles and is scalable. Supplemental mixing used to mix the upper portion of the vessels relies on recirculation pumps or spargers. This section describes that process.

4.1 Ultrafiltration Feed Process Vessel (UFP-VSL-00002A/B)

Normal operation (without leaching): Under normal operation (without leaching), the UFP operates at or below an aspect ratio of 1.4. The aspect ratio (H/D) is defined by the liquid height (H) divided by the vessel diameter (D). A combination of the PJMs and a recirculation jet will provide adequate mixing up to an $H/D=1.4$. The recirculation pump must provide 2200 gpm to the jet. The jet is sized to provide an exit velocity of 30 ft/sec. Each PJM provides at least a 12 m/s flow during the PJM drive cycle.

During this mode of operation, the sparge tubes will be run in an 'idling' air flow mode using 0.5 to 1.0 acfm (~2 scfm) of air per sparge point. The layout of PJMs, sparge lines, and recirculation lines is shown in Figures 4.1 and 4.2.

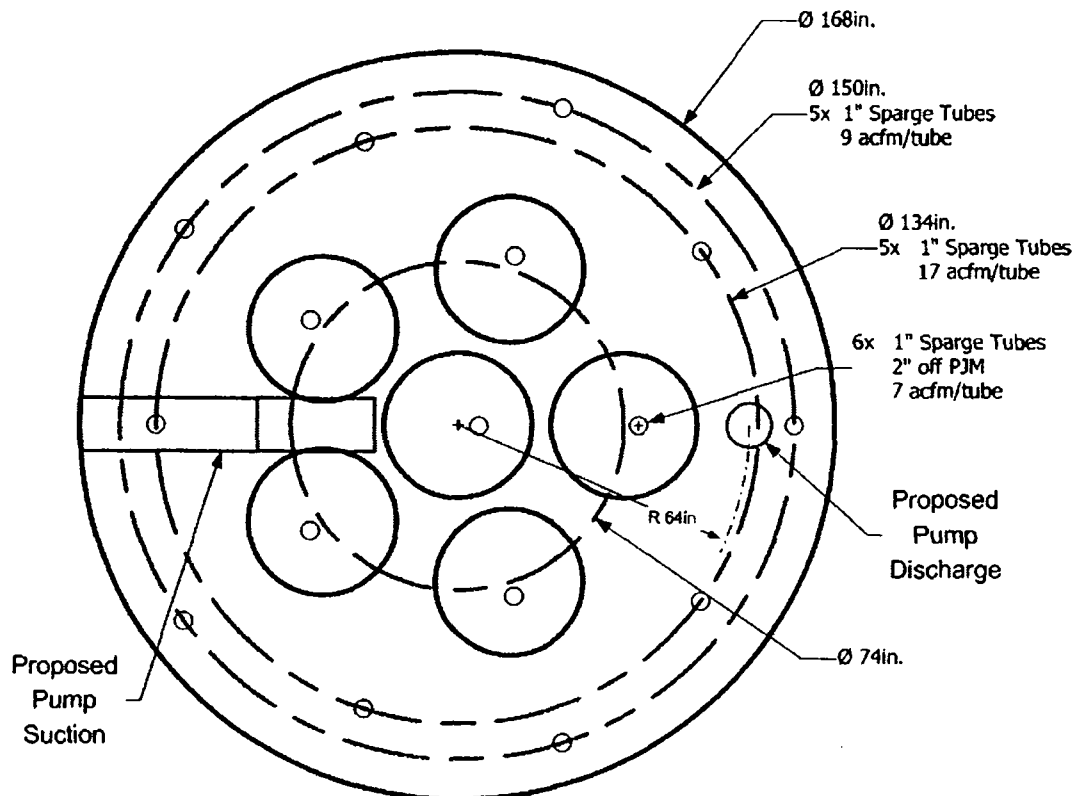


Figure 4.1. UFP-VSL-00002A/B Mixing System Layout – Plan View

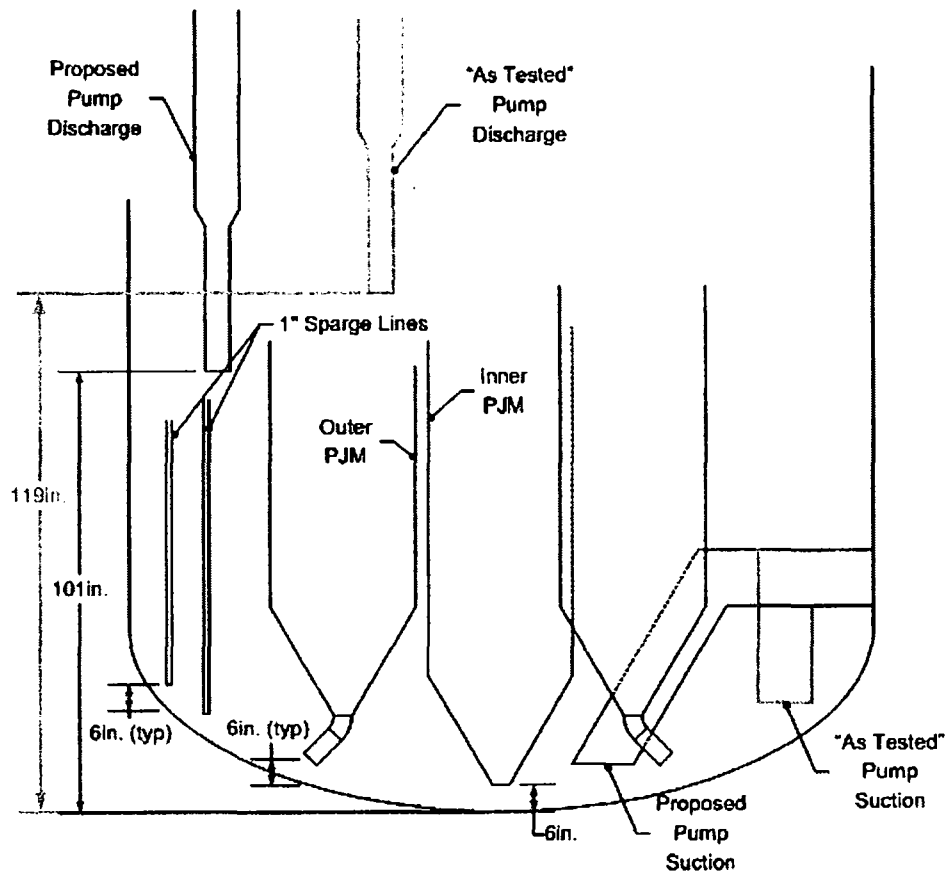
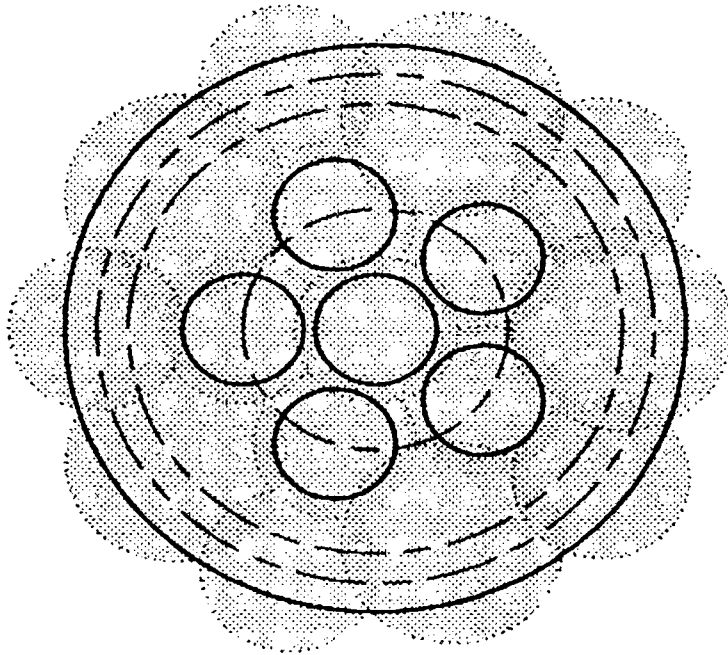


Figure 4.2. UFP-VSL-00002A/B Mixing System Layout – Elevation View

Normal operation (with leaching): When leaching is required, the liquid H/D is above 1.4, the pump is off, and the air sparge system must be used to provide mixing. Varieties of sparge tube layouts were considered for use. The configuration presented in this document was chosen by WTP Engineering to minimize impact with consideration to total air requirement and total number of sparge lines. The selected configuration is shown in Figure 4.3. Hatched circles overlaid in the plan view indicate the size of the ZOI of each particular sparge tube at the flow rates specified in the adjacent table.

The bubble size and resulting mixing zone is based on the air flow (in acfm) at the sparge line exit. The required flow rate measured in scfm is based on the level and density of the liquid in the vessel. The scfm values in Figure 4.3 are based on the overflow level in UFP-VS-00002A/B and a slurry specific gravity of 1.35.

Post-DBE/High Levels: The UFP vessel under post-DBE conditions, or H/D greater than 1.4, requires operation of PJMs and full sparging.



16 Sparge Tubes	# OF TUBES	2/3 ZOI (in.)	flow/tube (acfm)	subtotal (acfm)	Radial Position (in)	Nozzle Elevation (in)	Pressure Corr. Factor	subtotal (scfm)	UFP-VSL-00002A/B
	5	60	17	87	67	23	2.23	195	
	5	48	9	45	75	27	2.22	101	
	6	44	7	42	37	219	1.58	67	
Total Air Flow (scfm):								362	

Figure 4.3. Sparge Air Requirements and Resulting ZOIs – UFP-VSL-00002A/B

PJM Details: The central PJM has a downward-pointing 4-inch nozzle that is 1.5 nozzle diameters (6 inches) off the bottom of the vessel. The outer five PJMs are located on a pitch circle diameter (PCD) of 74 inches. All of their exit nozzles are pointed outward toward the vessel sidewall at 45 degrees; the nozzle openings are also 4 inches and are 6 inches off the bottom of the vessel. Elevation views of the pulse tubes are shown in Figures 4.2 and 4.4 with the outer pulse tube nozzle detail shown in Figure 4.5.

Sparge Line Details: The sparge lines are all 1 inch with the outer two PCD spargers, i.e., 134- and 150-inch PCD, all approximately 6 inches off the bottom of the vessel. Each of the spargers on a single PCD is spaced so that the angle between them is the same; e.g., for a PCD set of spargers equal to 5, the spargers are spaced every 72 degrees. None of the exit tube locations are near pulse tube nozzles or the recirculation pump intake. The spargers above the pulse tubes are approximately 2 inches above the top. Elevation views of the spargers are shown in Figures 4.2 and 4.4.

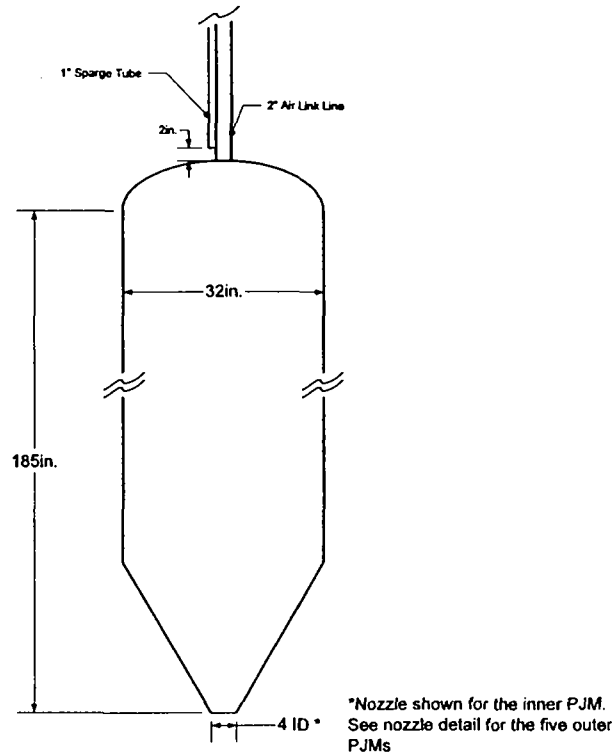


Figure 4.4. PJM Details – UFP-VSL-00002A/B

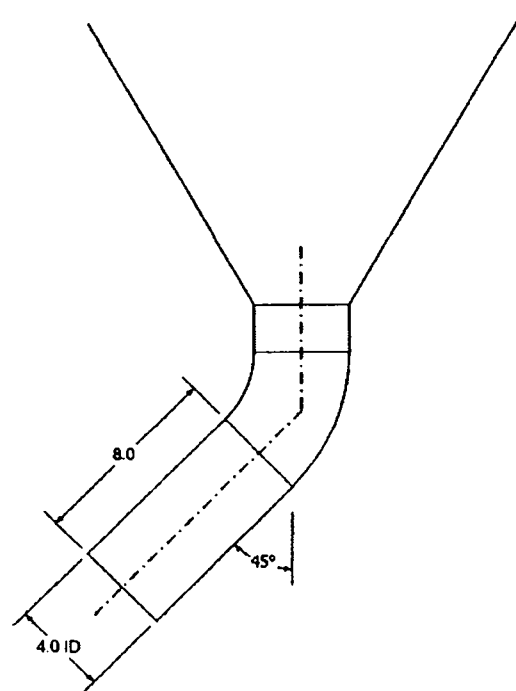


Figure 4.5. Outer PJM Nozzle Detail – UFP-VSL-00002A/B

4.1.1 HLW Lag Storage Vessel (HLP-VSL-00027A/B)

Normal Operation: The HLW LS vessel under normal operation will require eight PJMs and a 2200-gpm recirculation pump with one intake and two terminating nozzles configured to provide an exit velocity of 40 ft/sec to supply the required mixing. PJM nozzle velocities must be at least 12 m/s during the PJM drive cycle. Under normal operation the sparge tubes will be run in an 'idling' air flow-only mode, using 0.5–1.0 acfm (~2 scfm) of air per sparge point. Mixing with PJMs and recirculation jets has been tested up to an H/D of 0.74. The layout of PJMs, sparge lines, and recirculation lines is shown in Figures 4.6, 4.7, and 4.8.

Post-DBE/High Levels: The LS vessel under post-DBE conditions, or an H/D greater than 0.74 requires operation of PJMs and full sparging. Varieties of sparge tube layouts were considered for use. The configuration presented in this document was chosen by WTP engineering to minimize impact with consideration to total air requirement and total number of sparge lines. The selected configuration is shown in Figure 4.9. Hatched circles overlaid in the plan view indicate the size of the ZOI of each particular sparge tube at the flow rates specified in the adjacent table.

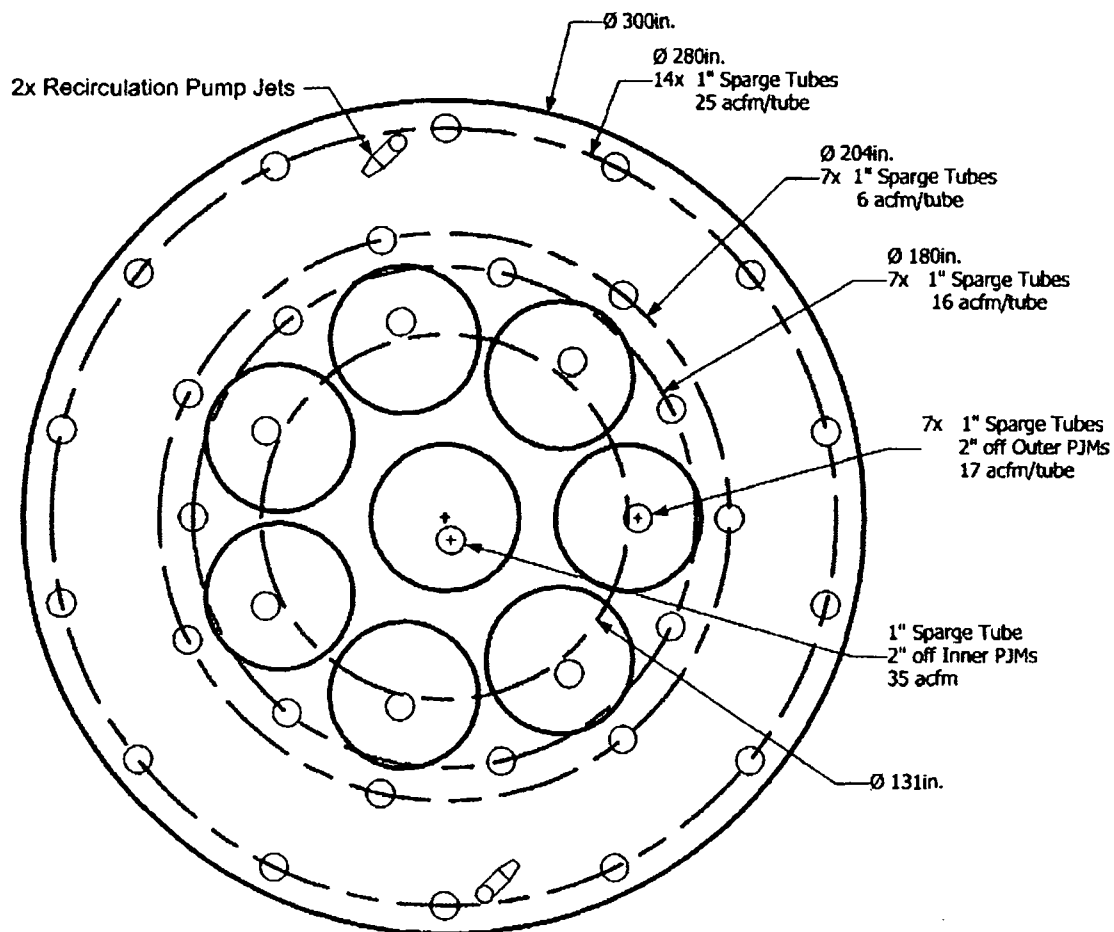


Figure 4.6. HLP-VSL-00027A/B Mixing System Layout – Plan View

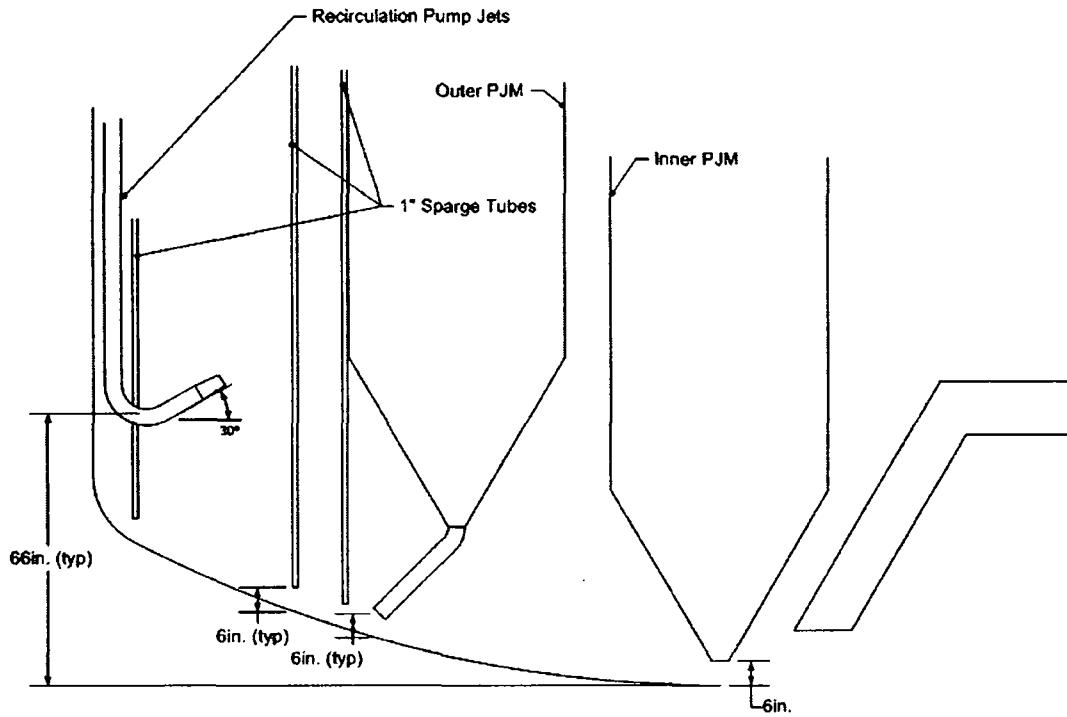


Figure 4.7. HLP-VSL-00027A/B Mixing System Layout – Elevation View

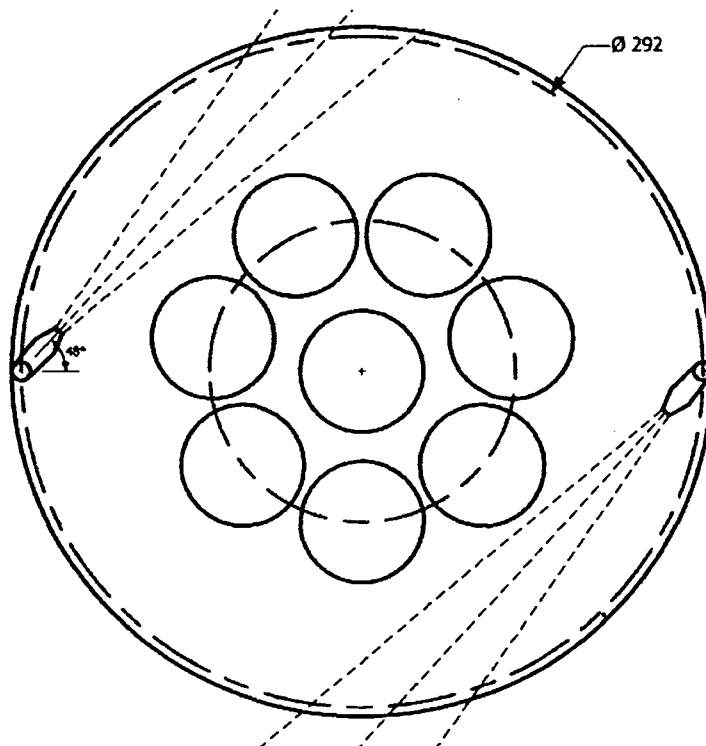
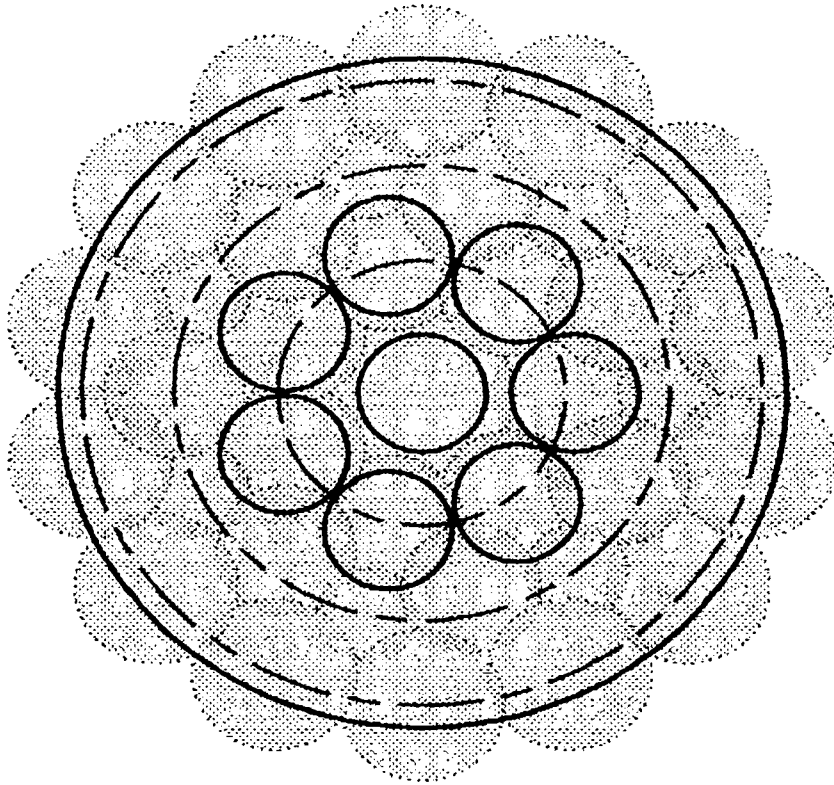


Figure 4.8. HLP-VSL-00027A/B, HLP-VSL-00028 Recirculation Jet Layout – Plan View



36 Sparge Tubes	# OF TUBES	2/3 ZOI (in.)	flow/tube (acfm)	subtotal (acfm)	Radial Position (in)	Nozzle Elevation (in)	Pressure Corr. Factor	subtotal (scfm)	HLP-VSL-00027A/B
	14	68	25	354	140	41	2.15	761	
	7	42	6	43	102	24	2.21	95	
	7	58	16	111	90	20	2.22	246	
	7	60	17	122	66	214	1.58	193	
	1	76	35	35	66	214	1.58	55	
Total Air Flow (scfm):								1351	

Figure 4.9. Sparge Air Requirements and Resulting Zones of Influence – HLP-VSL-00027A/B

The bubble size and resulting mixing zone are based on the air flow in acfm at the sparge line exit. The required flow rate measured in scfm is based on the level and density of the liquid in the vessel. The scfm values in Figure 4.9 calculated based on the overflow level in HLP-VS-00027A/B and a slurry specific gravity of 1.35.

PJM Details: The central PJM has a downward-pointing, 4-inch nozzle that is 1.5 nozzle diameters (6 inches) off the bottom of the vessel. The outer seven PJM are located on a PCD of 131 inches. All of their exit nozzles are pointed outward toward the vessel sidewall at 45 degrees; the nozzle openings are also 4 inches and are 6 inches off the bottom of the vessel. Elevation views of the pulse tubes are shown in Figures 4.7 and 4.10; the outer pulse tube nozzle detail is shown in Figure 4.11.

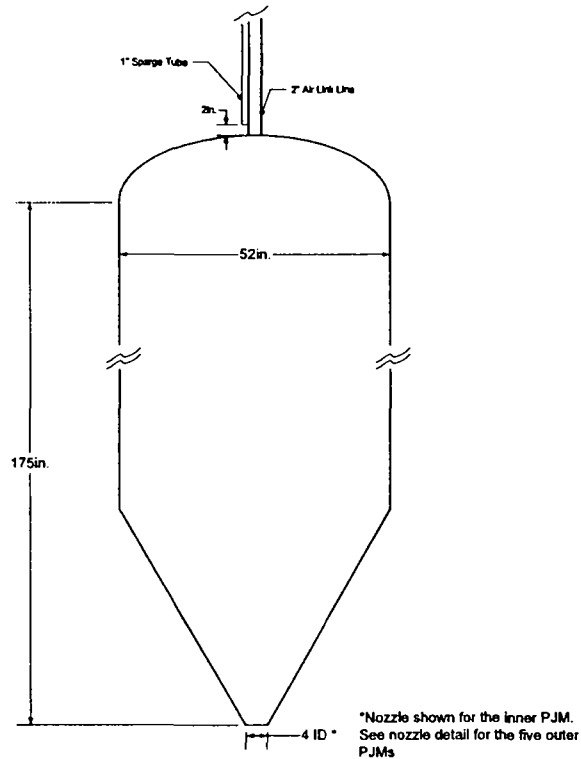


Figure 4.10. PJM Details – HLP-VSL-00027A/B, HLP-VSL-00028

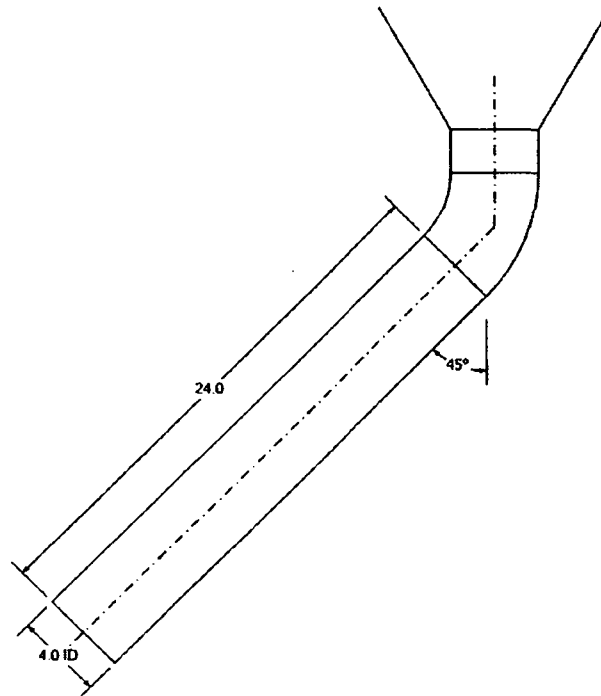


Figure 4.11. Outer PJM Nozzle Detail – HLP-VSL-00027A/B, HLP-VSL-00028

Sparge Details: The sparge tubes are all 1 inch; the outer three sets of PCD spargers, 180-, 204-, and 280-inch PCD, are approximately 6 inches off the bottom of the vessel. The spargers on a single PCD are spaced so the angle between them is the same; e.g., in a PCD set equal to 14 the spargers are spaced every 26 degrees. None of the exit tubes are near pulse tube nozzles or the recirculation pump intake. The spargers above the pulse tubes are approximately 2 inches above the top. Elevation views of the spargers are shown in Figures 4.7 and 4.10.

4.1.2 HLW Blend Vessel (HLP-VSL-00028)

Although a scaled prototypic test platform was not tested, the scaled testing in the LS prototypic platform can be applied to the blend vessel (BV) because the geometry and fluid properties are very similar.

Normal Operation: The HLW BV under normal operation will require eight PJMs and a 2200 gpm recirculation pump with one intake and two terminating nozzles configured to provide an exit velocity of 40 ft/sec to supply the required mixing. PJM nozzle velocities must be at least 12 m/sec during the PJM drive cycle. Under normal operation the sparge tubes will be run in an idling air flow-only mode, using 0.5-1.0 acfm (~2 scfm) of air per sparge point. Mixing with PJMs and recirculation jets has been tested up to an H/D of 0.74 in LS. This level scaled to the BV corresponds to an H/D of 0.70, which is more conservative than using the same H/D as in the LS, and is therefore recommended. The layout of PJMs, sparge lines and recirculation lines is shown in Figures 4.12, 4.7, and 4.8.

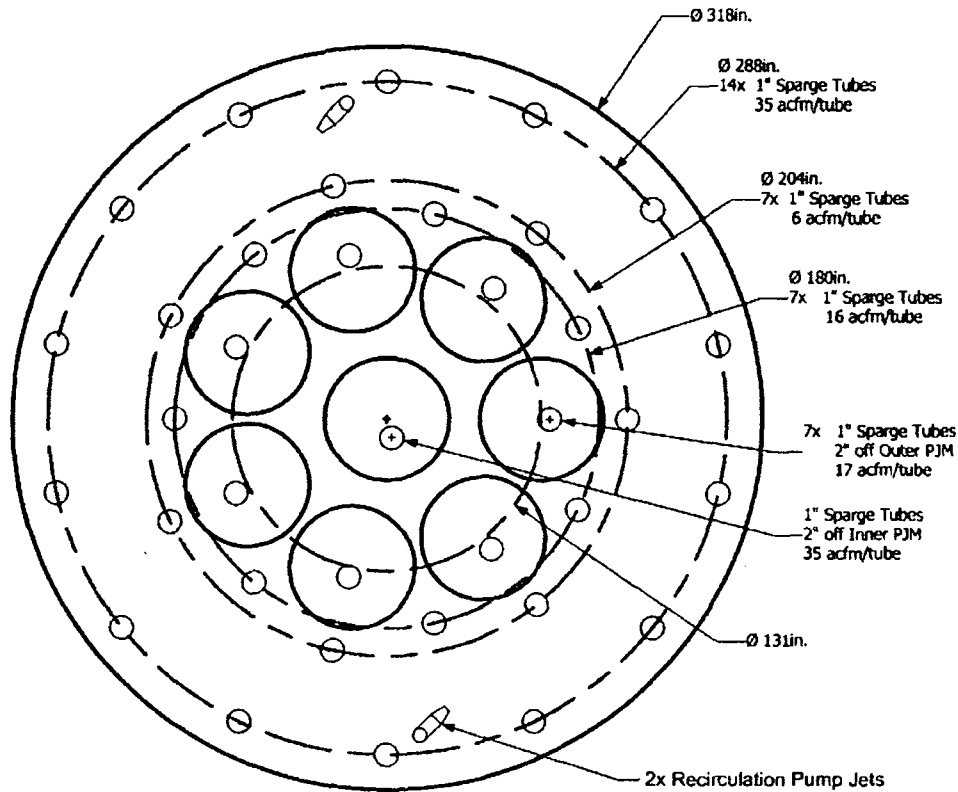
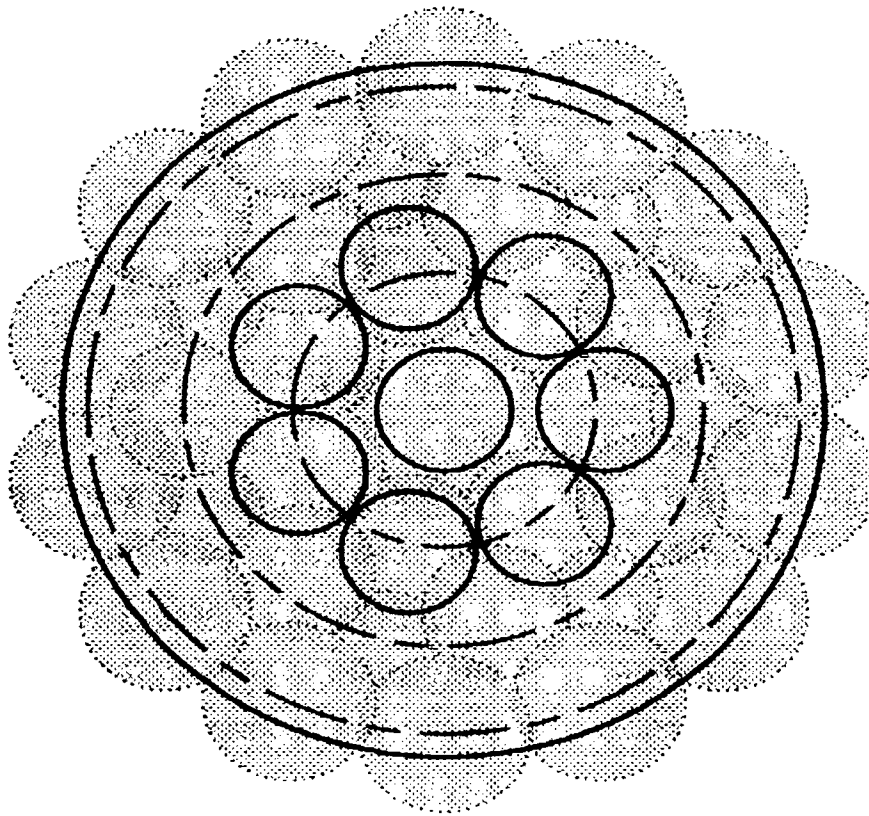


Figure 4.12. Specific Location of Spargers in the HLW BV Along with Air Flow Requirements

Post-DBE/High Levels: The BV under post-DBE conditions, or an H/D greater than 0.70, requires operation of PJMs and full sparging. Varieties of sparge tube layouts were considered for use. The configuration presented in this document was chosen by WTP engineering to minimize impact with consideration to total air requirement and total number of sparge lines. The selected configuration is shown in Figure 4.13. Hatched circles overlaid in the plan view indicate the size of the ZOI of each particular sparge tube at the flow rates specified in the adjacent table.

The bubble size and resulting mixing zone is based on the air flow in acfm at the sparge line exit. The required flow rate measured in scfm is based on the level and density of the liquid in the vessel. The scfm values in Figure 4.13 calculated based on the overflow level in HLP-VS-00028 and a slurry specific gravity of 1.35.



36 Sparge Tubes	# OF TUBES	2/3 ZOI (in.)	flow/tube (acfm)	subtotal (acfm)	Radial Position (in)	Nozzle Elevation (in)	Pressure Corr. Factor	subtotal (scfm)	HLP-VSL-00028
	14	76	35	491	144	40	2.14	1051	
7	42	6	43	102	23	2.20	94		
7	58	16	111	90	19	2.21	245		
7	60	17	122	66	214	1.57	192		
1	76	35	35	66	214	1.57	55		
Total Air Flow (scfm):								1638	

Figure 4.13. Number and Location of Spargers in the HLW BV Along with Air Flow Requirements

PJM Details: The central PJM has a downward-pointing, 4-inch nozzle that is 1.5 nozzle diameters or 6 inches off the bottom of the vessel. The outer seven PJM are located on a PCD of 131 inches. All of the exit nozzles are pointed outward toward the vessel sidewall at 45 degrees; the nozzle openings are 4 inches in diameter and 6 inches off the bottom of the vessel. Elevation views of the pulse tubes are shown in Figures 4.7 and 4.10 with the outer pulse tube nozzle detail shown in Figure 4.11.

Sparge Details: The sparge tubes are all 1 inch with the outer three sets of PCD spargers, 180-, 204-, and 288-inch PCD, all approximately 6 inches off the bottom of the vessel. All of the spargers on a single PCD are spaced so that the angle between them is the same; e.g., for a PCD set of spargers equal to 14, the spargers are spaced every 26 degrees. None of the exit tube locations are near pulse tube nozzles or the recirculation pump intake. The spargers above the pulse tubes are approximately 2 inches above the top. Elevation views of the spargers are shown in Figures 4.7 and 4.10.

5.0 Gas Retention and Release in Selected Prototype Vessel Configurations

The GR&R activity is focused on developing an understanding of flammable gas (e.g., hydrogen) retention and release in pulse-jet mixed tanks containing non-Newtonian wastes. Testing to date includes bench-scale development activities and experiments in PJM vessels covering a range of configurations and scales, all using non-Newtonian waste simulant. Several tests have been conducted to assess the volume fraction of gas retained in simulant during continuous gas generation and steady state PJM operation (i.e., gas holdup tests), and the gas release characteristics (volume and rate) after the restart of mixing following a stoppage (i.e., gas release tests). The following summarizes kaolin: bentonite clay simulant gas holdup and gas release tests completed in the UFP and lag storage prototype vessels using near-final design configurations and operating conditions. The basis for scale-up of the GR&R results is not fully reviewed and could not be included in this document.

5.1 Principle and Approach

To assess gas holdup and gas release in PJM tanks, gas bubbles are generated in situ in the simulant. The gas bubble generation technique is based on the decomposition of hydrogen peroxide (H_2O_2) on catalytic surfaces according to the following reaction:



Once sufficient H_2O_2 has decomposed to supersaturate the simulant in O_2 , bubbles nucleate and existing bubbles grow. Further decomposition of H_2O_2 leads to additional bubble nucleation and/or bubble growth as O_2 diffuses through the simulant to the bubbles. Generated gas will be retained or released depending on many factors, including the degree of mixing in the system, the retained gas volume fraction, the size of bubbles, and simulant rheology.

In gas holdup tests, H_2O_2 solution is added continuously for a period of time while the PJM system is operated normally to establish a constant gas generation rate. At steady state, the rate of gas generation equals the gas release rate (e.g., from bubbles migrating to the surface), and the steady-state gas volume fraction is termed the gas holdup. In gas release tests, the mixing system is shut down after an amount of H_2O_2 solution is added to allow gas bubbles to be retained in the quiescent simulant. The release of gas upon restart of the mixing system is tracked to assess gas release volumes and rates.

The primary data obtained in gas holdup and gas release tests are on the simulant surface level as a function of time. Through independently established correlations, the level measurements are used to calculate retained gas volume and gas volume fractions. The gas volume fraction α referenced to the initial simulant volume is defined as

$$\alpha = \frac{V_{gas}}{V_o} = \frac{V_{gas}}{V_{sim} + V_{sol}} \quad (5.2)$$

where V_{gas} is the volume of retained gas (e.g., O_2 bubbles), and the total initial slurry volume V_o includes the bubble-free simulant volume V_{sim} and the volume of H_2O_2 solution V_{sol} . In many cases V_{sol} is negligible compared with the large volume of gas-free simulant. However, in gas holdup experiments where H_2O_2 solution is added continuously for an extended period of time, a correction is made for the added solution volume.

According to the expected reaction stoichiometry (shown in Equation 5.1), two moles of H_2O_2 decompose to produce 1 mole of O_2 and 2 moles of H_2O . Using this relationship, the nominal H_2O_2 solution concentration (30 wt%), and ideal gas law considerations, the equivalent volumetric rate of O_2 gas generation can be determined for a given rate of H_2O_2 decomposition. Assuming instantaneous H_2O_2 decomposition or a steady process where a steady-state concentration of H_2O_2 is established in the slurry, O_2 gas is generated at a rate equivalent to H_2O_2 introduction. The latter is assumed to occur in gas holdup experiments, and reported steady-state volumetric gas generation rates (at 22°C and 1 atm) are calculated from measured H_2O_2 injection rates. Normalizing the gas volume generation rate by the volume of simulant in the vessel gives the specific volumetric gas generation rate (volume of O_2 gas/volume of simulant/time).

5.2 Gas Holdup in Normal Operations

This section demonstrates that gas is released regularly and controllably in normal operation of the LS and UFP prototype systems, resulting in relatively low gas holdup. Figure 5.1 plots the measured gas volume fraction as a function of time during and after a gas holdup test in the LS scaled prototype (Sequence 15). At elapsed time 0, a hydrogen peroxide addition rate was established to provide an effective O_2 gas generation rate of 0.18 vol%/min (normalized to atmospheric pressure and 22°C). The specific gas generation rates used in the prototype experiments exceed the expected maximum actual

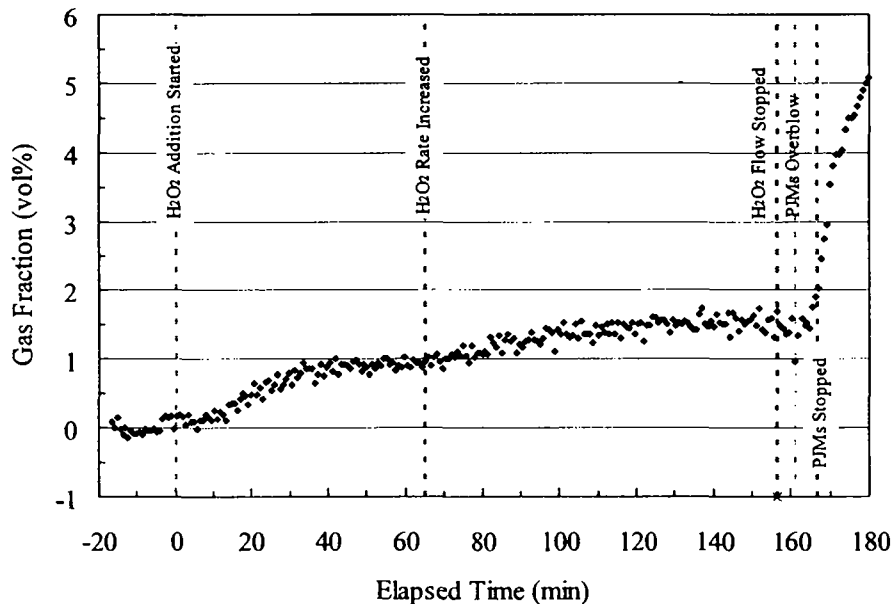


Figure 5.1. Gas Fraction as a Function of Time During and After a Gas Holdup Test in the APEL LS Prototype (Sequence 15). Events are marked on the plot by vertical lines.

waste gas generation rates (e.g., 2-4 vol%/day) by a factor of ~100 or more. Section 2.5 contains additional experimental details.

A steady-state gas fraction of ~0.9 vol% was attained after ~45 minutes. The rate of H₂O₂ addition was increased to an effective gas generation rate of 0.37 vol%/min after ~65 minutes, and a second steady-state gas holdup of 1.5 vol% was measured after ~125 minutes. The average gas holdup values in the last 10 minutes of H₂O₂ injection at each rate are tabulated in Table 5.1. The table also shows the standard deviation of the results (one value per pulse cycle, as shown in Figure 5.1).

As would be expected for a well-mixed system, the measured gas holdup in lag storage increased with increasing gas generation (H₂O₂ addition) rate.^(a) Figure 5.2 and Table 5.2 summarize the gas holdup experimental results for the two UFP prototype test sequences (5 and 6). A single relatively high equivalent gas generation rate was used in these tests (0.4 to 0.5 vol%/min).

Table 5.1. Summary of Gas Holdup in the LS Vessel (Sequence 15, 8-PJM cluster with 7 45° plus one vertical nozzles at ~12 m/s; 4-nozzle recirculation at ~120 gpm; 0.74 H/D; Bingham plastic rheology: 35-37 Pa yield stress, 26-27 cP consistency)

Experimental Gas Generation Rate (vol%/min)	Measured Gas Holdup±Standard Deviation (vol%)
0.18	0.93±0.06
0.37	1.5±0.1

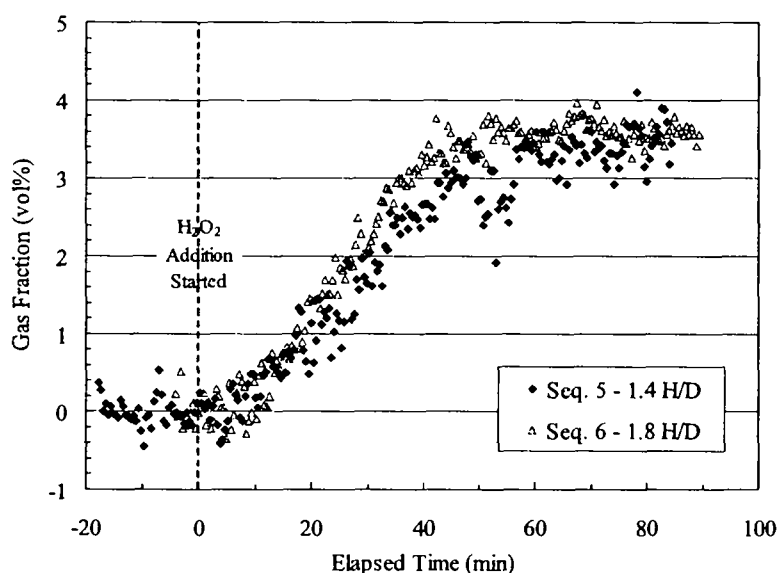


Figure 5.2. Gas Holdup Test Results in APEL UPF Prototype Using Two Sets of Operating Conditions: 1) Seq. 5, ~1.4 H/D, 4 PJMs + recirculation pump; and 2) Seq. 6, ~1.8 H/D, 4 PJMs + 1 sparger)

(a) Scaling arguments (not reported) suggest the holdup gas fraction should be proportional to the specific gas generation rate and the vessel scale factor (e.g., relative height for geometrically scaled vessels).

Table 5.2. Summary of Gas Holdup in UFP Vessel (Bingham plastic rheology: 33-36 Pa yield stress, 19-20 cP consistency)

Configuration	Experimental Gas Generation Rate (vol%/min)	Measured Gas Holdup±Standard Deviation (vol%)
Sequence 5: 1.4 H/D; 4 PJMs + recirculation pump at ~90 gpm	0.46	3.5±0.3
Sequence 6: 1.8 H/D; 4 PJMs + 1 sparge tube at ~3 cfm	0.41	3.6±0.1

5.3 Gas Release after Mixing System Restart

During a shutdown in which the air supply to PJMs and spargers is interrupted and recirculation pumps are idled, generated gas is expected to accumulate in the quiescent waste. In the extreme, all gas generated during the outage will be retained in the waste slurry. Upon restart of the mixing apparatus, accumulated gas is likely to be released. The release rate is dependent on many factors including waste rheology and mixing energy. Examples of gas release from gelled clay resulting from the restart of PJMs and spargers in the LS and UFP prototypes are provided below. In each of these tests the clay rheology exceeded the upper bounding Bingham plastic yield stress (>30 Pa). Based on preliminary matrix experiments conducted in the APEL 4PJM system, gas release rates may be faster as the simulant or waste slurry is thinned (lower rheological parameters).

Figure 5.3 shows the results of two gas release tests in the LS prototype. In the "overnight growth" test, H₂O₂ was introduced the day before the release began, and in the "30-min growth" test, the release experiment was started shortly after a preceding gas holdup test. The figure indicates relatively rapid initial gas release in both experiments. Nearly complete gas release was obtained after ~20 minutes (~27 pulses at 45 s/pulse cycle) in the overnight growth test with an initial gas volume fraction of ~4.5 vol%. Starting at a somewhat higher initial gas fraction (~5.8 vol%), gas was released to a retained gas volume fraction of ~1.8 vol% in 40 min in the "30-min growth" experiment, after which the gas fraction decreased slowly in time. In general, the characteristics of the initial gas release profiles (e.g., exponential decay) in the two experiments are similar. However, as noted above, the gas release rate decayed significantly in the "30-min growth" case before the retained gas was fully released.

Differences in gas release characteristics shown in Figure 5.3 are not fully explained at this point. Several contributing factors are under consideration: 1) differences in coincidental gas generation due to residual H₂O₂ decomposition; 2) differences in simulant rheological properties (e.g., strength) due to gel time, aging effects, and other factors (e.g., initial gas fraction); 3) differences in nominal bubble size, which may be a function of aging (bubble ripening), initial gas fraction, and rate of nucleation (a function of H₂O₂ concentration during bubble formation); 4) differences in initial gas fraction, which may make certain regions of the tank more difficult to mix due to buoyancy effects; 5) level sensors that do not measure tank level over the entire surface and therefore do not represent the total average gas fraction (note, however, that the sensor positions were not changed significantly between tests); and 6) other unaccounted differences in experimental variables such as sparger flow rate (e.g., no individual pressure gauges available on sparge tubes in the overnight growth test) and initial simulant depth (~5-cm lower in the overnight growth test).

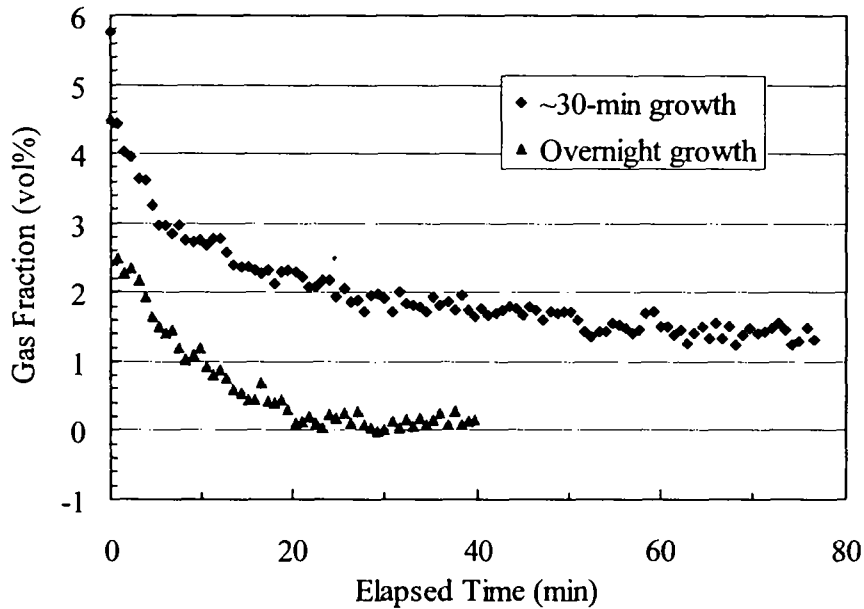


Figure 5.3. Gas Release from Gelled Clay in the APEL LS Prototype (Sequence 15, eight PJMs + four spargers).

After a near steady-state gas fraction (~ 1.4 vol%) was achieved in the "30-min. aging" test, the LS mixing system was stopped temporarily and restarted using eight (instead of four) sparge tubes at ~ 3 cfm each for ~ 10 min. These data are shown at long durations in Figure 5.4. The data indicate that retained gas volume fraction was reduced by ~ 1 vol% with the increased number of spargers.

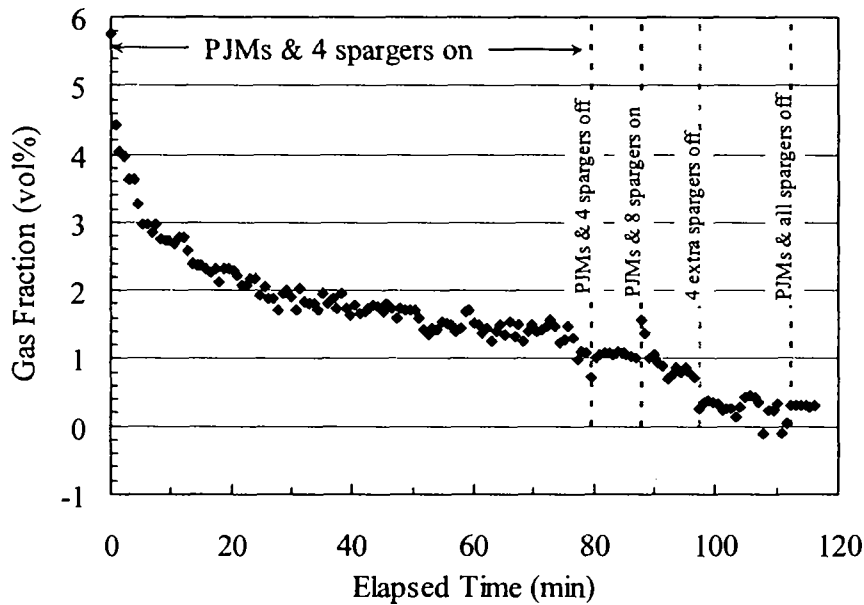


Figure 5.4. Gas Release from Gelled Clay in the APEL LS Prototype Showing Additional Gas Release Resulting from Operation of Eight Spargers (Sequence 15)

Gas release tests in the UFP prototypes were conducted with initial simulant loading to 1.4 and 1.8 H/D. The results are shown in Figures 5.5 and 5.6, respectively. The four-PJM tri-foil and single central sparge tube (~3 scfm) configuration was used in each test. As in the LS test, additional gas was released in some tests (e.g., Sequence 5, “~30-min. growth” shown in Figure 5.5) by using more spargers. However, Figures 5.5 and 5.6 only show the results for the specified baseline sparger operation (single sparge tube). At a given initial simulant loading (H/D value), the initial gas release profiles (i.e., change in gas fraction as a function of time) are consistent for the short and long growth cases. Differences at longer times, if any, may be due to factors identified above in the discussion of LS gas release tests.

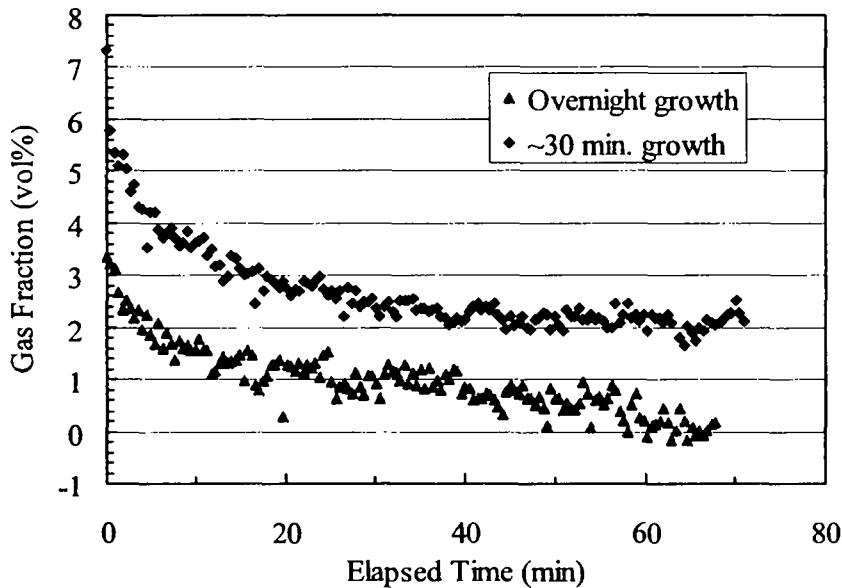


Figure 5.5. Gas Release from Gelled Clay at 1.4 H/D in the APEL UFP Prototype (Sequence 5)

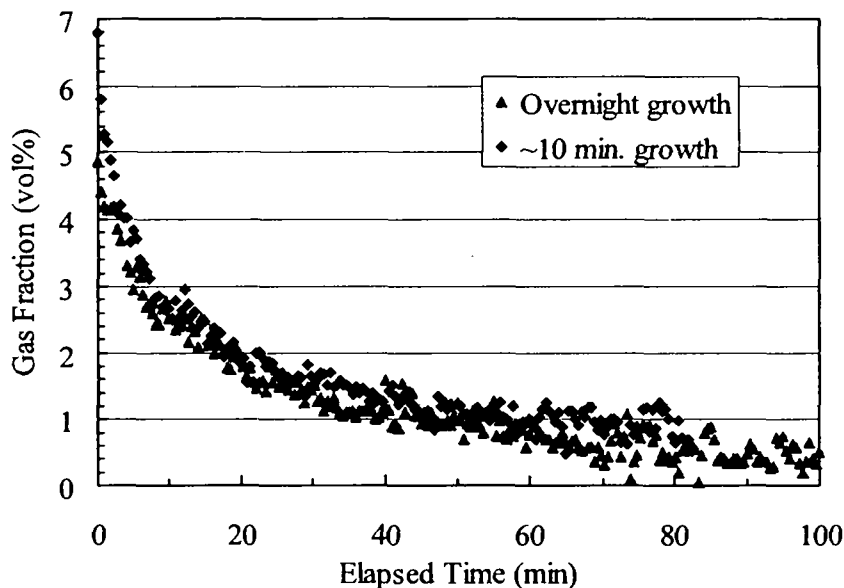


Figure 5.6. Gas Release from Gelled Clay at 1.8 H/D in the APEL UFP Prototype (Sequence 6)

6.0 References

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Appendix A

Technical Basis for Scaled Testing of WTP Mixing Vessels with Non-Newtonian Slurries

Appendix A

Technical Basis for Scaled Testing of WTP Mixing Vessels with Non-Newtonian Slurries

A.1 Introduction

Small-scale testing is a common approach used successfully in the many varied fields of applied fluid dynamics. The success of the approach depends greatly on the fact that system performance depends on certain non-dimensional groupings of physical parameters. If these parameter groupings can be preserved at different geometric scales (i.e., large and small), the essential behavior of the system will be the same at both scales. This principle is referred to as *similarity* in the theory of fluid dynamics engineering. Limitations of scaled testing are attributed to the inability to match important non-dimensional parameter groupings at both scales. In complex fluid dynamic problems, there can be many non-dimensional parameter groups; however, often the essential behavior of the phenomenon is dominated by only a few key groups. In this situation small-scale testing can produce results that are very close to large-scale behavior.

This appendix presents the approach used to establish the scalability of the scaled prototypic mixing tests. Section A.2 gives a brief introduction to the basics of pulse jet mixer (PJM) operation. Section A.3 gives a summary of the important properties and parameters involved in PJM mixing of non-Newtonian materials. Section A.4 explains the geometric scaling approach and how velocities and time are scaled. Section A.5 discusses the important non-dimensional parameters which, ideally, are to be preserved during scaled testing. Finally, Section A.6 summarizes the basis for scaled-testing.

A.2 Principles of PJM Operation

A schematic of a typical PJM system in a vessel is shown in Figure A.1. The tank has diameter D_T , volume V_T , and an operating level H . There are N PJMs in the tank, each with diameter D_{PT} and volume V_{PT} . Each PJM has a conical nozzle with diameter d_0 . For the baseline design, the total volume of the pulse tubes $N V_{PT}$ is approximately 10% the operating volume of the vessel.

There are three phases to the operation of the PJM. During the drive phase, the tube is pressurized and a volume of slurry is discharged. The level change in the tube during discharge is ΔL . The corresponding increase in waste level is ΔH where

$$\Delta H = N\Delta L \left(\frac{D_{PT}^2}{D_T^2 - ND_{PT}^2} \right) \quad (\text{partially submerged PJMs}) \quad (\text{A.1})$$

or

$$\Delta H = N\Delta L \frac{D_{PT}^2}{D_T^2} \quad (\text{fully submerged PJMs}) \quad (\text{A.2})$$

Typical values of ΔH are about 10% of the operating level H . The average velocity u_0 discharged during the drive phase is given by

$$u_0 = \frac{D_{PT}^2 \Delta L}{d_0^2 t_D} \quad (A.3)$$

where t_D is the drive time.

The drive pressure, p_D , required to produce the discharge velocity is given by

$$p_D = p_e + \frac{C_L}{2} \rho u_0^2 \quad (A.4)$$

where p_e is the pressure head at the exit of the nozzle, C_L is the nozzle loss coefficient, and ρ is the slurry density. The other two phases of PJM operation are the vent phase and suction phase.

Immediately after the drive phase, a vent is opened and excess pressure is allowed to vent to the atmosphere. During the suction phase, vacuum is applied to the pulse tube. The tube fills due to a combination of the applied vacuum and the difference in hydrostatic head between the waste level and the level in the tube. The vent time and suction time are given by t_V and t_S , respectively. The total cycle time for PJM operation is given by

$$t_C = t_D + t_V + t_S \quad (A.5)$$

It is important to emphasize that the average drive velocity given by Eq. (3) is both spatially and temporally averaged. Spatially, the velocity will vary over the cross section of the nozzle. Temporally, the velocity varies due to inertial effects. When the drive phase is over, some fluid continues to discharge due to the inertia of the moving column of fluid. These inertial effects are dependent on the physical size of the system. The actual velocity varies somewhat over the operating cycle, as shown in Figure A.2.

For comparing PJM operation at different scales, various average velocities can be considered. One is the area-averaged velocity, given by

$$\bar{u}_{\text{area}} = \frac{1}{t_p - t_m} \int_{t_m}^{t_D} u \, dt \quad (A.6)$$

Another is the true average velocity given by

$$\bar{u}_{\text{disch}} = \frac{D_{PT}^2 \Delta L_A}{d_0^2 t_{DA}} \quad (A.7)$$

where ΔL_A and t_{DA} are the actual measured level change and drive times in the pulse tube. Generally, Equation (A.6) will produce higher velocities than Equation (A.7).

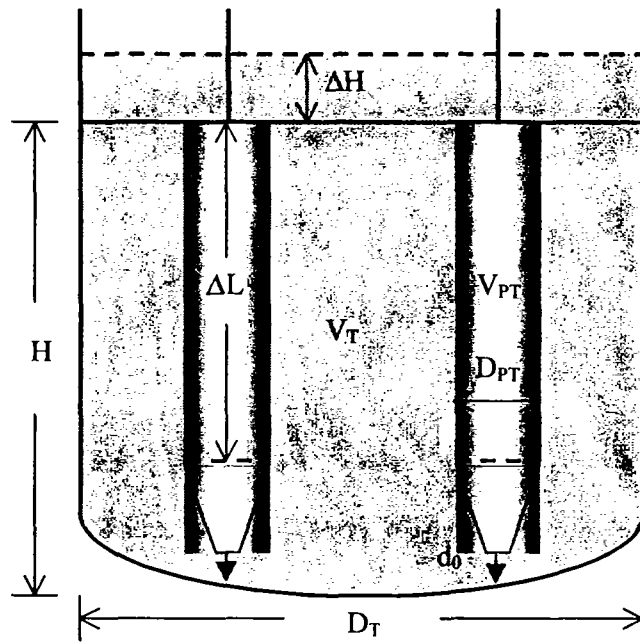


Figure A.1. Illustration of a Typical PJM System in a Waste Treatment Plant Vessel

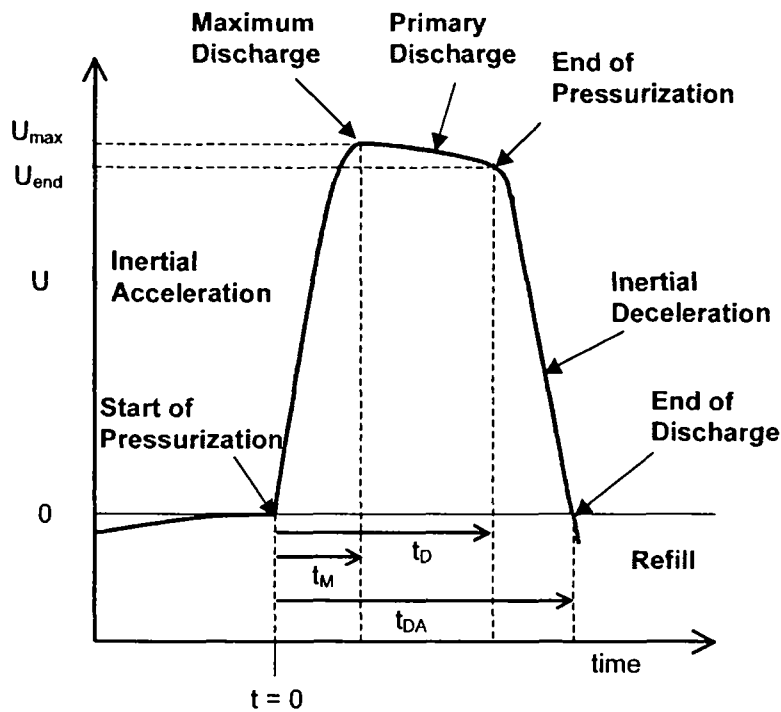


Figure A.2. Illustration of Temporal Variation of Velocity During PJM Operation

A.3 Important Properties, Parameters, and Non-Dimensional Groups

The following is a list of pertinent waste properties and system parameters to be used in forming non-dimensional parameter groups:

- Waste properties

ρ	slurry density (kg/m^3) (assumes well-mixed slurry with no settling)
τ_s	slurry shear strength (Pa)
τ_0	laminar flow yield stress (Pa) (from Bingham plastic fit of waste rheogram)
K	laminar flow consistency (mPa-s) (assumed to be effective Newtonian viscosity (μ) in turbulent region)
t_{rel}	slurry relaxation time (s) (characteristic response time of gelled slurry to an impulse)

- Physical parameters

u_0	nominal PJM jet velocity (m/s) (may be replaced with an averaged velocity)
d_0	PJM nozzle diameter (m)
t_D	PJM nominal drive time (s) (or actual drive time)
t_c	cycle time (s)
H	waste fill level (m)
V	vessel volume (m^3)
V_{PT}	pulse tube volume (m^3)
p	average hydrostatic pressure $\rho gH/2$ (Pa)
Q_0	PJM flow rate (per pulse) $(\pi/4)u_0d_0^2$ (m^3/s)
P_0	PJM hydraulic power (per pulse) $(\pi/8)\rho u_0^3d_0^2$ (W)

The relevant non-dimensional parameter groups for the physical system are as follows:

Yield Reynolds number:
$$\text{Re}_\tau = \frac{\rho u_0^2}{\tau_s}$$

This is the ratio of dynamic stress to slurry strength which directly affects size of the mixing cavern. It is considered a dominant non-dimensional parameter.

Jet Reynolds number:
$$\text{Re}_0 = \frac{\rho u_0 d_0}{\mu}$$

This is the ratio of dynamic stress to viscous stress. It affects the degree of turbulence in the mixed region as well as weakly affecting stresses at the cavern and boundary layers. It is considered a secondary non-dimensional parameter.

Non-Newtonian stress ratio:
$$N_\tau = \frac{\tau_s}{\tau_0}$$

This is the ratio of shear strength to Bingham yield stress. It may affect boundary layer structure and possibly the friction coefficient at the cavern boundary. The importance of this parameter is considered low.

Strouhal number:
$$S_0 = \frac{t_D u_0}{d_0}$$

This is the ratio of pulse time to flow time scale. It affects the degree to which flow approaches steady jet behavior and is considered a primary non-dimensional parameter. In the limit of steady jet flows, the Strouhal Number become infinite, and the effects of pulsation are no longer present. For small Strouhal number, the mixing behavior will be highly dominated by pulsation effects.

Deborah number:
$$D_0 = \frac{t_D}{T_s}$$

This is the ratio of pulse time to material response time. It affects how well non-steady flow at cavern mobilizes gelled slurry and is considered a secondary non-dimensional parameter.

Pressure ratio:
$$\frac{p_a}{\rho g H}$$

This is the ratio of ambient pressure to static head. It affects the scaling of gravity refill of a PJM but should not affect the discharge flow.

Densimetric Froude number:
$$F_0 = \frac{\rho u_0^2}{\Delta \rho g H}$$

This is the ratio of the potential energy to kinetic energy of flow. It requires density stratification and affects the ability of a jet to transport material upward. The importance of this parameter is considered low due to minimal solids settling in the turbulent region.

A.4 Geometric Scaling Approach

The non-Newtonian test program uses geometric scaling. We define the geometric scale factor s as

$$s = \frac{L_L}{L_S} \tag{A.8}$$

where L_L is any characteristic linear dimension of the large-scale system (such as tank diameter, nozzle diameter, waste level, etc.). At small scale, every linear dimension, L_S , is reduced or *scaled* by s

(i.e., $d_{0s} = d_{0L} / s$, $D_{Ts} = D_{TL} / s$, $H_s = H_L / s$). Hence the ideal small-scale test is an exact geometric miniature of the large system, with all areas scaled according to

$$A_s = \frac{1}{s^2} A_L \quad (\text{A.9})$$

and all volumes scaled according to

$$V_s = \frac{1}{s^3} V_L \quad (\text{A.10})$$

Typically in scaled fluid mixing tests, scale factors up to about 10 are considered acceptable, that is, much of the important physics can be captured at small scale. For the non-Newtonian test program, conservative scale factors in the range of 4 to 5 were selected due to the relatively new nature of the tests and the importance of the outcome.

When testing at small scale, one must determine how to scale velocity (i.e., PJM drive velocity u_0). One choice is to scale velocity by the scale factor. This is problematic, however, because it tends to reduce the Reynolds number by $1/s^2$ and introduce further difficulties with the scaling of time. A better choice is to keep jet velocity constant at both scales:

$$u_{0s} = u_{0L} \quad (\text{A.11})$$

With geometric scaling and constant velocity scaling, nozzle flow rates per pulse scale according to

$$Q_{0s} = Q_{0L} / s^2 \quad (\text{A.12})$$

Jet hydraulic power also scales similarly. However, power per unit volume scales according to

$$\left. \frac{P_0}{V} \right)_s = s \left. \frac{P_0}{V} \right)_L \quad (\text{A.13})$$

For steady jet mixing, time does not come into play. However, PJM operation is a periodic process. Therefore, the scaling of time must be addressed.

If velocity is held constant and the geometry is scaled, then it follows that all imposed time scales must be reduced at small scale. Similarly, to keep the jet discharge velocity the same while scaling pulse volume geometrically, the pulse time will be reduced by the scale factor according to

$$t_{Ds} = \frac{1}{s} t_{DL} \quad (\text{A.14})$$

Hence the PJM drive time (as well as refill time and cycle time) are all reduced by s at small scale.

A.5 Scaling Non-Dimensional Parameters

In general, for a given non-Newtonian PJM mixing test, the non-dimensional cavern position should depend on all of the non-dimension parameter groups:

$$\frac{H_C}{D_T} = f(\text{Re}_\tau, \text{Re}_0, N_\tau, S_0, D_0, F_0) \quad (\text{A.15})$$

Similarly, non-dimensional mixing time (time to steady cavern formation, time to break through, or time to full mobilization) should depend on the same parameters:

$$\frac{t_M}{t_D} = g(\text{Re}_\tau, \text{Re}_0, N_\tau, S_0, D_0, F_0) \quad (\text{A.16})$$

The ideal small-scale test is one where the measured non-dimensional cavern height and mixing time are the same as those at full scale. Hence, the extent to which the non-dimensional parameters scale will determine the success of the small scale test approach.

To this end, we consider how each of the non-dimensional parameters scale with the geometric scale factor s :

Yield Reynolds Number: $\text{Re}_{\tau S} = \text{Re}_{\tau L}$

The yield Reynolds number will be the same at both scales so long as the simulant used has the same shear strength τ_s :

Jet Reynolds Number: $\text{Re}_{0s} = \frac{1}{s} \text{Re}_{0L}$

The Reynolds number at small scale is reduced by the geometric scale factor. This should introduce only minor differences in test results since the Reynolds numbers in both tests are quite large. Whether the reduction in Reynolds number produces conservative results (i.e., lower caverns) at small scale is not clear due to the competing effects of Reynolds number on jet structure and friction coefficients. The potential need for a minor Reynolds number correction to small-scale results should be evident from the scaling tests. If necessary, the Reynolds number can be matched at small scale by reducing the consistency or viscosity by the factor $1/s$.

Non-Newtonian stress ratio: $N_{\tau S} = N_{\tau L}$

The non-Newtonian stress ratio will be the same at both scales if the same simulant is used.

Strouhal number: $S_{0S} = S_{0L}$

The Strouhal number will be the same at both scales.

Deborah number:
$$D_{0S} = \frac{1}{s} D_{0L}$$

The Deborah number will be smaller in the small-scale tests. If the Deborah number is large overall, the effect will be negligible. If Deborah is close to unity, then the small-scale results will be conservative.

Densimetric Froude number:
$$F_{0S} = sF_{0L}$$

The densimetric Froude number will be larger at small scale. This would produce non-conservative results at small scale should the effect be important. So long as simulants with very slow particle settling are used, this effect should be negligible.

A.6 Summary of Scaled Test Approach

By way of summary, the primary non-dimensional parameters required for small-scale testing are the yield Reynolds number Re_{τ} , and the Strouhal number S_0 . If these are matched at large and small scale, then we expect, to first order, non-dimensional cavern heights and mixing times to be the same:

$$\left. \frac{H_C}{D_T} \right)_S \approx \left. \frac{H_C}{D_T} \right)_L \quad (A.17)$$

and

$$\left. \frac{t_M}{t_D} \right)_S \approx \left. \frac{t_M}{t_D} \right)_L \quad (A.18)$$

Given that full-scale cavern heights are adequately predicted by reduced-scale testing, it follows that specification of PJM operation parameters sufficient to achieve complete mixing (no stagnant regions) at reduced scale will produce designs that also provide complete mixing at full-scale. Further, testing at reduced scale will provide a degree of conservatism so long as the consistency, k , of the simulant is the same as the full-scale bounding value. This is true since the jet Reynolds number will be smaller in the scaled-test than in the full-scale system:

$$Re_{0s} = \frac{1}{s} Re_{0L} \quad (A.19)$$

If adequate mixing is achieved in a reduced-scale test, then it can be expected that the degree of turbulence will be greater in the full-scale vessel due the associated effect of increased jet Reynolds number.

Appendix B

Dye Method

Appendix B

Dye Method

The concentration of dye [in this case Food Dye Color No. 1, (Brilliant Blue FCF) (BB FCF)] in an aqueous sample was determined through the correlation shown in Figure B.1. This correlation follows Beer's law, which says that the dye concentration is proportional to the optical absorbance value of the dye at the mode wavelength. The mode wavelength for BB FCF is approximately 633 nm. The results are only valid over a certain region of dye concentration. From visual inspection of Figure B.1, the linear region is present up to an absorbance value of 1.5 (~9 ppm FCD1). When the dye concentration is above this level the sample must be diluted with water and remeasured. The original dye concentration can be calculated by knowing the quantity of water used for the dilution.

Beer's Law Chart of Brilliant Blue (FD&C Blue 1) in Water

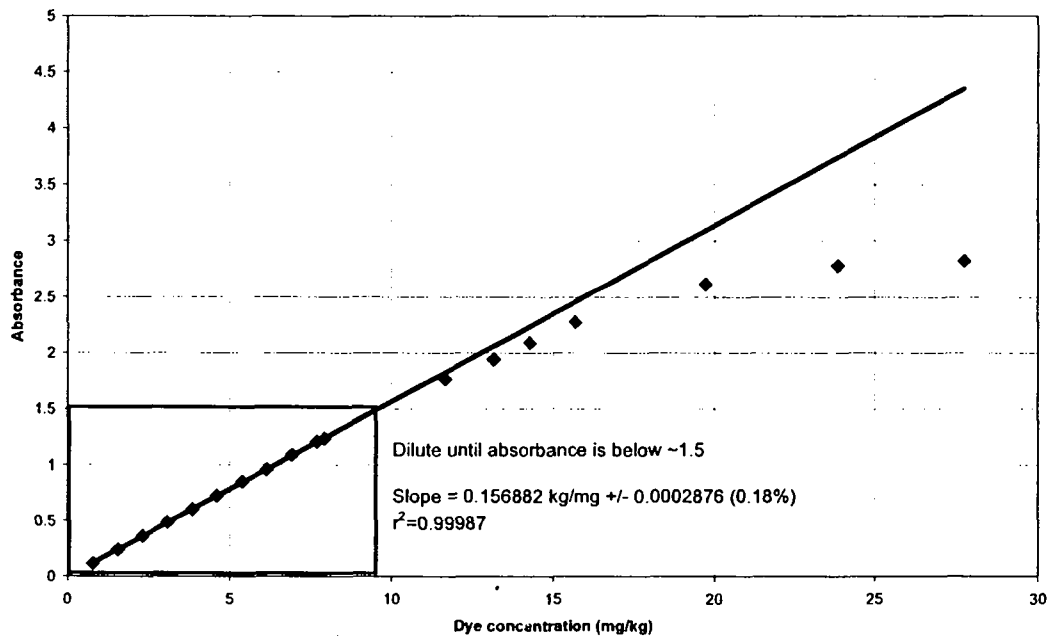


Figure B.1. Beer's Law Correlation of Optical Absorbance to BB FCF Dye Concentration in Water

Absorption of dye onto the surface of the clay particles can be estimated through a linear approximation. This correlation is shown in Figure B.2, where the dye concentration in the liquid phase is plotted against the dye concentration in the solid phase. Due to batch to batch variations of the clay composition, small differences in the amount of dye absorbed were measured from sample to sample. The linear isotherm assumption allows for the use of Equation B.1 to calculate percent mixed in a PJM test.

$$X_j = \frac{A_f - A_0}{A_j - A_0} \quad (\text{B.1})$$

where

X_j	is the fraction mixed of the j-th tank sample
A_f	is the optical absorbance of the final homogenized simulant
A_0	is the optical absorbance of the initial baseline simulant
A_j	is the optical absorbance of the j-th tank sample

Linear Approximation of Isotherm in Operational Dye Concentration Range

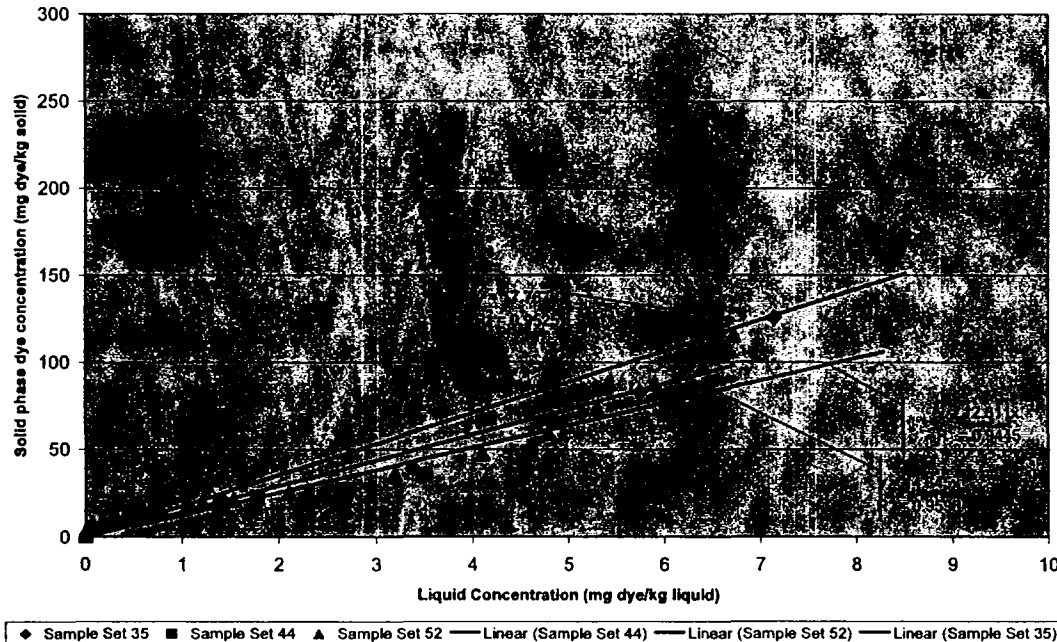


Figure B.2. Linear Fit of Isotherm Data over the Linear Beer's Law Region

A polynomial fit to one of the isotherm data sets is shown in Figure B.3. Use of this fit allows for an estimation of the error incurred through the assumption of a linear isotherm. This error is estimated by calculating the difference in the percent mixed between Equations B.1 and B.2. To perform this calculation the correlation shown in Figure B.3 is used to calculate the K_d values of each sample in the calculation. A conservative estimation of the solids loading in each sample is assumed at 30 wt% solids 70 wt% liquid.

$$X_j = \frac{Y_l(A_f - A_0) + Y_s(K_{df}A_f - K_{do}A_0)}{Y_l(A_j - A_0) + Y_s(K_{dj}A_j - K_{do}A_0)} \quad (\text{B.2})$$

where

K_{df}	is the distribution coefficient at the homogenized tank tracer concentration
K_{do}	is the distribution coefficient at the initial baseline tracer concentration
K_{dj}	is the distribution coefficient at the j-th tank sample tracer concentration

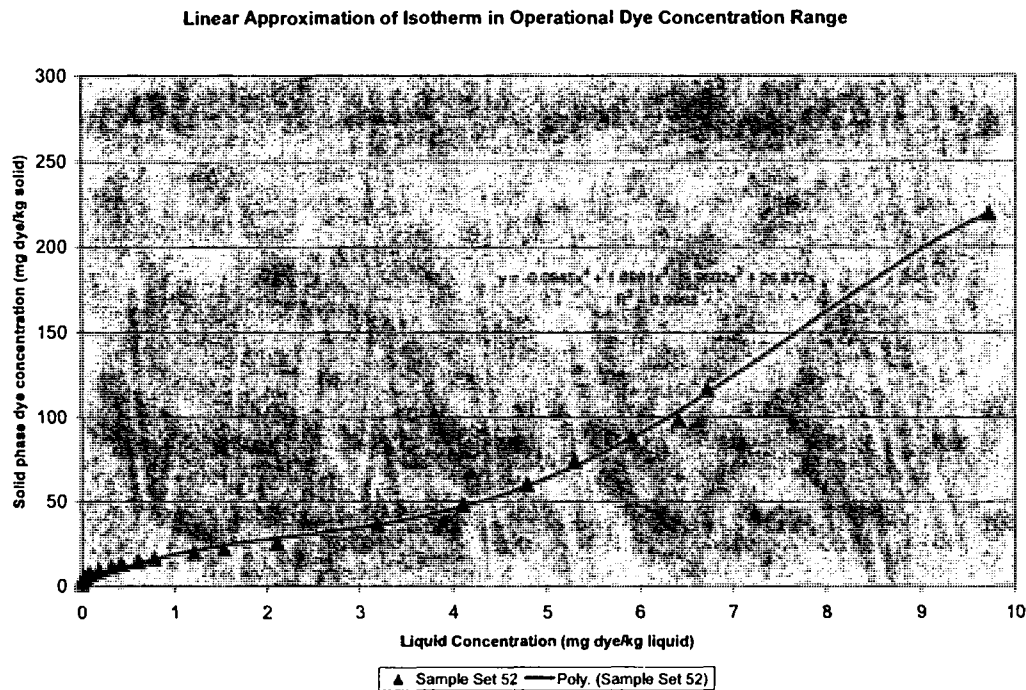


Figure B.3. Polynomial Fit of Isotherm Data over the Linear Beer's Law Region

During prototype testing, Equation B.1 was used to calculate a fraction mixed for each sample at each sample location. These samples were drawn from different locations in the testing vessel. Sample locations 1, 2, and 3 are from separate pulse tubes and represent the composition of the mixing cavern (see Section 1.4.6). Locations 4 and 5 were located near the tank wall at low and high elevations, respectively. During the first run of a test sequence, samples from locations 1, 4, and 5 were taken approximately every 10 minutes after completion of dye injection. After 50 minutes of operation, samples were drawn from all sample locations and the next run experimental condition was employed. During subsequent run conditions, samples from locations 1, 4, and 5 were taken every 15 minutes. After 45-90 minutes of operation, samples were drawn from all sample locations and the next run experimental condition was employed. The fraction of the tank mixed calculated from each sample is shown in Figures B.4 through B.7 for LS test sequences 4, 7, 11, and 20, respectively. Figures B.8 and B.9 show the fraction mixed results for UFP test sequences 2 and 3B.

The final fraction mixed value was determined as the minimum fraction mixed from the locations 1, 2, and 3 of the last sample of a test run. This represents the fraction mixed value associated with highest dye concentration in the cavern after approximately 45-50 minutes of operation. As discussed above, the error associated with the linear isotherm approximation is estimated through the use of Equation B.3. In the worst case, typical errors due to this assumption are approximately less than ± 0.15 fraction mixed; the error goes to zero as the fraction mixed approaches unity. The final fraction of the tank mixed calculated from each run is shown in Tables B.1 through B.4 for LS test sequences 4, 7, 11, and 20, respectively. Tables B.5 and B.6 show the fraction mixed results for UFP test sequences 2 and 3B. Although the NaCl tracer technique is discussed in this document, the NaCl tracer results are supplementary to the BB dye results and will be discussed in a future report.

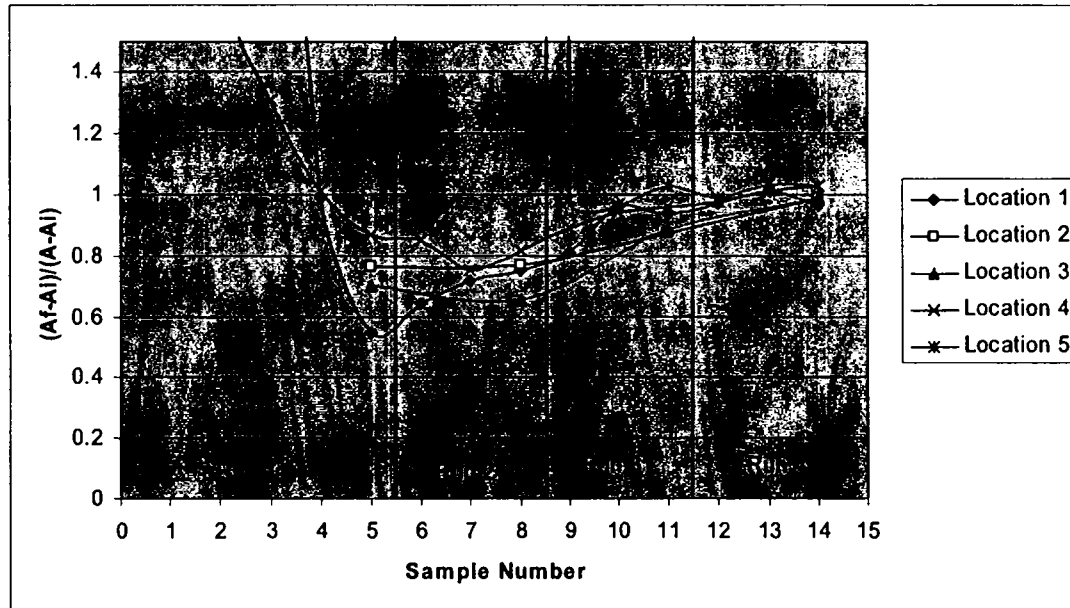


Figure B.4. Fraction Mixed Chart for LS Test Sequence 4

Table B.1. Final Fraction Mixed and Error Estimate Due to Linear Isotherm Assumption for LS Test Sequence 4

Run	Fraction Mixed	Linear Isotherm Estimated Error (\pm) ^(a)
1	0.54	0.15
2	0.65	0.13
3	0.87	0.052
4	0.97	0.014

(a) Estimated error due to assumption of linear isotherm for dye absorption. Experimental error not included.

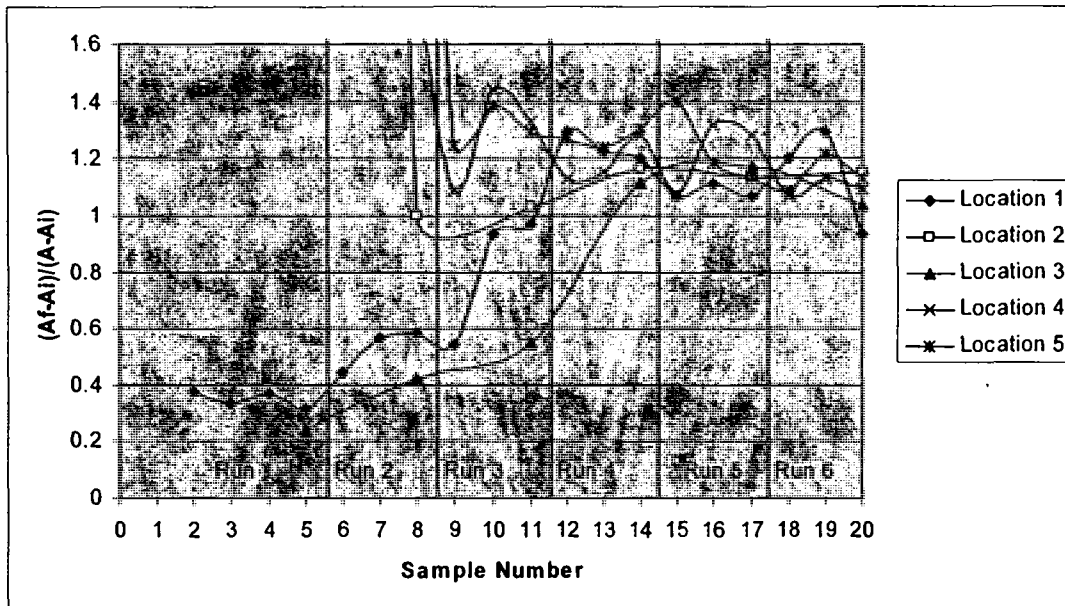


Figure B.5. Fraction Mixed Chart for LS Test Sequence 7

Table B.12. Final Fraction Mixed and Error Estimate Due to Linear Isotherm Assumption for LS Test Sequence 7

Run	Fraction Mixed	Linear Isotherm Estimated Error (\pm) ^(a)
1	0.24	0.11
2	0.42	0.085
3	0.55	0.060
4	1.1	0.010
5	1.1	0.0058
6	0.93	0.0067

(a) Estimated error due to assumption of linear isotherm for dye absorption. Experimental error not included.

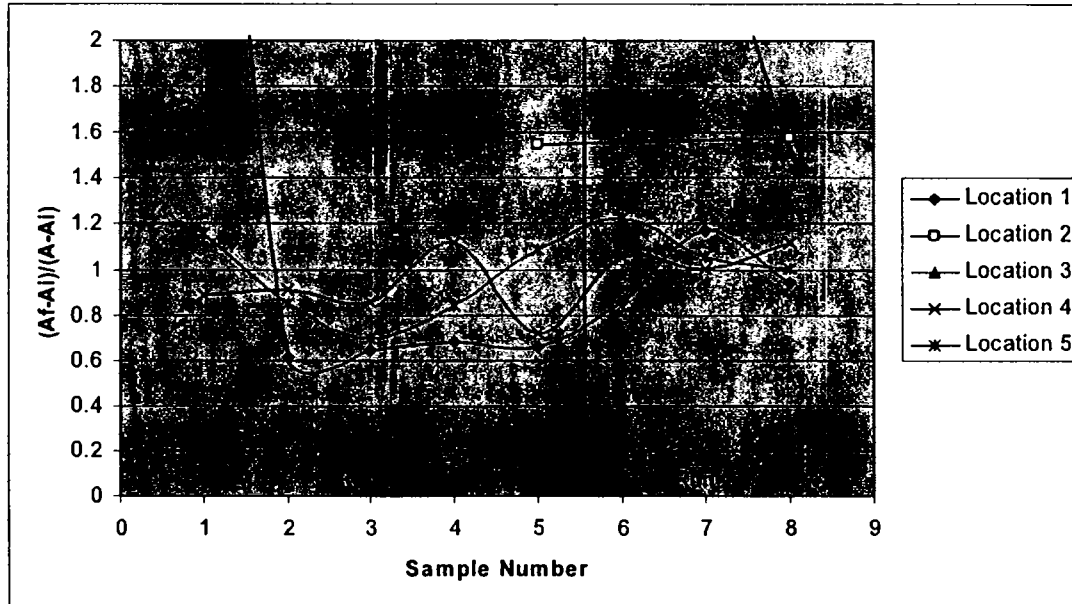


Figure B.6. Fraction Mixed Chart for LS Test Sequence 11

Table B.3. Final Fraction Mixed and Error Estimate Due to Linear Isotherm Assumption for LS Test Sequence 11

Run	Fraction Mixed	Linear Isotherm Estimated Error (\pm) ^(a)
1	0.66	0.033
2	0.95	0.0055

(a) Estimated error due to assumption of linear isotherm for dye absorption. Experimental error not included.

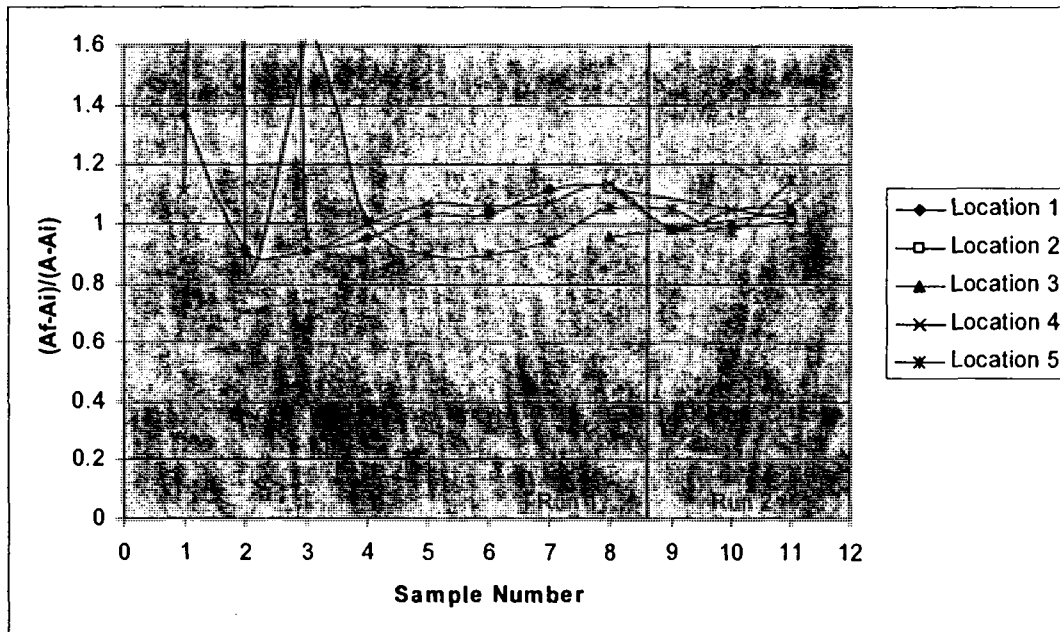


Figure B.7. Fraction Mixed Chart for LS Test Sequence 20

Table B.4. Final Fraction Mixed and Error Estimate Due to Linear Isotherm Assumption for LS Test Sequence 20

Run	Fraction Mixed	Linear Isotherm Estimated Error (\pm) ^(a)
1	0.96	0.0097
2	1.0	0.00069

(a) Estimated error due to assumption of linear isotherm for dye absorption. Experimental error not included.

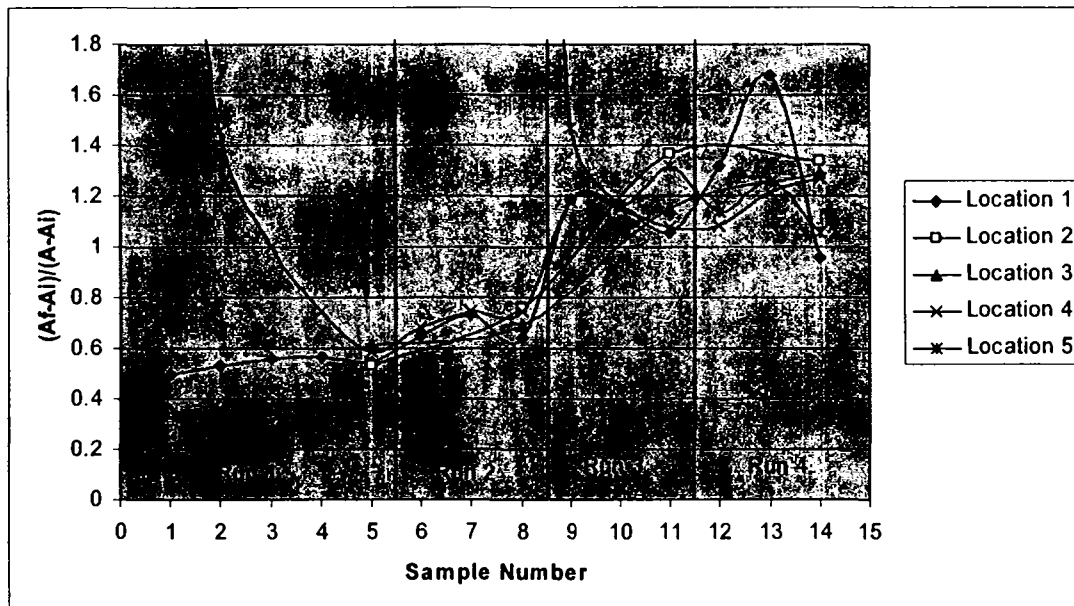


Figure B.8. Fraction Mixed Chart for UFP Test Sequence 2

Table B.5. Final Fraction Mixed and Error Estimate Due to Linear Isotherm Assumption for UFP Test Sequence 2

Run	Fraction Mixed	Linear Isotherm Estimated Error (\pm) ^(a)
1	0.53	0.093
2	0.64	0.074
3	1.1	0.013
4	0.96	0.0088

(a) Estimated error due to assumption of linear isotherm for dye absorption. Experimental error not included.

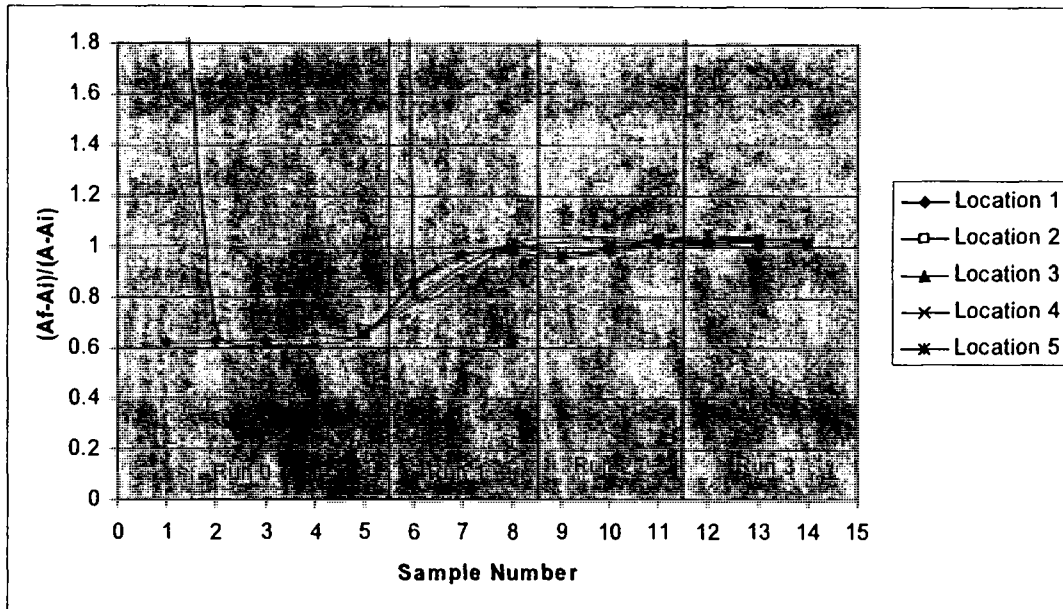


Figure B.9. Fraction Mixed Chart for UFP Test Sequence 3B

Table B.6. Final Fraction Mixed and Error Estimate Due to Linear Isotherm Assumption for UFP Test Sequence 3B

Run	Fraction Mixed	Linear Isotherm Estimated Error (\pm) ^(a)
1	0.65	0.12
2	0.98	0.0074
3	1.0	0.0019
4	1.0	0.0038

(a) Estimated error due to assumption of linear isotherm for dye absorption. Experimental error not included.

SEPARATION

PAGE



Memorandum

To: R. E. Smith MS4-A2 Date: March 22, 2004
From: G. M. Duncan MS4-D2 CCN: 085014
Ext: 371-3822
Fax: 371-3508

Subject: **PRETREATMENT NON-NEWTONIAN FLUID MIXING DESIGN REQUIREMENTS**

The following provides input to the Hanford Tank Waste Treatment and Immobilization Plant (WTP), Pretreatment Project Engineering regarding pulse jet mixer (PJM) and sparger design requirements for vessels containing non-Newtonian fluids. This input supports continuation of detailed physical design activities associated with piping, equipment, and structural interfaces.

Background

Mixing requirements for pretreatment (PT) vessels containing non-Newtonian fluids have been developed in a testing program conducted by WTP Research and Technology (R&T) department, with the close involvement of WTP Engineering. The PT vessels are as follows:

- Ultrafilter process (UFP) vessels (two vessels) [UFP-VSL-00002A, UFP-VSL-00002B]
- Lag storage (LS) vessels (two vessels) [HLP-VSL-00027A, HLP-VSL-00027B]
- PT blend vessel (one vessel) [HLP-VSL-00028]

Mixing system design requirements for both normal plant operation and for Important To Safety (ITS) operation have been determined. In each case, these requirements have been established based on a joint, Engineering and R&T, interpretation of the results of several months of 1/8-scale, 1/4-scale, and full-scale testing. These results have been presented in a series of project-wide meetings, with broad functional representation. These meetings are documented in correspondence control number (CCN) 077889.

R&T is in the process of developing a comprehensive series of reports to describe and document the testing results. A top-level summary report from R&T is planned for the end of March 2004, with a series of more detailed test reports being planned for issuance over the third quarter of 2004. In advance of this necessary documentation, these design requirements contained herein have been developed, to support continuation of detailed design and to support determination of related project cost and schedule impacts. For completeness, these design requirements will be supplanted by a system description.

Summary Specifications

For each group of PT non-Newtonian vessels, a mixing system has been selected. It should be noted that a combination of PJM's, spargers, and pumps are required to be deployed. The adopted specifications for each vessel group are summarized in Table 1.

Table 1: Summary Mixing System Specification

	SPECIFICATION	UFP	LS/BLEND
1	Recirculation Pump Required?	Yes	Yes
2	Recirculation Pump Flow (gallons per minute)	2200	2200
3	Pump discharge nozzle configuration within vessel	Single discharge nozzle (nozzle velocity - 30 feet per second)	2 discharge nozzles (nozzle velocity - 40 feet per second)
4	Spargers (quantity, size)	16, 1" (nominal diameter tubing)	36, 1" (nominal diameter tubing)
5	Sparger flow	(See Tables 3 and 4)	(See Tables 3 and 4)
6	Sparger minimum design pressure for cleaning	100 pounds per square inch gage (capability for water and compressed air cleaning shall be provided)	100 pounds per square inch gage (capability for water and compressed air cleaning shall be provided)
7	Antifoam	Required (supply from existing antifoam addition system)	Required (supply from existing antifoam addition system)
8	Waste Rheology	Non-Newtonian [Bingham Plastic yield stress of 30 Pascal, and a consistency viscosity of 30 centipoise (Reference: CCN 065607)]	Non-Newtonian [Bingham Plastic yield stress of 30 Pascal, and a consistency viscosity of 30 centipoise (Reference: CCN 065607)]
9	PJM (quantity)	6	8
10	PJM Configuration	Cluster (with inner shroud)	Cluster (with inner shroud)
11	PJM Nozzle diameter (inches)	4	4
12	PJM Nozzle velocity (meters per second)	12.0 minimum/14.2 maximum	12 minimum/13.4 maximum
13	PJM Air line diameter (inches) from jet pump pair (JPP) to vessel	2 (bounding line length of 200 feet)	2 (bounding line length of 200 feet)
14	PJM air demand (peak, average)	(See Tables 3 and 4)	(See Tables 3 and 4)
15	JPP Outline Dimensions	Same as AEA Technology (AEA) Model L50M (original JPP for this application)	Same as AEA Model L50M (original JPP for this application)

Vessel Mixing Mode Specifications

Table 2 provides a top-level summary of the required mixing modes of operation, for each vessel.

Table 2: Summary Operating Modes

	SPECIFICATION	UFP	LS/BLEND
1	Normal Mixing Mode	<ul style="list-style-type: none"> • Pump - ON • PJM's - ON • Spargers - IDLE [up to 1.4 aspect ratio for 30 Pascal fluid; up to 1.8 aspect ratio with leached (lower yield strength) fluid]	<ul style="list-style-type: none"> • Pump - ON • PJM's - ON • Spargers - IDLE [up to 0.74 aspect ratio for 30 Pascal fluid; spargers to be activated at higher fluid level]
2	Leach Mixing Mode	<ul style="list-style-type: none"> • Pump - OFF (high temperature leach mode) • PJM's - ON • Spargers - ON [> 1.4 aspect ratio, temperature up to ~194 degrees F]	N/A
3	Pump Failure Mixing Mode [normal power available]	<ul style="list-style-type: none"> • Pump - Not available • PJM's - ON • Spargers - ON 	<ul style="list-style-type: none"> • Pump - Not available • PJM's - ON • Spargers - ON
4	ITS Mixing Mode (loss of normal power) [Note: ON/OFF frequency and duration of this mode to be determined]	<ul style="list-style-type: none"> • Pump - Not Available • PJM's - ON • Spargers - ON 	<ul style="list-style-type: none"> • Pump - Not Available • PJM's - ON • Spargers - ON

Important to Safety (ITS) Requirements

ITS compressed air to the sparger system and the PJM system for each vessel shall be provided. Compressed air shall be provided to all PJM's and all spargers, assuming a single active failure. [See PT Preliminary Safety Analysis Report, Section 4.3.4 for project application of single failure criterion.]. The ITS compressor size, number, and operational sequences will be documented separately from this memorandum.

During ITS operation of the PJMs and the spargers, exhaust air may be directed towards the Zone 5 heating, ventilation and air conditioning system.

Additional safety criteria, generically applicable to the design of all WTP ITS systems, structures, and components (SSCs), are addressed within the *Safety Requirements Document* and are not repeated herein.

It should be noted that HLP-VSL-00022, although containing Newtonian fluids, generates sufficient hydrogen that its PJM system is ITS. Therefore, these same ITS requirements as noted above shall apply to this Newtonian vessel.

Spargers

Each sparger shall be capable of individual controlled flow. Two operating modes of sparger operation are necessary. In the IDLE mode, at least a minimum flow of air shall be provided (on the order of 1 standard cubic feet per minute) to maintain sparger nozzle clear of obstructions. In the ON mode, each sparger shall operate at its specified flowrate. Each sparger shall have the capability of being flushed, using either full pressure compressed air or water.

Compressed Air Consumption

Air consumption for PJM and sparger operation, for normal mixing operation, and for ITS mixing operation, is shown in Tables 3 and 4, respectively. It should be noted that sparger air flow is based on maximum operating volume, corresponding to an overflow condition, and is therefore considered to be bounding. With respect to PJM air flow, the noted figures in Tables 3 and 4 are the direct result of AEA's calculations, and do not contain any contingency allowance. A margin of 10% should be applied to these PJM flows, to account for uncertainty regarding JPP air consumption.

Table 3: Compressed Air Consumption - Routine System Operation

Vessel	# of Spargers	Sparger Mode	Total Sparge Air (SCFM)	# of PJMs	Peak PJM Air (SCFM)	Cycle Ave PJM Air (SCFM)	Total Air Required (SCFM)
UFP-VSL-00002A	16	idle	32	6	5016	1140	1172
UFP-VSL-00002B	16	idle	32	6	5016	1140	1172
UFP-VSL-00002A*	16	ON	394	6	5016	1140	1534
UFP-VSL-00002B*	16	ON	394	6	5016	1140	1534
HLP-VSL-00027A	36	idle	63	8	5936	1288	1351
HLP-VSL-00027B	36	idle	63	8	5936	1288	1351
HLP-VSL-00028	36	idle	63	8	5936	1288	1351

*The recirculation pump is not running during the (high temperature) leaching process. If a batch requires leaching, the spargers will be used for mixing during this process.

Table 4: Compressed Air Consumption - ITS System Operation

Vessel	# of Spargers	Sparger Mode	Total Sparge Air (SCFM)	# of PJMs	Peak PJM Air (SCFM)	Cycle Ave PJM Air (SCFM)	Total Air Required (SCFM)
UFP-VSL-00002A	16	ON	394	6	5016	1140	1534
UFP-VSL-00002B	16	ON	394	6	5016	1140	1534
HLP-VSL-00027A	36	ON	1449	8	5936	1288	2737
HLP-VSL-00027B	36	ON	1449	8	5936	1288	2737
HLP-VSL-00028	36	ON	1745	8	5936	1288	3033

Open Items

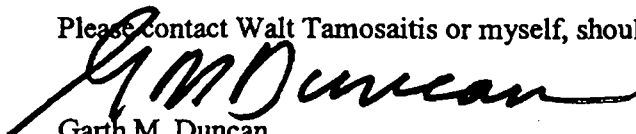
There are several additional actions that are being taken to complete closeout of this technical issue. These key items are summarized below:

- Post-Design Basis Event (DBE) Operating Requirements - Intermittent mixing requirements, ITS air demand, and required emergency diesel generator capacity will be defined separately, and will be included in the system description.
- Operation and Controls - Detailed operating and control modes for spargers and PJMs, and associated vessel level controls will be defined separately, and will be included in the system description.
- R&T Test Reports - These key project documents are in preparation by R&T, supported by Battelle and Savannah River Technology Center. A separate schedule has been developed to track and status these deliverables.
- Additional R&T Testing - There is some additional testing to be performed, primarily to support Engineering in developing a complete operating sequence for ITS mixing which minimizes compressed air demand.
- Vessel drawings - Several drawings are in preparation by Central Engineering, to support PT Plant Design layout of the sparge and PJM air supply piping. The LS and Blend vessel drawings should be available March 24, 2004 by close of business, with the Ultrafiltration Process System (UFP) vessel drawings available by March 26, 2004 close of business.
- Heat Transfer - Vessel heat transfer capacity is under evaluation and will be the subject of separate correspondence.

SUMMARY

The foregoing provides interim definitive design direction regarding Non-Newtonian vessel mixing. This has been determined to be sufficient to support near-term physical design activities for the PT facility.

Please contact Walt Tamosaitis or myself, should you have any questions.



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CEC/cec/cd

CONCURRENCE:



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Manager, Research & Technology
Operations

085014

Distribution

Anderson, Steve	MS5-K.1	Julyk, John	MS8-B
Barnes, Steve	MS1-B	Lynch, Steve	MS4-A2
Beckman, Al	MS4-A1	PDC	MS11-B
Chiaramonte, Gerry	MS9-A	Tamosaitis, Walt	MS1-B
Corriveau, Clarence	MS6-P2	Tosetti, Rich	MS4-A2
Duncan, Garth	MS4-D2	Voke, Robert	MS9-A
Garrett, Richard	MS4-B1	Wilson, Jim	MS12-B
Hoffmann, Mark	MS4-D2		