

BNL-64931

CONF-9709138--

TREATABILITY STUDY ON THE USE OF BY-PRODUCT SULFUR IN KAZAKHSTAN
FOR THE STABILIZATION OF HAZARDOUS AND RADIOACTIVE WASTES¹

BY

RECEIVED
DEC 03 1997
OSTI

P. D. Kalb, L. W. Milian
Environmental and Waste Technology Center
Brookhaven National Laboratory
Upton, N.Y. 11973-5000

S. P. Yim
Korea Atomic Energy Research Institute

R. S. Dyer, W. R. Michaud
U. S. Environmental Protection Agency

This work was performed under the auspices of the U.S. Environmental Protection Agency under Contract No. DE-AC02-76CH00016.

¹Presented at the American Chemical Society's Emerging Technologies in Hazardous Waste Management IX, Pittsburgh, PA, September 15-17, 1997

MASTER

DISTRIBUTION OF THIS DOCUMENT IS UNLIMITED

W

DISCLAIMER

This report was prepared as an account of work sponsored by an agency of the United States Government. Neither the United States Government nor any agency thereof, nor any of their employees, makes any warranty, express or implied, or assumes any legal liability or responsibility for the accuracy, completeness, or usefulness of any information, apparatus, product, or process disclosed, or represents that its use would not infringe privately owned rights. Reference herein to any specific commercial product, process, or service by trade name, trademark, manufacturer, or otherwise does not necessarily constitute or imply its endorsement, recommendation, or favoring by the United States Government or any agency thereof. The views and opinions of authors expressed herein do not necessarily state or reflect those of the United States Government or any agency thereof.

DISCLAIMER

Portions of this document may be illegible in electronic image products. Images are produced from the best available original document.

ABSTRACT

The Republic of Kazakhstan generates significant quantities of excess elemental sulfur from the production and refining of petroleum reserves. In addition, the country also produces hazardous, and radioactive wastes which require treatment/stabilization. In an effort to find secondary uses for the elemental sulfur, and simultaneously produce a material which could be used to encapsulate, and reduce the dispersion of harmful contaminants into the environment, BNL evaluated the use of the sulfur polymer cement (SPC) produced from by-product sulfur in Kazakhstan. This thermoplastic binder material forms a durable waste form with low leaching properties and is compatible with a wide range of waste types. Several hundred kilograms of Kazakhstan sulfur were shipped to the U.S. and converted to SPC (by reaction with 5 wt% organic modifiers) for use in this study. A phosphogypsum sand waste generated in Kazakhstan during the purification of phosphate fertilizer was selected for treatment. Waste loadings of 40 wt% were easily achieved. Waste form performance testing included compressive strength, water immersion, and Accelerated Leach Testing.

INTRODUCTION

Chevron Oil Corp. together with the Kazakhstan Ministry of Oil and Gas has formed a partnership (Tengizchevroil) to explore, develop, and market oil and gas reserves in the Tengiz region of Kazakhstan on the northern shore of the Caspian sea. The growing oil industry in Kazakhstan is already one of their major industries. However, due to limited resources and the required infrastructure to manage resulting toxic and hazardous wastes, there is a growing potential for environmental consequences arising from oil production, refining operations and other environmentally sensitive industries that will need to be addressed. This project, sponsored by the U.S. Environmental Protection Agency Office of International Activities is investigating potential environmental solutions to foster both commercial and environmental sustainable development in Kazakhstan.

Large quantities of by-product sulfur are currently generated by the cleanup of hydrogen sulfide in the production of petroleum and natural gas at the Tengizchevroil fields in Kazakhstan. Currently about 1,200 metric tons/day of sulfur are generated from processing 60,000 barrels of oil/day, but oil production is expected to grow rapidly. The by-product sulfur has little commercial or social benefit and presently, much of it is disposed as waste. Tengizchevroil has obtained special permission from the Kazakhstan Ministry of the Environment to "block and store" the sulfur by-product, a process in which the molten sulfur is cooled into large solid blocks for storage. But as the volume of sulfur residue increases with increased oil and gas production, this practice may no longer be sound or acceptable. In addition, hazardous oil and gas residuals (e.g., incinerator ash, blowdown solutions), as well as toxic and hazardous wastes generated by other past, currently operating and emerging industries throughout Kazakhstan are produced and require treatment prior to disposal.

This paper evaluates the feasibility of using sulfur polymer, (produced from by-product sulfur in the Tengiz region of Kazakhstan) to encapsulate hazardous and radioactive wastes generated on

site, at other sites in Kazakhstan, and elsewhere in eastern Europe. Phospho-gypsum sand waste generated in Kazakhstan from the production of fertilizers was used for this treatability study. This waste was characterized, encapsulated in sulfur polymer and subjected to selected standardized performance tests (e.g., NRC, ASTM) to evaluate mechanical integrity, durability, and leaching properties.

SULFUR POLYMER ENCAPSULATION

Using techniques developed by the U.S. Bureau of Mines [1,2], by-product sulfur can be successfully converted into a stable, durable alternative to concrete with numerous environmental and commercial applications including stabilization of toxic and hazardous wastes. The sulfur is reacted with an organic oligomer (e.g., dicyclopentadiene) and other polymers to form a thermoplastic material known as sulfur polymer or sulfur polymer cement (SPC) with mechanical properties and chemical durability greater than conventional cement products. Its strength and resistance to harsh chemical environments makes sulfur polymer useful for encapsulation of radioactive, hazardous and mixed wastes, as well as for general construction, paving, piping, and coatings for tanks and pads. Since the processing temperature of SPC is relatively low (melting temperature of 120°C) compared with thermal processes such as vitrification (process temperatures in excess of 1200°C), volatilization of contaminants is minimized or eliminated. The commercial cost of sulfur polymer produced in the U.S. is about \$0.12/lb., but due to the large inventory of waste sulfur and low operating expenses, the anticipated cost in Kazakhstan would be lower.

The application of sulfur polymer for encapsulating radioactive, hazardous, and mixed wastes was developed over the last 10 years at the BNL Environmental and Waste Technology Center under sponsorship of the U.S. DOE. [3,4,5,6,7] The process uses a heated double planetary orbital mixer to heat and melt the waste and sulfur polymer binder to form a homogeneous molten mixture. The mixture is then poured into suitable molds for cooling to a solid monolithic final waste form. Contrary to conventional cement processes, the sulfur polymer encapsulation process does not rely on a chemical reaction for hardening, so it is more compatible with a wide range of waste types and can accept greater volumes of waste. Thus, overall waste volumes for storage and disposal are reduced and treatment costs are significantly decreased.

The process has been successfully applied to a wide range of waste types including evaporator concentrates, ash, and sludges. Improved waste loadings have been achieved while still exceeding waste form performance standards specified by the Nuclear Regulatory Commission (NRC) and EPA. A process flow diagram of the sulfur polymer microencapsulation process is shown in Figure 1.

KAZAKHSTAN SULFUR

Twelve drums containing approximately 44 kg. each of by-product elemental sulfur was shipped to the U.S. by Tengizchevroil in Kazakhstan. Six drums of the Kazakhstan sulfur were shipped to McBee and Associates (Lebanon, OR) for conversion to sulfur polymer cement (SPC).

The material was reacted in a 15 gallon stainless steel, steam-jacketed reactor at a temperature of 140°C. The molten sulfur was reacted with a mixture of 2.5 wt% polyester grade dicyclopentadiene and 2.5 wt% of a proprietary reactive polymer manufactured by Exxon Chemical Co. The reaction was conducted for about 4 hours until a viscosity of 50 centipoise, measured at 135°C was achieved. The molten SPC material was then cooled and solidified in 5 gallon steel containers for shipment. Unlike commercially available SPC which is marketed in flaked or granular form, the SPC was received in large blocks from which smaller quantities were chipped off as needed for processing.

Small representative samples of the source sulfur and the SPC produced from Kazakhstan sulfur were analyzed for purity by ACTLABS, Inc., using ICP, INAA, ICP/MS, and XRF techniques. Results are summarized in Table 1. Comparison of carbon content before (<3 ppm) and after conversion to SPC (5 wt%) indicates that the organic polymer modification was accomplished successfully. The source sulfur and the SPC contained a small quantity of chromium which is defined by Environmental Protection Agency (EPA) as a toxic metal. However, leach testing of these materials according to the EPA Toxicity Characteristic Leaching Procedure (TCLP) [8] resulted in concentrations of chromium (0.05 ppm) below the allowable limit (5 ppm) that define a characteristic hazardous waste.

CHARACTERISTICS OF PHOSPHO-GYPSUM SAND WASTE

Analyses of phospho-gypsum sand waste were conducted to determine density, moisture content, acidity in water, particle size, elemental composition, and radionuclides. Density was measured at room temperature using a Multi Pycnometer MVP-1 (Quantachrome Co.). Moisture content was determined at 105°C using a Moisture Analyzer MA30 (Sartorius Co.). Acidity in water was obtained by soaking 1g of the sand into 10ml of demineralized water and measuring the pH of the supernatant water after 24 hours. Particle size distribution was analyzed using ASTM standard sieves according to ASTM C 136-80. After 1 g of phospho-gypsum sand waste was dissolved in 30 ml of acid solution (conc. HCl: conc. HNO₃ = 1:1), elemental composition of dissolved solution were analyzed for various elements by Inductively Coupled Plasma-Atomic Emission Spectroscopy (ICP). Non-dissolved material was filtered and weighed after drying at 110°C. The concentration of phosphorus in dissolved solution was measured using a Hach kit (Hach Co.).

The properties of phospho-gypsum sand waste are summarized in Table 2. The elemental composition of phospho-gypsum sand waste is presented in Table 3. Particle size distribution of waste is shown in Figure 2.

The moisture content of the "as-received" phospho-gypsum sand waste was 12.3 wt%. The phospho-gypsum sand waste was then dried by heating overnight in a convection oven at 110°C, prior to encapsulation in SPC. Dried waste was encapsulated in SPC directly without any size screening or size reduction because the content of relatively large particles (> 2 mm) was less than 4 wt%. The major elements of phospho-gypsum sand waste were calcium, phosphorus, iron, and aluminum. Non-dissolved materials in acid solution might be silicon compounds. Hazardous (toxic) elements are not present in the waste in significant quantities. In addition to analysis for non-

radioactive elements, radionuclides analysis was conducted qualitatively by counting 100g of sample for 24 hours on an intrinsic Ge gamma-spectrometer (with a Canberra computer system). However, no evidence was found for the existence of any radionuclides other than background levels of natural uranium. Although analyses on elements and radionuclides revealed no toxic and radioactive constituents, handling and disposal of this material may provide environmental concern due to its dispersibility.

DETERMINATION OF OPTIMUM LOADING WITH SURROGATE

Only a limited supply of the Kazakhstan phospho-gypsum sand waste was available for this study. Therefore, waste form loading optimization was performed using calcium sulfate reagent as a surrogate for Kazakhstan phospho-gypsum sand waste. Surrogate calcium sulfate was a fine powder of $\text{CaSO}_4 \cdot 1/2\text{H}_2\text{O}$ (calcium hemihydrate). Its density and moisture content were 2.61 and 5.6 wt%, respectively. It was dried at 110°C prior to encapsulation. The density of the dry powder was 2.613.

SPC waste forms were formulated using an electrically heated, stirred mixer by: 1) heating of the SPC at 130°C until the SPC has completely melted, 2) adding dry surrogate to molten SPC and mixing the constituents into a homogeneous slurry by mixer, 3) pouring into a suitable mold and cooling to a monolithic solid. Formulations were prepared with increasing quantities of waste until the limits of processibility were reached.

SPC waste forms containing 10, 15, 20, 30, 40, 50, and 56 wt% surrogate calcium sulfate were prepared according to above mentioned steps. Above 50 wt% surrogate, the molten SPC and surrogate mixture was so viscous that it hardly flowed into the mold. Maximum loading of surrogate determined by the limit of processibility in this experiment was 56 wt%. The molten mixtures containing 30 and 40 wt% surrogate, respectively, had a good fluidity and homogeneity at 130°C. The waste forms were also homogeneous after cooling. In the mixtures containing 10, 15 and 20 wt% surrogate, some of the particles of surrogate seemed to be agglomerated in the melted sulfur. These agglomerates tended to settle during solidification in the SPC as shown in Figure 3.

All waste forms prepared with the surrogate waste were immersed in water and were observed for changes in their physical appearance over a period of 90 days. Results are shown in Figure 4. SPC waste forms containing 50 wt% and 56 wt% surrogate cracked within 2 weeks and 1 week, respectively. The other waste forms maintained their original forms after 90 days.

As a result of processibility and immersion tests, optimum loading of surrogate calcium sulfate was determined at the range of 30-40 wt% in the SPC waste form.

WASTE FORM PERFORMANCE

Sample Preparation

Based on determination of optimum loading with surrogate, SPC waste forms containing 40 wt% Kazakhstan phospho-gypsum sand waste were prepared by the same steps and conditions described above. The molten mixture of SPC and waste was poured into a PVC tube measuring 1.6" in diameter by 3' in height. The same molten mixture was also poured into another PVC tube measuring 1" in diameter by 1' in height. After cooling to ambient temperature, they were cut into specimens measuring 1.6" in diameter by 3.2" in height and 1" in diameter by 1" in height, respectively, within 7 days. Eight specimens measuring 1.6" in diameter by 3.2" in height and five specimens measuring 1" in diameter by 1" in height were obtained. Of the eight specimens measuring 1.6" in diameter by 3.2" in height, four specimens were used in the compressive strength testing and another three specimens were used in water immersion testing. The last specimen was cross-sectioned for observation. Three of five samples measuring 1" in diameter by 1" in height were used in an Accelerated Leach Test (ALT) and the other two samples were archived. Also, SPC waste forms containing 40 wt% surrogate calcium sulfate were prepared in the same fashion. Compressive strength testing, water immersion testing, and ALT were performed on SPC waste form specimens containing surrogate as well as on SPC waste form specimens containing actual phospho-gypsum sand waste.

Homogeneity

A thorough homogenization of waste within the solidified waste form is desirable so that the waste form will exhibit uniform performance characteristics. As a measure of homogeneity, the apparent density of the waste form was obtained for all specimens by dividing the weight of the waste form by the volume calculated from the form dimensions. Apparent densities for the specimens measuring 1.6" in diameter by 3.2" in height are presented in Table 4.

SPC waste form specimens containing phospho-gypsum sand waste have similar apparent densities of approximately 1.89, regardless of their level in the mold. Waste form specimens containing surrogate have almost the same apparent density of 2.06 at any level within two molds. This is indicative of good homogeneity within SPC waste form specimens. The insides of SPC waste forms containing phospho-gypsum sand waste and surrogate calcium sulfate are shown in Figure 5. Particles of phospho-gypsum sand and surrogate calcium sulfate are dispersed homogeneously in SPC. No aggregation and settling were observed in the samples.

Compressive Strength

Compressive strength testing was performed in accordance with the standard method ASTM C-39 (Compressive Strength of Cylindrical Concrete Specimens). [9] Results are summarized in Table 5.

The neat SPC forms containing no waste tested for this study had a mean compressive strength of 3,616 psi. This compressive strength is higher than those of neat SPC forms (approximately 1,800-2,600 psi) previously reported [3, 10, 11]. While the compressive strength of the waste form containing 40 wt% surrogate increased to 5,134 psi by the addition of surrogate to the SPC, the compressive strength of the SPC waste form containing phospho-gypsum sand waste was 3,470 psi, similar to that of neat SPC form. After compressive testing, the insides of the SPC waste forms containing 40 wt% phospho-gypsum sand waste and 40 wt% surrogate, respectively, are shown in Figure 6. The SPC waste form containing phospho-gypsum sand waste had more voids than that of surrogate. Voids in the SPC waste form may be decreased by careful sample preparation. If the voids decrease, the compressive strength of the real waste form will increase. Although the compressive strength of the SPC waste form containing phospho-gypsum sand waste was not higher than those of the neat SPC form and the SPC waste form containing surrogate, the SPC waste form containing phospho-gypsum sand waste possessed excellent mechanical integrity.

Water Immersion Testing

Water immersion testing was performed on three replicate specimens for SPC waste forms containing 40 wt% phospho-gypsum sand waste and 40 wt% surrogate calcium sulfate, respectively. Each specimen was immersed completely in demineralized water at ambient temperature (25°C) and was examined periodically. Upon completion of 90 days, they were removed from the water and checked for variation in weight and dimension. Then, compressive strength testing was conducted according to ASTM C-39.

All specimens had negligible changes in weight and dimensions after 90 day water immersion. Their compressive strength were also retained after water immersion. Results are summarized in Table 5. Improvements of compressive strength may be attributed to experimental scatter associated with the limited sample population.

Accelerated Leach Test

Leach testing for the SPC waste form containing phospho-gypsum sand waste was conducted in accordance with Accelerated Leaching Test (ALT) [12, 13], adopted by ASTM as Standard Test Method C-1308-95 [14]. Three replicate specimens were leach tested at 25°C. Each specimen was leached in a leachant volume that was 100 times the geometric surface area of the specimen. In this experiment, 2,900 ml of distilled water was used in leachant for each specimen because each specimen had a surface area of 29 cm². It was changed twice on the first day, and then daily for 11 days. Leachates were analyzed for major elements of phospho-gypsum sand waste, e.g. Ca, Al, Fe, Na by ICP. Phosphorous in the leachate was checked by Hach kit.

Concentrations of Al, Fe, Na, and phosphorous in the leachate were under, or near, the detection limit for all leachates so they were omitted from leach rate calculation. Calcium leach rates were calculated using a computer program that was developed to accompany the Accelerated Leach Test (the ALT program). Figure 7 shows leaching data and modeling curves for three replicate SPC

waste forms containing 40 wt% phospho-gypsum sand.

The release of calcium from the SPC waste form was very slow. Release data were in close agreement with those predicted by diffusion model. The goodness-of-fit value of the diffusion model to the experimental data was less than 1% for all specimens. The average diffusion coefficient of calcium was $2.02 \times 10^{-10} \text{ cm}^2/\text{sec}$. It is interesting to note that release of calcium from the SPC waste form was controlled by diffusion while previous studies indicate the release of calcium from hydraulic cement waste forms is probably controlled by solubility [12]. Table 6 shows the projected cumulative fraction release of calcium from the SPC waste form containing 40 wt% phospho-gypsum sand waste as a function of waste form size after 10 years, 50 years, 100 years, and 300 years, using the average diffusion coefficient of $2.02 \times 10^{-10} \text{ cm}^2/\text{sec}$ at 25°C obtained from this experiment. This indicates clearly that the SPC waste form is stable and releases calcium very slowly. Releases of radioactive and hazardous contaminants are therefore also expected to be low.

SUMMARY AND CONCLUSIONS

Large quantities of by-product sulfur with little commercial or social benefit are generated in Kazakhstan and are currently being treated as waste. Conversion of this material to sulfur polymer will facilitate beneficial use of the material for treatment of hazardous, radioactive and mixed wastes and for various construction applications. By-product sulfur produced at Tengizchevroil oil production facilities in the Republic of Kazakhstan was shipped to the U.S. and was successfully converted to SPC as evidenced by the presence of 5 wt% carbon detected in the SPC product. Small quantities of chromium were also detected in the Kazakhstan sulfur and SPC product (approximately 6 ppm and 12 ppm, respectively), but neither material failed TCLP testing for hazardous waste.

The Kazakhstan SPC was then used to conduct a treatability study for the encapsulation of phospho-gypsum waste generated in Kazakhstan as a result of fertilizer production. Process development activities resulted in maximum waste loadings of 56 wt%, but optimal mixing and pouring was limited to mixtures containing 40 wt% waste. At 40 wt% waste loading, homogeneity was demonstrated by the close agreement of apparent density measurements throughout the waste forms. Performance testing of SPC waste forms included compression, water immersion, and accelerated leaching. Compressive strength for neat Kazakhstan SPC forms was about 1.5 times greater than previous data for domestic SPC forms. The compressive strength of SPC waste forms containing phospho-gypsum sand waste was 3,470 psi, similar to that of neat SPC forms. Although the compressive strength of the SPC waste forms containing phospho-gypsum sand waste was not higher than those of the neat SPC forms and the SPC waste forms containing surrogate, the SPC waste forms containing phospho-gypsum sand waste possessed excellent mechanical integrity. Waste forms containing 40 wt% waste did not show any deterioration under saturated conditions in water immersion. Their compressive strength were also retained after water immersion. Accelerated Leach Testing revealed diffusion is the dominant leaching mechanism and projected leaching for a full size waste form (2 m diameter x 2 m in height) is $< 5\%$ after 300 years.

REFERENCES

1. Sullivan T. A., and W. C. Mcbee, "Development and Testing of Superior of Sulfur Concretes," RI-8160, Bureau of Mines, U.S. Dept. of the Interior, Washington, DC, 1976
2. Mcbee, W. C., T. A. Sullivan, and B. W. Jong, "Modified Sulfur Cements for Use in Concretes, Flexible Pavings, Coatings and Grouts," RI-8545, Bureau of Mines, U.S. Dept. of the Interior, Washington, DC, 1981
3. Kalb P. D., and P. Colombo, "Modified Sulfur Cement Solidification of Low-Level Waste, Topical Report," BNL-51923, Brookhaven National Laboratory, NY, October 1985
4. Kalb P. D., J. H. Heiser III, and P. Colombo, "Modified Sulfur Cement Encapsulation of Mixed Waste Contaminated Incineration Fly Ash," Waste Management Vol. 11, Number 3, pp.147-153, Pergammon Press, New York, 1991
5. Adams J. W., and P. D. Kalb, "Thermoplastic Stabilization of a Chloride, Sulfate, and Nitrate Salts Mixed Waste Surrogate," Proceedings of the I&EC Special Symposium, Emerging Technologies for Hazardous Wastes, American Chemical Society, Atlanta, GA, September 19-20, 1995
6. Kalb P. D., J. W. Adams, M. L. Meyer, and H. H. Burns, "Thermoplastic Encapsulation Treatability Study for a Mixed Waste Incinerator Off-Gas Scrubbing Solution," Stabilization and Solidification of Hazardous, Radioactive, and Mixed Wastes, ASTM STP1240, T. M. Gilliam, and C. C. Wiles, eds., American Society for Testing and Materials, Philadelphia, PA, 1995
7. Kalb P. D., L. W. Milian, A. J. Grebenkov, and S. P. Rutenkroger, "Thermoplastic Process Treatability for Contaminated Hearth Ash from the Republic of Belarus," Proceedings of the 1996 International Conference on Incineration and Thermal Treatment Technologies, Savannah, GA, May 6-10, 1996
8. USEPA, "Toxicity Characteristic Leaching Procedure (TCLP)," 40 CFR 261, Appendix II, 55 FR 11863, U. S. Environmental Protection Agency, Washington DC, March 29, 1990
9. ASTM, "Standard Method of Test for Compressive Strength of Cylindrical Concrete Specimen," ASTM C39-72, American Society for Testing and Materials, Philadelphia, PA, 1975
10. Heiser III J. H., Pietrzak R., and Colombo P., "Durability of Incineration Ash Waste Encapsulated in Modified Sulfur Cement," Proceedings of the 1991 Incineration Conference, Knoxville, TN, May 13-17, 1991
11. Kalb P. D., J. H. Heiser III, and P. Colombo, "Modified Sulfur Cement Encapsulation of Mixed Waste Contaminated Incineration Fly Ash," Waste Management Vol. 11, Number 3, 1991
12. Fuhrmann M., R. F. Pietrzak, J. Heiser III, E. M. Franz, and P. Colombo, "Accelerated Leach Test Development Program," BNL-52270, Brookhaven National Laboratory, Upton, NY, 11973, 1990.
13. Fuhrmann M., R. F. Pietrzak, J. Heiser III, E. M. Franz, and P. Colombo, "User's Guide for the Accelerated Leach Test Computer Program," BNL-52267, Brookhaven National Laboratory, Upton, NY, 11973, 1990.
14. ASTM, "Standard Method of Test for Accelerated Leach Test for Diffusive Releases from

Solidified Waste and a Computer Program to Model Diffusive, Fractional Leaching from Cylindrical Waste Forms," ASTM C1308-95, American Society for Testing and Materials, West Conshohocken, PA, 1996

Table 1 Chemical Analysis of Kazakhstan Sulfur and Sulfur Polymer

Element	Concentration in Elemental Sulfur, ppm	Concentration in Sulfur Polymer, ppm
Ag	<0.5	<0.5
Al ₂ O ₃	300	300
As	<1	<1
Au	<0.001	8
Ba	7	6
Be	<2	<2
Bi	<5	<5
Br	<0.5	<0.5
CaO	<100	<100
Cd	<0.5	<0.5
Co	0.8	1.1
Cr	6.9	11.9
Cs	<0.2	<0.2
Cu	5	6
Fe ₂ O ₃	300	200
Hf	<0.2	<0.2
Ir	<0.001	<0.001
K ₂ O	<100	<100
MgO	100	400
MnO	<100	<100
Mo	<2	<2
Na ₂ O	<100	<100
Ni	3	3
P ₂ O ₅	<100	<100

<Continued>

Element	Concentration in Elemental Sulfur, ppm	Concentration in Sulfur Polymer, ppm
Pb	<5	<5
Sb	<0.1	<0.1
Se	<0.5	<0.5
SiO ₂	1,400	2,700
Sr	<1	<1
Ta	<0.3	<0.3
TiO ₂	<100	<100
V	<2	<2
W	<1	<1
Y	<2	<2
Zn	5	7
Zr	3	3
Sc	<0.01	0.01
La	0.2	0.2
Ce	1	2
Nd	<1	4
Sm	0.03	0.05
Eu	<0.05	<0.05
Tb	<0.1	<0.1
Yb	<0.05	0.06
Lu	<0.01	<0.01
U	<0.1	<0.1
Th	<0.1	0.1
C	<3	50,020

Table 2 Properties of phospho-gypsum sand waste

Property	Value
Density, g/cm ³ ^{a)}	2.57
Moisture Content, wt%	12.3
Acidity in Water, pH	5.1
Non-Dissolved Materials in Acid Solution, wt% ^{b)}	7.4

a) Density measured by Multi-Pycnometer after drying at 105°C

b) Acid solution (conc. HCl:conc. HNO₃ = 1:1)

Table 3 Elemental composition of phospho-gypsum sand waste

Element	Concentration, ppm
Al	4,754
Ba	210.4
Ni	8.9
Fe	5,390
Cr	5.1
Zn	27.3
Cu	32.7
Cd	-
Ca	45,770
Sr	2,515
Mg	377.6
Na	1,703
K	2,512
Rb	598.1
As	103.6
Se	9.7
Ti	52.4
Mn	24.5
Sb	68.8
Mg	421.3
Mo	7.2
Pb	1.0
V	1.2
P	9,350

Table 4 Apparent densities of SPC waste forms containing 40 wt% waste

Sample Number	Apparent Density of SPC Waste Form, g/cm ³	
	Phospho-Gypsum Sand Waste	Surrogate Calcium Sulfate
#1	1.908	2.068
#2	1.893	2.052
#3	1.897	2.070
#4	1.925	2.061
#5	1.901	2.056
#6	1.898	-
#7	1.855	-
#8	1.890	-
#1 ^(a)	-	2.051
#2 ^(a)	-	2.062
#3 ^(a)	-	2.060
#4 ^(a)	-	2.066
Mean ^{b)}	1.895 ± 0.017	2.060 ± 0.005

* The larger sample number it has, the closer it was to the bottom of the mold.

** Same sample number for each waste does not mean that the samples were taken from same level of the mold, but refers to the relative level in the mold for each waste.

a) Replicate specimens obtained using another PVC mold with the mixture in the same batch.

b) Error expressed at the 95% confidence limit

Table 5 Compressive strengths of SPC waste forms

Waste Type and Loading	Compressive Strength before Water Immersion, psi (Mpa)	Compressive Strength after Water Immersion, psi (Mpa)
Neat ^{a) b)}	3,616 (24.9)	-
Phospho-gypsum sand waste, 40 wt%	3,470 ± 287 ^{c)} (23.9 ± 2.0)	3,794 ± 1,680 ^{c)} (26.1 ± 11.6)
Surrogate calcium sulfate, 40 wt%	5,135 ± 632 ^{d)} (35.4 ± 4.4)	6,541 ± 435 ^{e)} (45.1 ± 3.0)

a) Result reflects mean value for two specimens

b) Standard deviation for two specimens : ± 286 psi(± 2.0 Mpa)

Error expressed at 95% confidence limit for two replicate specimens : ± 2,571 psi
(± 17.7 Mpa)

c) Mean value and error expressed at 95% confidence limit for four replicate specimens.

d) Mean value and error expressed at 95% confidence limit for five replicate specimens.

e) Mean value and error expressed at 95% confidence limit for three replicate specimens.

Table 6 Projected cumulative fractional release of calcium from the SPC waste form containing 40 wt% phospho-gypsum sand waste

Dimension of Waste Form	Projected Cumulative Fractional Release, % ^{a)}			
	10 years	50 years	100 years	300 years
57 cm in I. D. x 70 cm in height	2.8	6.2	8.7	14.6
1 m in I. D. x 1 m in height	1.7	3.8	5.3	9.2
2 m in I. D. x 2 m in height	0.86	1.9	2.7	4.6

a) Projection was made by the diffusion model using a diffusion coefficient of $2.02 \times 10^{-10} \text{ cm}^2/\text{sec}$ measured at 25°C.

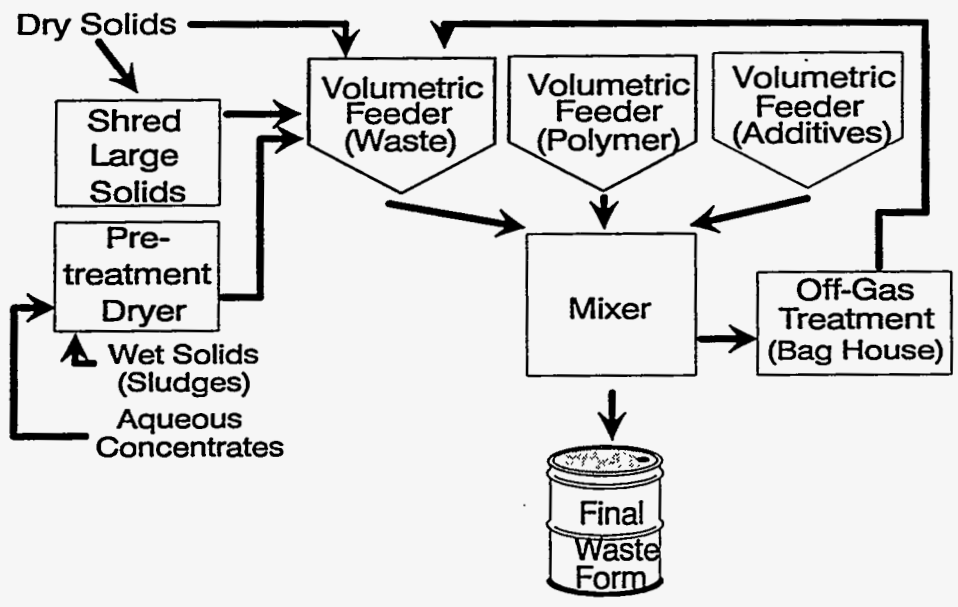


Figure 1 Sulfur polymer encapsulation process flow diagram

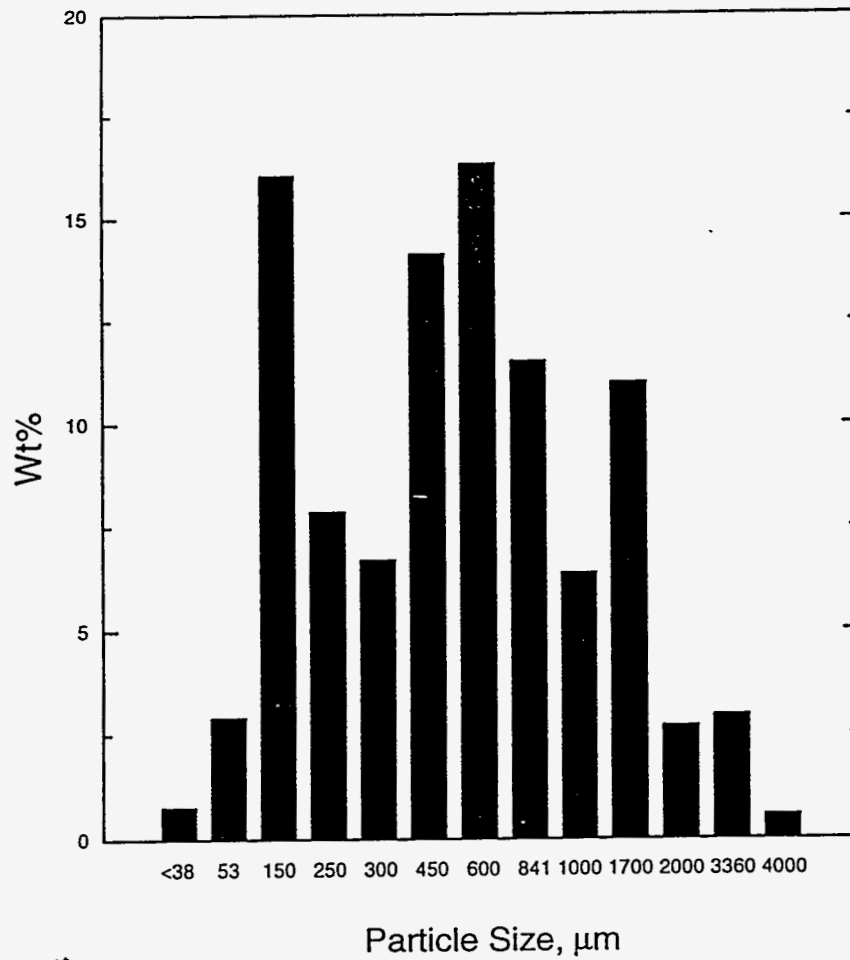


Figure 2 Particle size distribution of phospho-gypsum sand waste

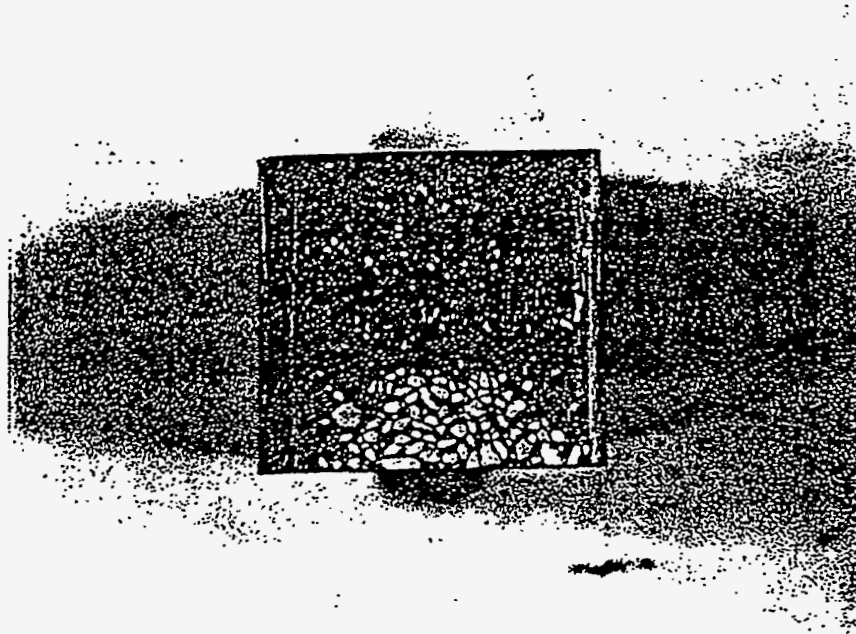


Figure 3 Agglomerates of surrogate calcium sulfate particles and its settling in an SPC waste form containing 20 wt% surrogate calcium sulfate.

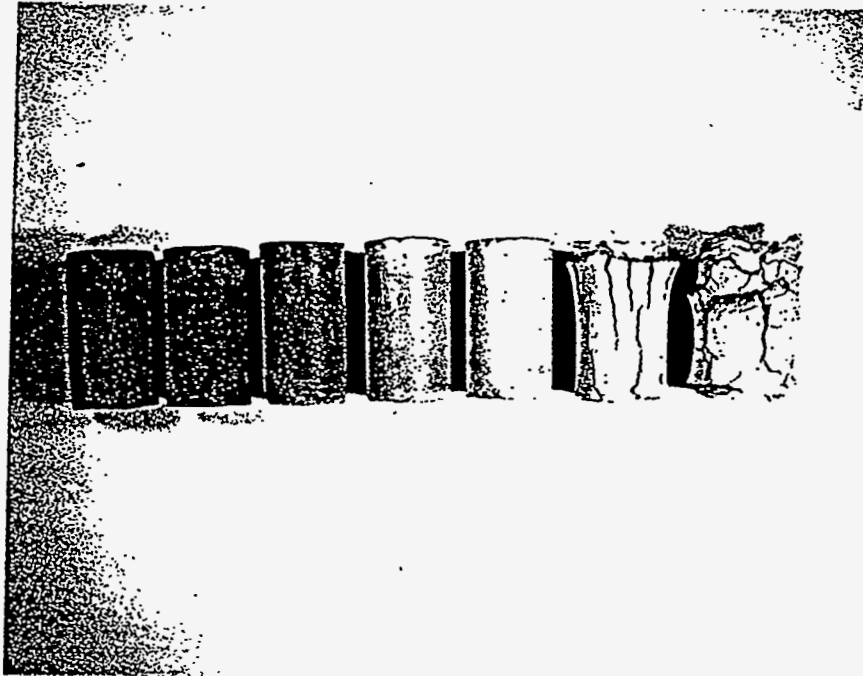


Figure 4 The results of 90 day water immersion test for specimens containing (from left to right) 10 wt%, 15 wt%, 20 wt%, 30 wt%, 40 wt%, 50 wt% and 56 wt% surrogate calcium sulfate, respectively.

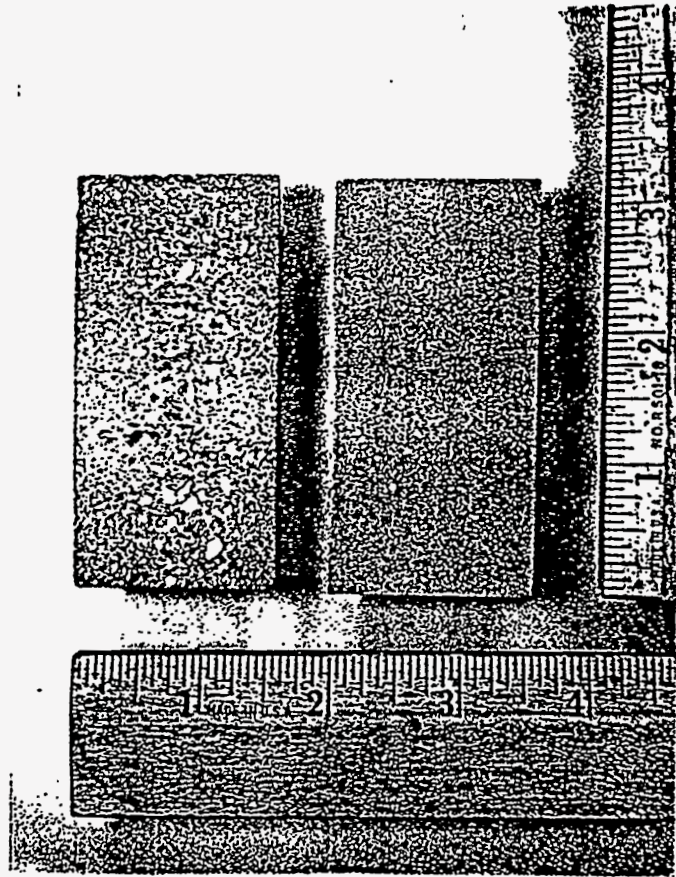
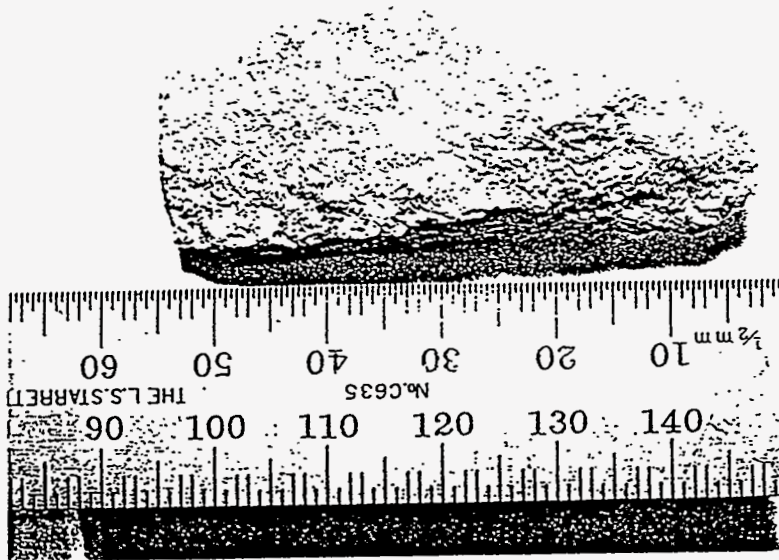
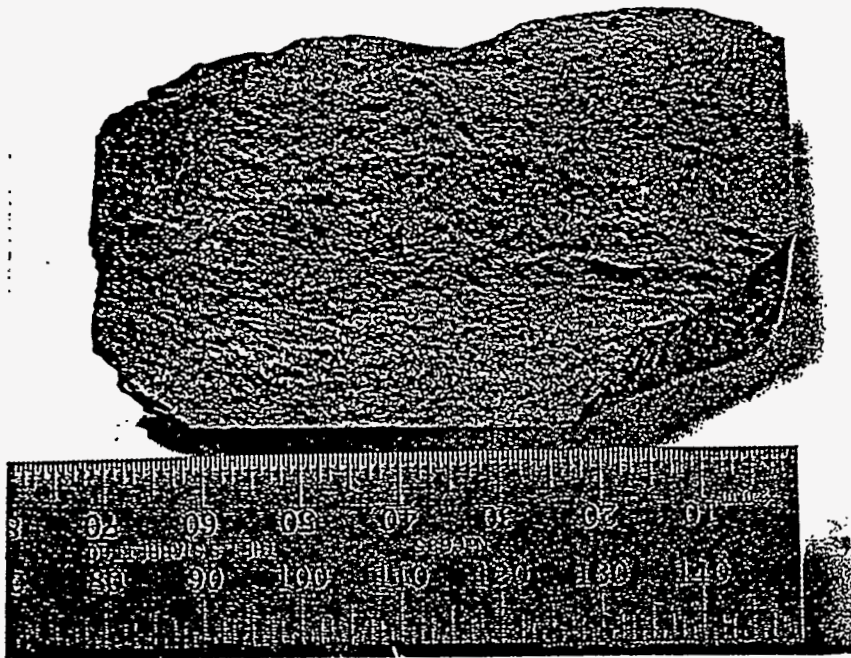


Figure 5 The insides of SPC waste forms containing 40 wt% phospho-gypsum sand waste and 40 wt% surrogate calcium sulfate, respectively.



(a)



(b)

Figure 6 The insides of SPC waste forms after compressive strength testing
(a) An SPC waste form containing 40 wt% phospho-gypsum sand waste
(b) An SPC waste form containing 40 wt% surrogate waste

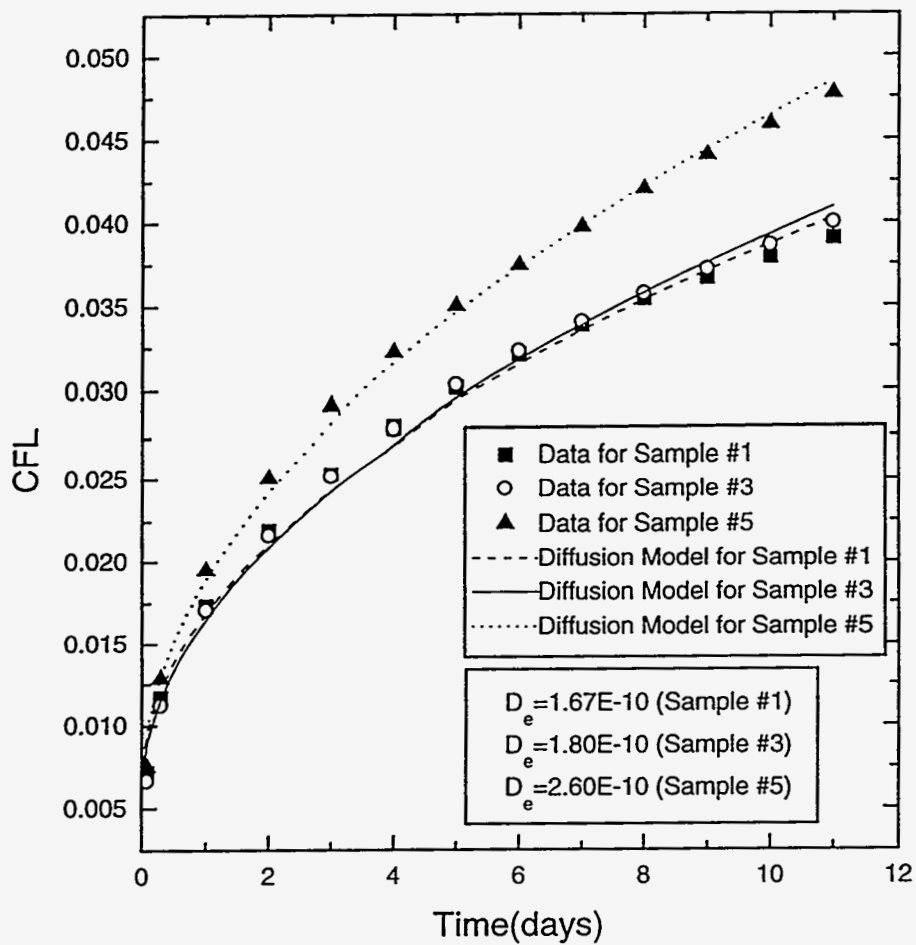


Figure 7 Leaching data points and diffusion model curves for three replicate SPC waste forms containing 40 wt% phosphogypsum sand waste at 25°C