

## **TECHNICAL/ECONOMIC ASSESSMENT OF SELECTED PCB DECONTAMINATION PROCESSES**

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### **Summary**

Eleven emerging alternative treatments for polychlorinated biphenyl (PCB) contaminated sediments have been compared and ranked using technical performance, status of development, test and evaluation data needs, and cost as factors. In ranking the processes, weights were assigned the factors to emphasize the extent of decontamination, the estimated cost of treatment, and the versatility of the process.

The emerging treatment processes are based on six different technologies: one on low-temperature oxidation, two on chlorine removal, one on pyrolysis, three on removing and concentrating, one on vitrification, and three on microorganisms. Types of technologies not developed are chlorinolysis, stabilizing, and enzymes.

On the basis of the comparisons made, the treatment processes were ranked in the following order from highest to lowest: KPEG, LARC, Acurex, Bio-Clean, Supercritical Water, Advanced Electric Reactor, Vitrification, OHM Extraction, Soilex, Composting, and Dybron Bi-Chem 1006. The first eight processes show potential for reduction of PCB concentrations to the desired background levels (1-5 ppm) or less, with minimum environmental impacts and low to moderate cost. All the technologies except the advanced electric reactor required further development and testing.

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### **Introduction**

The PCB contamination problems in the Hudson River and New Bedford, Massachusetts, are reported to be among the worst in the United States in terms of concentration and total quantity of PCBs. It is estimated that 290,000 kg of PCBs are contaminating 382,000 m<sup>3</sup> (500,000 yd<sup>3</sup>) of sediments of the Hudson River. During the 70s, approximately 907,000 kg of PCBs were used in the New Bedford area annually, of which an estimated 45,500 kg were improperly disposed. The PCB contamination problems pose threats to both drinking water and the fishing industry. There are also numerous industrial lagoons contaminated with large quantities of PCBs.

The only available proven technology is dredging and expensive incineration. Land disposal of the sediments untreated has legal restrictions. Biodegradation is a possibility, but sufficient information does not exist to design and operate such a system. There is little experience in the application of encapsulation technology to PCB-contaminated sediments.

This study was undertaken to identify the most technically feasible processes that have been proposed by research concerns for the removal of PCBs from sediments; to identify their extent of development, effectiveness, limitations and probable costs; and to determine needs for further development. The study involved four phases: data acquisition, screening and selection of the most technically feasible processes, development of criteria for process assessment, and process assessment.

### **Data acquisition**

Three major sources of data were: EPA's file of proposals and correspondence concerning problems of PCB contamination and possible approaches to alternative solutions; the open literature; and direct contacts with proponents of treatment technologies.

A reference list was prepared, which included treatment feasibility study reports, process test and evaluation reports, process development proposals, and patents. As processes were identified, direct contacts were made with the investigators for details of their process studies.

### **Screening and selection of most technically feasible processes**

Alternative destruction/detoxification/removal (DDR) processes were subjected to screening to identify those to be assessed further. The processes were categorized according to their generic technology so that their potential performance could be judged appropriately. Processes with undesirable aspects were rejected from further assessment. For example, lack of tolerance for water by a process is undesirable because extensive sediment drying is required. Processes showing insufficient tolerance for water were therefore rejected from further consideration as a primary treatment process in favor of more tolerant alternatives.

Table 1 lists the processes screened, identifies those selected for further assessment and gives the reasons for rejection of the rest. Some of the technologies (e.g. nucleophilic substitution) have provided several processes. Some (e.g. enzymes) have not yet provided any processes. A process evaluated as "1" in Table 1 was selected for further assessment. Other evaluation numbers assigned to the rest of the screened processes refer to footnotes that identify the reason for rejection of the process for further assessment. References cited are identified fully in the reference section.

**TABLE 1**  
**Screening of PCB treatment processes**

Generic technology	References	Process	Evaluation*
<b>CHEMICAL</b>			
<i>General</i>	[1-11]		
<i>Low-temperature oxidation</i>			
Wet air oxidation	[12-14]	Uncatalyzed, general Zimpro Process, Santa Maria, CA Waste Site	2 4, 13
		Catalyzed Dow Chemical Co. Patent 3,984,311	2
		IT Environmental Science	2
Supercritical water oxidation	[15]	Modar	1
Chemical oxidants	[16]	Potassium permanganate plus chromic acid and nitric acid	6
		Chloriodides	4, 7
		Ruthenium tetroxide	3, 4, 8
Ozonation	[17-20]	GE UV/ozonation process	2
<i>Chlorine removal</i>	[21]	Molten aluminum/distillation	14
Dehydrochlorination	[22,23]	Catalytic:	2, 3
		Nickel on kieselguhr	2, 3
		Pd on charcoal	2, 3
		Lithium aluminum hydride	2, 3
		Butyl lithium	2, 3
		Raney nickel	2, 3
Reducing agents	[22,24]	Sodium in liquid ammonia	7, 9
		Nickel-catalyzed zinc reduction	7, 9
		Hydrazine	7, 9
		UV light plus hydrogen	2
		Mildly acidic zinc powder [25]	2, 14
Nucleophilic substitution	[25-36]	Sodium-based processes:	
		Goodyear, sodium in naphthalene (1980)	10
		Acurex, proprietary solvent	10
		PCBX/Sun Ohio	10
		PPM	10
		Ontario Hydro Power	10
		Potassium poly(ethylene glycolate) based:	
		EPA in-house KPEG	19
		KPEG Terrclean-C1	1
		GE KOH-PEG	11
		New York University KPEG	12
Radiant energy	[9,10,37-46]	UV/Photolysis	3
		Syntex photolytic	3, 4
		Thermal corona glow	5
		Microwave plasma	9, 17
		RF in-situ heating	18
		Gamma radiation [40]	9
		LARC	1
Electromechanical reduction	[8]	Electromechanical research process	14
<i>Chlorinolysis</i>	[24]	Hoechst	9
		Goodyear catalytic hydrogenolysis	9
		Exhaustive chlorination	9

TABLE 1 (continued)

Generic technology	References	Process	Evaluation <sup>a</sup>
<i>Pyrolysis</i>	[47-49]	Advanced electric reactor Wright-Malta alkaline catalyst fuel-gas process	1 12
<b>PHYSICAL</b>			
<i>Removing and concentrating</i>	[50-58]		
Heated Air Stripping Extraction		American Toxics Disposal, Inc. Critical Fluid Systems, CO <sub>2</sub> Furfural Acurex solvent wash O.H.M. extraction Soilex process	14 14 15 1 1 1
Adsorption		Carbon adsorption, general Neoprene rubber adsorption	13 15
Vitrification	[59]	Batelle vitrification process	1
<i>Stabilizing</i>	[60-64]	Asphalt with lime pretreatment Z-Impremix Sulfur-asphalt blends (K-20) Ground freezing	16 15 16 13
<i>Bottom recovery</i>	[65-69]	Dredging	13
<b>BIOLOGICAL</b>			
<i>Microorganisms</i>	[70-77]	Bio-Clean Sybron B-Chem 1006 PB Composting Bio-Surf	1 1 1 4, 13
	[48,49,78-80]	Ecotrol, Inc. Wormes Biochemical's Phenobac Rhee anaerobic degradation	4, 13 11, 13 14
<i>Enzymes</i>	[80-82]	No processes found	

<sup>a</sup>Explanation of process rating:

1. Identified emerging sediment treatment process.
2. Destruction efficiency appears to be too low to meet environmental goals.
3. Processing time appears to be extremely long for practical timely cleanup.
4. Data available for dioxin, other chlorinated compounds, or other contaminants, but not PCBs.
5. Process has been shown to destroy PCBs in gas streams only. It may be feasible for sediments, but has not been shown to be.
6. PCBs with 5-7 chlorine atoms per molecule are not destroyed.
7. Products of partial degradation may be toxic.
8. Reagent is very costly/toxic or both.
9. Process costs appear to be excessively high compared with other emerging treatment processes.
10. Water destroys the reagent with its action, thus the process would require excessive drying of sediments and, probably, extraction in pretreatments. The process would therefore have application only as a subordinate final step to several extraction and concentration operations.
11. This particular process was not evaluated because data were not available for assessment.
12. This process is an alternative to another process using the same generic technology, but it is in very early stages of development, and data were not available for assessment.
13. This technique is basically applicable to preliminary operations prior to treatment or to treatment of wastestreams (e.g., wastewaters) from chemical or physical treatments.
14. This process is in the concept stage and data are insufficient to assess it for PCB-contaminated sediments.
15. This process has been found to be ineffective.
16. This technology provides only for encapsulation of the PCB-contaminated sediments.
17. This process supports incineration of PCBs.
18. The process does not appear to be feasible for submerged sediments.
19. Basic data support to identify emerging treatment process.

## Development of criteria for process assessment

The PCB contamination problem in the Hudson River is representative of the type of PCB destruction/detoxification problems focused on in this study. It is expected that the contaminated sediments will have to be dredged from all sites and that the dredged sediments will have high water content.

Criteria for assessment of alternative treatments were chosen which relate to a broad range of principles of operation of diverse applied technologies, yet can be used effectively in comparing one treatment process with another. Additional factors, specific to a technology, were included to help portray the inherent strengths and limitations of a process. Table 2 lists the seven criteria used in comparative process evaluation and three additional factors relating to the needs for further process development and evaluation. The table also includes an overall description of the findings for the processes evaluated.

The goal set for process performance is to reduce the PCB concentration in treated sediments to levels of 1 to 5 ppm. Several of the processes were found to meet this goal. Those that showed reduction to less than 2 ppm were assigned a rating of "6". Those that attained a level between 2 and 10 ppm were assigned a "4". Those with residual concentrations greater than 10 ppm were rated "2".

Available capacity was found not to exist for any of the processes. However, several were developed sufficiently to permit projections of the time required to build a facility for application of the treatment. Those for which such projections could not be made were rated "2". Those requiring 24 or more months were rated "4". Those requiring 12 to 16 months were rated "6".

Conditions/limitations that were rated were tolerance for water, required processing time, and controllability of process conditions. Those treatments that could tolerate water up to about 40 percent would not require a drying step with its attendant fines' control problems. Those requiring only 1 day for treatment could generally show a faster rate of cleanup than those requiring 3 days. Some biological processes required more than 3 weeks. The treatments generally provided control of the processing conditions; however, a few (e.g., composting) would not necessarily do so. The three conditions/limitations were ranked as follows:

<u>Conditions/limitations</u>	<u>Rank</u>
Tolerates to 40 percent water and treats in 1 day	6
Sediment needs to be dried	5
Tolerates to 40 percent water and treats in 3 days	4
Tolerates water and treats in > 3 weeks	3
Sediment needs to be dried, treats in > 3 weeks	2
Processing conditions uncontrollable	1

TABLE 2

## Criteria and technical factors used in process assessment

Criteria/factor	Description
<i>Criteria</i>	
Estimated residual PCB	The goal set for process performance was to reduce the PCB concentration in treated sediments to levels of 1 to 5 ppm. Several of the processes were found to meet this goal.
Available capacity	Although available capacity was found not to exist for any of the processes, several were developed sufficiently to permit projections of the time required to build a facility for application of the treatment.
Conditions/limitations	These included tolerance for water, required processing time, controllability, extent of destruction/decontamination, number of stages of extraction required, and limits on the concentration of PCBs that could be treated. Some processes required one day or less for cleanup; some biological processes required more than 3 weeks.
Concentration range handled	The PCB concentration of the sediments treated ranged from unknown to 3000 ppm. Some processes had limits inherent in the technology.
Status of development	Processes were found to range from concept stage to completed field test stage. Most were in the pilot stage of testing.
Test and evaluation data needs	Data needs varied with the status of the process development. At worst, data were available showing tests of the concept. At best, the process had been field tested, and only permits and checkout were needed.
Estimated cost	The estimated costs of treatments were made in terms of the cost per cubic meter of dry sediment treated, assuming a density of 1.68 Mg/m <sup>3</sup> , plus costs of associated operations — dredging, transportation, handling of treated sediments, as required. All costs are stated in 1985 dollars.
<i>Factor</i>	
Unit operations	The process technology was described, including the active agents, the principles and mechanisms of PCB destruction, and complete characterization of all unit operations.
RCRA waste generated	Some processes have hazardous wastes as residuals from the treatments applied.
Estimated D/D/R efficiency	All the processes achieved a better than 90% destruction/detoxification/removal (D/D/R) efficiency.

Concentration range handled in data developed for the processes ranged from unknown to 3,000 ppm. Ratings were assigned based on the upper limit of feed concentration. The ratings were as follows:

<u>PCB concentration treated, ppm</u>	<u>Rank</u>
≥ 3,000	6
2,000 to 3,000	5
1,500 to 2,000	4
500	3
250 to 350	2
Unknown	1

Status of development ratings were “1” for no data, “2” for laboratory-scale tests completed, “3” for bench-scale tests completed, “4” for pilot-scale tests completed, “5” for field tests completed; and “6” for commercial system designed and ready for construction.

Test and evaluation data needs could be rated differently, depending upon the purpose. For indicating the extent to which a treatment process is readied for use, the more data that are available the better. For indicating the need to support a very promising technology that lacks sufficient progress, the potential and the data needs should be rated in combination. The ratings used here are for the former purpose and are as follows:

<u>Test and evaluation data needs</u>	<u>Rank</u>
None except permits and checkout	6
Field tests	5
Pilot tests and costs	4
Laboratory and bench tests	3
Conceptual treatment process design	2
D/D/R data, residual PCB data, RCRA waste data	1

The application of any treatment process can involve the need for one or more of the following unit operations: dredging, transport, storage, landfill disposal, land treatment disposal, incineration, and/or alternative treatment. Estimates were developed for all of these so that, in any given process evaluation, the proper elements could be added to obtain an estimate of the cost of application. The estimates were made in terms of the cost per cubic meter of sediment treated. The sediment was assumed to have a density of 1.68 Mg/m<sup>3</sup>.

Dredging costs for those treatments requiring removal of the sediment before treatment are estimated at \$20/m<sup>3</sup> based on the recent experience of the U.S. Army Corps of Engineers in contracting for dredging in the New York State area [83].

Transport costs are given as a range. The Corps' experience is \$13/m<sup>3</sup> for

short hauling distances [83]. A cost of \$126/m<sup>3</sup> was used for long hauling distances, which represents an assumed 483-km average transport distance to RCRA landfills capable of accepting PCB-contaminated wastes [84].

Storage cost will sometimes be incurred to hold the dredged sediments pending treatment; e.g., where dredging rates exceed the rates at which the treatment can be applied. These have been set arbitrarily at \$10/m<sup>3</sup>.

Land treatment was used in one of the processes to degrade residual solvent left in the soil after treatment. This involves the controlled application of wastes to the surface of the soil. At land-treatment facilities, wastes are either spread on or injected into the soil, followed by tilling into the soil with farm equipment. The physical and chemical properties of the soil, in unison with the biological component of the soil and sunlight work together to immobilize, degrade, and transform portions of the wastes. The application and tilling process can be repeated many times on the same plot, making land treatment a dynamic system designed to reduce and ultimately eliminate a portion of the waste, as opposed to permanent storage such as landfills.

The American Petroleum Institute [85] has reported that there were 213 land-treatment facilities in operation handling waste from 16 different industry sectors. The most extensive use of land treatment is for petroleum refinery wastes, with 105 land-treatment facilities, many of which are located on the same site as the refinery. More recently, EPA verified the existence of 114 land-treatment facilities and obtained information on operating parameters at some of these sites [86].

Wastes are typically mixed to a depth of 0.5–1.0 ft, where biochemical reactions take place. Application frequencies can range from daily to yearly, with tilling occurring as frequently as daily.

The average cost of controlled, managed land treatment cited by the American Petroleum Institute, \$60/ton, equates to \$111/m<sup>3</sup> of sediments. For short-term land treatment of readily-degradable solvents remaining in treated sediments free of PCBs after they are washed or dried, the cost is estimated at \$33/m<sup>3</sup> [87].

Redeposition costs of decontaminated sediments were also estimated at \$33/m<sup>3</sup>. Slightly lower costs might be expected in special cases.

Because the regulations permit the use of incineration or chemical waste landfill and the application costs of these two methods are available from firms engaging in their practice, these costs were used as lower and upper limits with which to compare the costs of applying new alternative technology.

Landfill disposal costs, incurred when the sediments must be placed in authorized chemical waste landfills, are estimated as ranging from \$260/m<sup>3</sup> for the Michigan area (EPA Regional Office) to \$490/m<sup>3</sup>, based on the highest prices charged for hazardous wastes by commercial facilities [84]. This range includes an intermediate value of \$420/m<sup>3</sup> reported by the Corps of Engineers.

Costs for incineration techniques capable of achieving 99.9999 percent de-



TABLE 3

Unit cost estimates for steps involved in treatment and disposal of PCB-contaminated sediments

Operation	Cost, \$/m <sup>3</sup>
Dredging	20
Transport	13 to 126
Storage	10
Landfill and Disposal	260 to 490
Landfarming	33
Restricted Land Disposal	111
Incineration	1680

struction and removal efficiencies for PCBs are difficult to predict. Even more difficult is prediction of the price commercial facilities will charge to accept the responsibility of handling such a sensitive waste. Surveys made to determine the likely charges to incinerate dioxin-containing wastes resulted in a reported price on the order of \$1,000/Mg [88]. This translates to \$1,680/m<sup>3</sup>, the value adopted for this evaluation, and the cost of disposal of residue from incineration is included. The total cost of use of incineration including dredging at \$20/m<sup>3</sup> and transport at \$13 to \$126/m<sup>3</sup> is \$1713 to \$1826/m<sup>3</sup>.

When available, alternative treatment costs were obtained from the proponent of the process. Otherwise, they were estimated based on the types of unit processes involved and the environmental controls required, or they were determined not to be estimable considering the status of development of the process.

While all costs are in 1985 dollars, the treatment costs are not all necessarily based upon the same labor rates, corporate fixed charges, or profit. These costs vary from one firm to another. The cited estimates are costs of purchasing the treatments. Further costs analyses will be needed to provide a basis for comparison of processes on the basis of individual cost elements.

Table 3 shows the unit cost estimates used to develop cost ranges for the emerging treatments.

Estimated costs were rated by comparing the range of the cost estimates obtained with the cost of placing them into a chemical waste landfill. Treatment processes showing the lowest estimated cost range were rated "6"; those showing a probable cost lower than landfill were rated "4"; those showing an estimated cost equal to landfill were rated "2"; and those showing an estimated cost range greater than landfill were rated "1".

Overall ranking was accomplished through the use of weighting factors assigned to each rated factor. The weighted average rank was then obtained by summing the products of the weighting factors and the ratings and dividing by the sum of the weighting factors. The weighting factors were:

<u>Factor</u>	<u>Weight</u>
Residual PCB concentration	5
Capacity	2
Conditions/limitations	3
Concentration range handled	2
Status of development	2
Test and evaluation data needs	1
Estimated costs	4

The weightings tend to give greatest emphasis to the ability of the treatment to reduce the PCBs and to the probable cost of the treatment. Much less emphasis is placed on the status of development. Thus, an almost fully developed process with an extremely high cost would be ranked lower by application of the weighting process than a less developed process with a much lower potential cost. Test and evaluation data needs have not been heavily weighted because nearly all the alternative treatment processes that show low potential cost require more data to be proven.

Under this procedure, the perfect process for treating PCB-decontaminated sediments would show the following levels for each ranking factor and would receive, using the ratings given, a weighted rating of 6.0:

<u>Factor level</u>	<u>Rating, R</u>	<u>Wt</u>	<u>R × Wt</u>
1. Residual PCB, treated sediment less than 1 ppm	6	5	30
2. Capacity adequate for site cleanup available in 12–16 months	6	2	12
3. Tolerates to 40 percent water and treats in 1 day (24 h)	6	3	18
4. Handles concentrations greater than 3,000 ppm	6	2	12
5. Commercial system designed and ready for construction	6	2	12
6. No test and evaluation data needs except permits and checkout	6	1	6
7. Lowest estimated cost range among alternative emerging technologies	6	4	<u>24</u>
Total $R \times Wt$	$\Sigma R \times Wt$		114
Weighted rating ( $\Sigma R \times Wt$ ) / $\Sigma Wt$			6

### **Process assessment**

The processes were assessed by characterization and ranking. Characterization provided for objective comparison of the processes. Ranking provided a subjective comparison of the processes based on the seven criteria.

## Characterization

Table 4 summarizes five characteristics of the processes: unit operations, available capacity, conditions/limitations, concentration handled, and any generated RCRA wastes. The unit operations employed are given, and each is identified by a number. Generally, a greater number of unit operations will mean a greater effect on treatment costs.

None of the processes has currently available capacity approaching that required for major cleanups. Therefore, the time required to build capacity is listed. Construction time ranges from 12 to 24 months.

Certain conditions that typify the process or limit its versatility are given in column 4 of Table 4. Table 4 also identifies any RCRA waste streams generated by the process.

The data from studies of the processes were examined for ranges of PCB concentrations handled to date. Generally, the values are not limitations on the process, but only on the data acquired. The value  $\leq 300$  ppm for the Bio-Clean process may, however, be a limitation requiring process adjustment to control.

Table 5 lists five additional characteristics of the processes and the rating developed in the ranking process. The characteristics shown here relate to the needs for further process development and evaluation. The process status is given in terms of stages of development completed. The processes range in stages completed from concept to pilot plant.

Both PCB destruction and residual PCB concentration in treated sediments are given to the extent available. Certain processes may require extension of the unit operations, employed (e.g., more stages of extraction) to attain the required performance levels.

Test and evaluation data needs are indicated for each process. Needs vary from none (AER process) to complete site-specific evaluation.

The estimated costs of applying the process are listed in  $\$/\text{m}^3$ . Although cost estimates lack the necessary accuracy at this stage of development of the alternative processes to serve as the sole criterion of potential, they nevertheless indicate that seven of the processes may cost no more than landfill and five could cost less. (Cost estimates could not be made for the Sybron process and composting.)

The processes varied in complexity as evidenced by the number of unit operations employed. Supercritical water oxidation, Bio-Clean and vitrification each employed three unit operations; KPEG employed eight. Operator training requirements were not evaluated comparatively due to insufficient data. However, for the scaled-up treatment processes, the operating labor requirements are expected to be quite similar.

TABLE 4

## Treatment process assessment

Process	Unit operations	Available capacity (or time to provide)	Conditions and limits	Concentration handled	RCRA waste generated
<i>Chemical</i>					
Supercritical water oxidation	1, 4, 10	—	20–40% solids; 374 °C, 23.3 MPa	> 3000 ppm	None
KPEG, Terraclean-CL	1, 3, 4, 7	(24 mo)	organic content > 5% or supplemental fuel 150 °C, 0.5–2 h	500 ppm or greater	w.w.tr. act. carbon
KPEG, NYU	1, 2, 3, 4, 5, 6, 7, 9	—	—	—	—
KPEG, EPA in-house LARC	1, 2, 5, 15	(24 mo)	tolerates 25% water	480 ppm	None
Advanced electric reactor (I.M. Huber)	7, 8, 12, 13, 14	(16 mo)	2204 °C, 2,400 kWh/m <sup>3</sup> needs predryer	> 3000 ppm	None
<i>Physical</i>					
O.H. Materials methanol extraction	2, 7, 8, 14, 15	—	predry to < 1% moisture	> 400 ppm	PCB-loaded carbon from solvent cleanup
“Soilex” kerosene/water	1, 2, 5, 15	—	25% of kerosene solvent retained in soil; 3 d per batch	to 350 ppm tested	Concentrated PCB from still to incineration
Acurex solvent wash	2, 4, 5, 6, 10, 11	—	3–12 washes, tolerates < 40% water	up to 1,983 ppm	Concentrated PCBs to KPEG
Vitrification	8, 12, 14	—	Electrical power usage increases with soil moisture; submerged sediments dredged and treated	500 ppm	None
<i>Biological</i>					
Composting	15, 16	(16 mo)	Seasonal effects, reaction time must be > 4 weeks	1,590 ppm	Treated material is still a RCRA waste
Bio-Clean	1, 2, 17	27 m <sup>3</sup> /d available, 12 mo for full-size	Proved for PCP, laboratory confirmed for PCBs	≤ 300 ppm	None
Sybron Bi-Chem 1006	15, 17	—	Unknown	Unknown	Unknown

Note – Unit operations key:

1. Liquid/solids separation
2. Extraction/solubilization (liquid-solids)
3. Liquid/liquid extraction
4. Chemical reactor
5. Stripping still
6. Solvent recovery still
7. Adsorption
8. Dryer (solids)
9. Dryer (liquids)

10. Filtration
11. Steam cleaning
12. Thermal reactor
13. Grinding
14. Air pollution controls
15. Landfarm
16. Inoculation/digestion
17. V light reactor

TABLE 5  
Treatment process assessment

Process	Status <sup>a</sup>	Estimated D/D/R efficiency, % <sup>b</sup>	Estimated residual PCB, ppm	Test and evaluation data needs	Estimated costs, \$/m <sup>3</sup>	Rating <sup>c</sup>
<i>Chemical/physical</i>						
Supercritical water oxidation, Modar	Field test with PCB liquids	> 99.9995	<0.1 ppb	1, 2, 3, 4, 5, 6, 7	250-733	4.58
KPEG Terraclean-CL	Pilot tests	> 98	<1 ppm	1, 6	208-375	5.42
LARC	Lab tests	> 90	38-50	2, 3, 4, 5, 6, 7	223-336	5.26
Advanced electric reactor	Pilot tests	> 99.9999	<1 ppb	None <sup>d</sup>	830-943	4.58
<i>Physical</i>						
O.H. Materials, methanol extraction	Field tests under way	97	<25 ppm	2, 3, 6, 7	401-514	4.16
Soilex	Pilot tests	95 (3 stages)	6-9 ppm	5, 6, 7	856-913	3.26
Acurex solvent wash	Pilot-scale (field tests planned)	99.9	<2 ppm	Identity of mixed solvent, 6, 7	196-569	5.21
In-situ vitrification Battelle Pacific NW for EPRI	Pilot test of soil	99.9	None in vitrified block, 0.7 ppm in adjacent soil	6	255-548	4.53
<i>Biological</i>						
Composting, aerobic	Lab-scale	62	504-908	4, 5, 6	—	2.47
anaerobic	Lab-scale	18-47	825-1268	4, 5, 6	—	2.47
Bio-Clean, aerobic	Bench-scale	99.99	25 ppb	3, 5, 6, 7	191-370	4.84
Sybron Bi-Chem 1006	Lab-scale and concept	50	—	3, 4, 5, 6, 7	—	1.48

Note - Data needs key: 1. D/D/R data; 2. Residual PCB data; 3. Unit operations data; 4. Bench-scale data; 5. Pilot-scale data; 6. Field test data; 7. Cost data; 8. RCRA waste

<sup>a</sup>Status is defined in terms of the types of studies completed.

<sup>b</sup>D/D/R = destruction/detoxification/removal.

<sup>c</sup>The rating was obtained as shown by the example, under Section Characterization.

<sup>d</sup>AER is fully permitted under TSCA in EPA Region IV for destruction of PCB.

<sup>e</sup>Treatment is continued until a residual of <2 ppm PCBs is obtained.

TABLE 6

Subjective ranking of treatment processes on overall suitability, and estimated cost of application

Process	Cost of application, \$/m <sup>3</sup> treated
KPEG	\$211- 378
LARC	\$223- 336
Acurex solvent wash	\$196- 569
Bio-Clean	\$191- 370
Modar supercritical water	\$250- 733
Advanced electric reactor	\$830- 942
Vitrification	\$255- 548
OHM methanol extraction	\$400- 514
Soilex solvent extraction	\$856- 913
Composting	Unable to estimate cost
Sybron Bi-Chem 1006	Unable to estimate cost
Chemical waste landfill	\$260- 490
Incineration	\$1713-1826

### Ranking of treatment processes

In contrast to process characterization which involves all factors listed in Tables 4 and 5, ranking is subjective and is based on the seven criteria previously described. An attempt was made to define and determine a single number that could represent the overall position of each process relative to an arbitrarily defined perfect process.

Based on the weighted ratings, the processes rank as follows from highest to lowest: KPEG, LARC, Acurex, Bio-Clean, Modar-Supercritical Water, Advanced Electric Reactor, Vitrification, OHM Extraction, Soilex, Composting, and Sybron Bi-Chem 1006 PB/Hudson River Isolates.

### Conclusions

Emerging treatment processes for decontamination of sediments containing PCBs that show potential as alternatives to incineration and chemical waste landfill have been identified. Eleven alternative treatments were compared and ranked using technical performance, status of development, test and evaluation data needs, and cost as factors. The first eight processes show potential for reduction of PCB concentrations to the desired background levels (1-5 ppm) or less, with minimum environmental impacts and low to moderate cost. The sediments must be dredged for application of these treatments.

Of the eleven processes assessed, all but the Advanced Electric Reactor (AER) are in various stages of development for laboratory-scale through field tests. The AER is a permitted treatment under TSCA in EPA Region VI, based

on completed trial burns. There is no immediately available capacity for any of the treatment processes. Further data are needed in most cases to define the final system designs for the processes.

At this stage, estimated costs of application of these eleven processes are less than or within the range of costs of chemical waste landfill, except for the AER estimated cost, which exceeds that of landfill, but is less than incineration. These costs are planning estimates only. In most cases, further research is needed to provide data suitable for more definite cost estimates.

The emerging treatment processes are based on five types of generic technologies: low-temperature oxidation, chlorine removal, pyrolysis, removal and concentration, and microorganisms. Types of generic technologies not yielding competitive emerging processes are: chlorinolysis, stabilization, and enzymes. A search of these technologies yielded no suitable candidate processes at this time.

On the basis of the comparisons made, the treatment processes were ranked in order, from highest to lowest, as shown in Table 6. The estimated cost range (1985 dollars) per cubic meter of sediment treated is also shown for each process. Costs of chemical waste landfill and incineration are given for comparison.

## Notice

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## References

- 1 L.F. Centofanti, Analysis of PCB Chemical Destruction effluents, Div. Environ. Chem., 185th meeting American Chemical Society, St. Louis, MD, April 8-13, 1984, Vol. 24 (11), pp. 11-12.
- 2 K. Chen, The reclamation of transformer oils containing PCBs by the Sunohio PCBX process, Final Report, Prepared for the Tennessee Valley Authority Office of Power, Division of Energy Demonstrations and Technology, Chattanooga, TN, TVA/OP/EDT-83/3, DE83 901683, 1982, 112 pp.
- 3 K. Childs, PCB treatment and destruction technologies, In: Proc. 28th Ontario Industrial Waste Conference, Ottawa, Ontario, Canada, June 13-16, Ontario Ministry of the Environment, Ontario, Canada, 1982, pp. 121-148.
- 4 J.H. Craddock, Polychlorinated biphenyls (PCBs) disposal and treatment technologies: an update, Paper presented at the Fertilizer Institute Environmental Symposium, San Antonio, TX, March 8-10, 1982, p. 161.
- 5 B.H. Edwards, J.N. Paullin and K. Coghlan-Jordan, Emerging technologies for the control of hazardous wastes, EPA-600/2-82-011, NTIS PB 82-236993, U.S. Environmental Protection Agency, Cincinnati, OH, 1982, 145 pp.
- 6 Environment Canada, Proc. Technical Seminar on Chemical Spills, Toronto, Canada, October 25-27, 1983, 291 pp.

- 7 A.W. Hornig, Destruction of PCB-contaminated soils with a high-temperature fluid-wall (HTFW) reactor, In: Proc. 1984 Hazardous Material Spills Conference; Prevention, Behavior, Control, and Cleanup of Spills and Waste Sites, Nashville, TN, April 9-12, Government Institute, Rockville, MD, 1984, pp. 73-79.
- 8 M.J. Massey and F.M. Walsh, An electrochemical process for decontaminating PCB-contaminated transformer coolants, PCB Seminar, EPRI, Seattle, WA, October 22-25, 1985.
- 9 C.J. Rogers and A. Kornel, Laboratory and field tests of chemical reagents for in situ detoxification of chlorinated dioxins in soils, Paper presented at the American Chemical Society National Meeting, Chicago, IL, September 1985.
- 10 C.J. Rogers, Chemical treatment of PCBs in the environment, In: Proc. 8th Annual Research Symposium, Fort Mitchell, KY, March 8-10, 1982, EPA-600/9-83-003, U.S. Environmental Protection Agency, 1983, pp. 197-201.
- 11 C.J. Rogers, Letter to J.B. White, July 19, 1985.
- 12 C.R. Baillod, R.A. Lamparter and D.G. Leddy, Wet oxidation of toxic organic substances, presented at the Purdue Industrial Waste Conference, West Lafayette, IN, 1978.
- 13 R.A. Miller and M.D. Seviontowski, The destruction of various organic substances by a catalyzed wet oxidation process, Report for EPA, Contract 68-03-2568, Work Directive T-7016, no date.
- 14 R.A. Miller and R.D. Fox, Catalyzed wet oxidation of hazardous wastes from detoxification of hazardous wastes, In: J.H. Exner (Ed.), Ann Arbor Science, Ann Arbor, MI, 1982.
- 15 M. Modell, G.G. Gandet, M. Simson, G.T. Hong and K. Blemen, Supercritical Water, Testing Reveals New Process Hold Promise, Solid Waste Management, August 1982.
- 16 FMC Corporation, Industrial waste treatment with hydrogen peroxide, Industrial Chemical Group, 2000 Market Street, Philadelphia, PA, no date.
- 17 R.K. Arisman, R.C. Musick, J.D. Zeff and T.C. Crase, Experience in operation of a ultraviolet-ozone (Ultrox<sup>®</sup>) pilot plant for destroying polychlorinated biphenyls in industrial waste influent, In: Proc. 35th Industrial Waste Conference, Purdue University, West Lafayette, IN, May 11-13, Ann Arbor Science, Ann Arbor, MI, 1981, pp. 802-808.
- 18 W.J. Lacy and R.G. Rice, The status and future of ozone for water and wastewater treatment, *Ind. Water Eng.*, 14(5) (1977) 14-19.
- 19 R.G. Rice and M.E. Browning, Ozone for Industrial Water and Wastewater Treatment, A Literature Survey, Jacobs Engineering Groupe, Washington, D.C.; and Allied Chemical Co., Syracuse, N.Y., for the Environmental Protection Agency, Ada, OK, EPA-600/2-80-060, April 1980.
- 20 H.W. Prengle and C.E. Mauk, New technology: Ozone/UV chemical oxidation wastewater process for metal complexes, organics, and disinfection, *AICHE Symp. Series*, 7 (1978) 228-243.
- 21 United States Patent Office, Destruction of polychlorinated biphenyls and other hazardous halogenated hydrocarbons, U.S. Patent No. 4,469,661, Issued to C.E. Shultz, September 4, 1984.
- 22 N.S. Chu and S.C. Vick, Chemical Destruction of PCBs, PCB Seminar, EPRI, Seattle, WA, October 27, 1985.
- 23 R.B. Lapiere, E. Birin, D. Wu, L. Guezi and W.L. Kranick, Catalytic hydrochlorination of polychlorinated pesticides and related substances, Worcester Polytechnic Institute, Prepared for Municipal Environmental Research Lab, NTIS PB-262804, 1977.
- 24 E.M. Sworzen and D.G. Ackerman, Interim guidelines for the disposal destruction of PCBs items by non-thermal methods, NTIS Publication No. PB82-217498, EPA-600/2-82-069, U.S. Environmental Protection Agency, Washington, DC, 1982.
- 25 K.H. Sweeny and J.R. Fischer, Investigation of means for controlled self-destruction of pesticides, NTIS Publication No. PB-198224, U.S. Environmental Protection Agency, Washington, DC, 1970, 131 pp.



- 26 M.P. Brown, Characterization of Hudson river sediment provided to American Toxics Disposal, Inc. by the New York State Department of Environmental Conservation, Office of Special Projects New York State Department of Environmental Conservation, Albany, New York, 1985, 7 pp.
- 27 D.J. Brunelle and D.A. Singleton, Chemical reaction of polychlorinated biphenyls on soils with poly(ethylene glycol)/KOH, *Chemosphere* 14(2) (1985) 173-181.
- 28 J. March, *Advanced Organic Chemistry*, McGraw-Hill, New York, 1968.
- 29 New York University, Development and evaluation of a low energy process technology for extraction and chemical destruction of polychlorinated biphenyls (PCBs) from contaminated soils and sludges, NYU/DASP 84-02, New York, NY, 1984, 30 pp.
- 30 B. Ruzz, M. Iqbal, W. Brenner and C.J. Rogers, A low energy process for the extraction and chemical destruction of PCBs from contaminated soils and sludges, Paper presented at the American Chemical Society National Meeting, Chicago, IL, September 1985.
- 31 J.G. Smith and L.B. Gurbacham, The use of sodium naphthalenide to chemically destroy polychlorinated biphenyl by dechlorination, Preprint Extended Abstract, Presented before the Division of Environmental Chemistry at the 182nd Meeting of the American Chemical Society, New York, NY, August 23-28, 21 (2) (1981) 88-89.
- 32 Sunohio, PCBX: chemical destruction of PCBs, Brochure, Sunohio, 1700 Gateway Boulevard, SE, Canton, OH 47707, no date.
- 33 United States Patent Office, Methods for decontaminating soil, U.S. Patent No. 4,447,541, Issued to R.L. Peterson, May 8, 1984.
- 34 L. Weitzman, Treatment and destruction of PCB and PCB contaminated materials, Preprint Extended Abstract, Presented before the Division of Environmental Chemistry at the 182nd Meeting of the American Chemical Society, New York, NY, August 23-28, 21 (1981) 42-44.
- 35 L. Weitzman, Cleaning of PCB contaminated soils, Proc. 1983 PCB Seminar, EPRI, Atlanta, GA, December 6-8, 1984.
- 36 L. Weitzman, Solvent cleaning of PCB contaminated soils, Proc. PCB Seminar, EPRI, Seattle, WA, October 22-25, 1985.
- 37 L.J. Bailin and B.L. Hertzler, Development of microwave plasma detoxification process for hazardous wastes - Phase I, NTIS PB-268526, EPA-600/2-77-030, U.S. Environmental Protection Agency, Cincinnati, OH, 1977, 82 pp.
- 38 L.J. Bailin and B.L. Hertzler, Development of microwave plasma detoxification process for hazardous wastes - Part I, *Environ. Sci. Technol.*, 12(6) (1978) 673-679.
- 39 L.J. Bailin, B.L. Hertzler and D.A. Oberacker, Detoxification of Pesticides and Hazardous Waste by the Microwave Plasma Process, ACS Symposium Series, No. 73, American Chemical Society, Washington, DC, 1978, p. 49.
- 40 T.F. Craft, R.D. Kimbrough and C.T. Brown, Radiation treatment of high strength chlorinated hydrocarbon wastes, NTIS PB-244388, EPA-660/2-75-017, U.S. Environmental Protection Agency, Athens, GA, 1975, 33 pp.
- 41 H. Dev, J.E. Bridges, G.C. Stesty and C.J. Rogers, In-situ decontamination of spills and landfills by radio frequency heating, Paper presented at the American Chemical Society National Meeting, Chicago, IL, September 1985.
- 42 E.V. Kalmaz, R.B. Craig and G.W. Zimmerman, Kinetics model and simulation of the concentration variations of the species of polychlorinated biphenyls (PCBs) involved in photochemical transformation, Preprint Extended Abstract, Presented before the Division of Environmental Chemistry at the 182nd Meeting of the American Chemical Society, New York, NY, August 23-28, 21 (2) (1981) 100-103.
- 43 J.M. Meuser and W.C. Weimer, Amine-enhanced photodegradation of polychlorinated biphenyls, Final Report, EPRI-CS-2513, Batelle Pacific Northwest Labs, Richland, WA, 1982.
- 44 J.R. Plimmer, Approaches to Decontamination or Disposal of Pesticides: Photodecomposition, ACS Symposium Series, No. 73, American Chemical Society, Washington, DC, 1978, p. 13.

- 45 J.G. Trump, K.A. Wright, A.J. Sinskey, D.N. Shan and R. Fernald, Disinfection of municipal sludge and wastewater by energized electrons, Report No. INIS-MF-6718, In: Proc. International Seminar on Radioisotopes and Radiation Applied to Environmental Protection, Sao Paulo, Brazil, September 17, 1979.
- 46 P.R. West, S.K. Chaudhary and R.H. Mitchell, Photodechlorination of polychlorinated biphenyls induced by hydroquinone in basic media, Presented before the Division of Environmental Chemistry at the 186th Meeting of the American Chemical Society, Washington, DC, August 28-September 2, 23 (2) (1983) 384-387.
- 47 J.W. Boyd, Thermal Treatment of PCBs and PCB-contaminated Soils, EPRI, PCB Seminar, October 21-25, Seattle, WA, 1985.
- 48 New York State Department of Environmental Conservation, Status Report: PCB biodegradation using Sybron Bi-Chem 1006 PB/Hudson river isolates, Office of Special Projects, Albany, NY, May 21, 1985, 2 pp.
- 49 New York State Department of Environmental Conservation, Status Report: Wright-Malta steam gasification process, Office of Special Projects, Albany, NY, May 17, 1985, 2 pp.
- 50 A.J. Angiola and J.M. Soden, Predicting and controlling downwind concentrations of PCB from surface impoundments, In: Proc. Annu. Meet. Air Poll. Control Assoc., 75(4) (1982) 82.
- 51 R. Caron, Superfund cleanup of PCB contaminated soil, Minden, WV, Personal communication with Ben H. Carpenter, December 1985.
- 52 H. Gilmer and F.J. Freestone, Cleanup of an oil and mixed chemical spill at Dittmer, MO, April-May 1977, In: Proc. 1978 National Conference on Control of Hazardous Material Spills, Miami Beach, FL, April 11-13, Information Transfer, Rockville, MD, 1978, pp. 131-134.
- 53 G.D. Githens, Carbon adsorption onsite for remedial actions, Pollut. Eng., 16(1) (1984) 22-25.
- 54 C.W. Hancher, J.M. Napier and F.E. Kosinski, Removal of PCB from oils and soils, Preprint prepared for submission to: 5th Department of Energy Environmental Protection Conference, November 6-8, Albuquerque, NM, DE85 002619, 1984, 7 pp.
- 55 H.S. Hawthorne, Solvent decontamination of PCB electrical equipment, In: IEEE Conference Record of the Industrial and Commercial Power Systems Technical Conference, Philadelphia, PA, May 10-13, Institute of Electrical and Electronics Engineers, Inc., New York, NY, 1982, pp. 74-78.
- 56 M.C. Lee, R.A. Griffin, M.L. Miller and E.S.K. Chian, Adsorption of water-soluble polychlorinated biphenyl Aroclor 1242 and used capacitor fluid by soil materials and coal chars, J. Environ. Sci. Health, A, 14(5) (1979) 415-442.
- 57 M.B. Saunders, Pilot studies for solvent extraction of PCB from soil, Proc. PCB Seminar, EPRI, Seattle, WA, October 22-25, 1985; see also D.E. Schwinn, D.F. Storrier, R.J. Moore and W.S. Carter, PCB removal by carbon adsorption, Pollut. Eng., 16(1) (1984) 20-21.
- 58 Versar, Inc., Work plan for PCB analyses, Versar Proposal No. 84-989. Submitted to the New York State Department of Environmental Conservation, Albany, NY, June 28, 1984.
- 59 C.L. Timmerman, In situ vitrification of PCB-contaminated soils, Proc. PCB Seminar, EPRI, Seattle, WA, October 22-25, 1985.
- 60 M. Ghassemi and M. Haro, Hazardous Waste Surface Impoundment Technology, J. Envir. Eng., 111(5) (October 1985), pp. 602-617.
- 61 Law Engineering Testing Company, Unsolicited proposal for the application of ground freezing technology to subsurface contamination control and clean-up, Law Engineering Proposal No. 2032.91, 1982.
- 62 F.B. Stroud, R.T. Wilkerson and A. Smith, Treatment and stabilization of PCB contaminated water and waste oil: a case study, In: Proc. 1978 National Conference on Control of Hazardous Material Spills, Miami Beach, FL, April 11-13, Information Transfer, Rockville, MD, 1978, pp. 135-144.

- 63 R.V. Subnamanian and R. Mahalingam, Immobilization of hazardous residuals by encapsulation (Semi-Annual Technical Report), NTIS PB-271410, National Science Foundation, Washington, DC, 1977, 93 pp.
- 64 M.E. Tittlebaum, R.M. Seals, F. Cartledge and S. Engels, State of the art on stabilization of hazardous organic liquid wastes and sludges, CRC Critical Reviews in Environ. Control, 15(2) (1985) 179-211.
- 65 I.G. Carcich and J. Tofflemire, PCB cleanup activities on the upper Hudson River, In: Management of Bottom Sediments Containing Toxic Substances, Proc. 7th U.S./Japanese Experts Meeting, National Technical Information Service, Springfield, VA, 1983, pp. 168-180.
- 66 T.D. Hand and A.W. Ford, The feasibility of dredging for bottom recovery of spills of dense, hazardous chemicals, In: Proc. 1978 National Conference on Control of Hazardous Material Spills, April 11-13, Miami Beach, FL, 1978, pp. 315-324.
- 67 Murakami and Takeishi, Behaviour of Heavy Metals and PCBs in the Removal and Treatment Operation of Bottom Deposits, presented at the Toxic Substances 2nd. U.S./Japanese Experts Meeting, Tokyo, October 25-29, 1976.
- 68 U.S. Army Corps of Engineers, Water Resources Support Center, Management of Bottom Sediments Containing Toxic Substances: Proc. 7th U.S./Japan Experts Meeting, November 2-4, 1981, New York, NY, Cont. Accession No. A136740, 1983, 421 pp.
- 69 T.F. Zimmie and T.J. Tofflemire, Maintenance dredging and toxic substances, In: Proc. 2nd Conference on International Waterborne Transport, ASCE Urban Transportation Division Specialty Conference, New York, NY, October 5-7, 1977, American Society of Chemical Engineers, New York, NY, 1978, pp. 704-719.
- 70 U.S. Bedard, R. Unterman, L.H. Bopp, M.J. Brennan, M.L. Haberl and C. Johnson, Rapid screening assay for PCB-degradative ability, Bio. Sci. Br., General Electric Company, Corporate Research and Development, Schenectady, NY 12301, 1985.
- 71 J.A. Bumpus, M. Tien, D. Wright and S.D. Aust, Oxidation of persistent environmental pollutants by a white rat fungus, Science, 228 (June 20, 1985) 1434-1436.
- 72 R.R. Clark, E.S.K. Chian and R.A. Griffin, Degradation of polychlorinated biphenyls by mixed microbial cultures, Appl. Environ. Microbiol., 37(4) (1979) 680-685.
- 73 I.W. Dawes and I.W. Sutherland, Microbial Physiology, John Wiley and Son, New York, 1976.
- 74 K. Furakawa, Microbial degradation of PCBs, In: A.M. Chakrabarty (Ed.), Biodegradation and Detoxification of Environmental Pollutants, CRC Press, Boca Raton, FL, 1982, pp. 33-57.
- 75 J.D. Isbister, G.L. Anspach and J.F. Kitchens, Composting for degradation of PCBs in soil, In: Proc. 1984 Hazardous Materials Spills Conference; Prevention, Behavior, Control, and Cleanup of Spills and Waste Sites, Nashville, TN, April 9-12, Government Institute, Rockville, MD, 1984, pp. 104-109.
- 76 H.L. Kong and G.S. Sayler, Degradation and total mineralization of monohalogenated biphenyls in natural sediment and mixed bacterial cultures, Appl. Environ. Microbiol., 46 (1983) 666-672.
- 77 D. McCormick, One bag's meat, Biotechnology, 3 (May 1985) 429-435.
- 78 G-Y. Rhee, Letter to R.Mt. Pleasant, May 16, 1985.
- 79 G-Y. Rhee, M. Chen and B. Bush, Anaerobic biodegradation of PCBs, Proposal submitted to the New York State Department of Environmental Conservation, Albany, NY, 1985, 17 pp.
- 80 R. Unterman, D.L. Bedard, L.W. Bopp, M.G. Brennan, C. Johnson and M.L. Haberl, Microbial degradation of polychlorinated biphenyls, Proc. Tenth Conf. on New Frontiers for Hazardous Waste Management, Pittsburgh, Pennsylvania, September 15-18, EPA/600-9-85-025, 1985, pp. 481-488.
- 81 D. Catelani, C. Sorlini and V. Treccani, The metabolism of biphenyl by *Pseudomonas Putida*, Experimentia, 27 (1971) 1173-1174.

- 82 M.L. Rochkind, G.S. Saylor and J.W. Blackburn, Microbial decomposition of chlorinated hydrocarbons, EPA Contract 68-03-3074-3, 1985.
- 83 J. Wheeler, U.S. Army Corps of Engineers, Buffalo District, Contact Summary, Research Triangle Institute, January 15, 1986.
- 84 Industrial Economics, Inc., Regulatory analysis of proposed restrictions on land disposal of certain dioxin-containing wastes, Studies and Methods Branch, EPA, Washington, DC, 1985.
- 85 American Petroleum Institute, Land Treatment - Safe and Efficient Disposal of Petroleum Waste, API, Washington, DC, 1983, 21 pp.
- 86 S. Thorneloe, Land treatment data base, Memorandum to J. Durham, January 31, Environmental Protection Agency, Office of Air Quality Planning and Standards, 1986.
- 87 J. Caplan, Personal communication to Ben H. Carpenter, February 6, 1986.
- 88 Pope-Reid Associates, personal communication to Ben H. Carpenter, March 12, 1986.

### Further readings

- G.P. Adams and R.L. Peterson, Non-Sodium Process for Removal of PCBs from Contaminated Transformer Oil, PCB Seminar, EPRI, Seattle, WA, October 22-25, 1985.
- G. Addis and J. Marks (Eds.), Proc. 1981 PCB Seminar, Dallas, TX, December 1-3, 1981, EPRI EL-2572, Electric Power Research Institute, Palo Alto, CA, 1982, 331 pp.
- R.W. Agnew, Removal and treatment of contaminated river bottom muds: field demonstration, NTIS Publication No. PB84-129022, EPA-600/52-84-006, U.S. Environmental Protection Agency, Cincinnati, OH, 70 pp.
- J. Ball, F. Prizmar and P. Peterman, Investigation of chlorinated and nonchlorinated compounds in the lower Fox river watershed, NTIS PB-292-818/2, EPA-905/3-78-004, U.S. Environmental Protection Agency, Great Lakes National Program Office, Chicago, IL, 1978, 235 pp.
- M.J. Bartos, NYC's plan to meet the water quality challenge, Civ. Eng. (Amer. Soc. Civ. Eng.), 48(11) (1978) 80-84.
- O.W. Berg, P.L. Diosady and G.A.V. Rees, PCB contamination of the aquatic environment in Ontario, Paper presented before the Division of Water, Air, and Waste Chemistry at the 173rd Meeting of the American Chemical Society, August 28-September 1, 12(2) (1972) 59.
- A.K. Bergh and R.S. Peoples, Distribution of polychlorinated biphenyls in a municipal wastewater treatment plant and environs, Sci. Total Environ., 8(3) (1977) 197-204.
- W.N. Billings, T.F. Bidleman and W.B. Vernberg, Movement of PCB from a contaminated reservoir into a drinking water supply, Bull. Environ. Contam. Toxicol., 19(2) (1978) 215-222.
- J.F. Brown, M.E. Lynch, J.C. Carnahan and J. Singleton, Chemical destruction of PCB's in transformer oil, Preprint Extended Abstract, Presented before the Division of Environmental Chemistry at the 182nd Meeting of the American Chemical Society, New York, NY, August 23-28, 21(2) (1981) 90-92.
- M.P. Brown, Letter to C.J. Rogers, May 24, 1985.
- M.P. Brown, Letter of April 16 to Donald L. Wilson, 1986.
- Business Week, The lagging cleanup of great lakes pollution, Bus. Week, May 29 (1978) p. 76.
- I.G. Carcich and T.J. Tofflemire, Distribution and concentration of PCB in the Hudson river and associated management problems, Environ. Int., 7(2) (1982) 73-85.
- L.F. Centofanti, Analysis of PCB chemical destruction effluents, Preprint Extended Abstract, Presented before the Division of Environmental Chemistry at the 185th Meeting of the American Chemical Society, St. Louis, MO, April 8-13, 24(1) (1984) 11-12.
- V.S. Chang and D.A. Fast, PCBs spill at federal pioneer limited's regina plant, In: Proc. Environment Canada 1st Technical Chemical Spills Seminar, Toronto, Canada, October 25-27, 1983, p. 225.
- G. Counsel, Wrench-Lok®: Grounding connections simplified, Paper presented before the Elec-

- trical Equipment Committee at the Pennsylvania Electric Association Fall Conference, Lancaster, PA, September 14, 1982.
- G. Daillaire, Toxics in the N.J. environment: Microcosm of U.S. IL, *Civ. Eng. (Amer. Soc. Civ. Eng.)*, 49(9) (1979) 74-80.
- Electrical World, One solution to the PCB-disposal dilemma, *Elect. World*, 190(9) (1978) 52-53.
- Environment Canada, Destruction technologies for polychlorinated biphenyls (PCBs), Economic and Technical Review Report EPS 3-EC-83-1, Environmental Protection Service, Ottawa, Ontario, July (reprint), 1983, 81 pp.
- T.J. Erler, J. Dragun and D.R. Weider, Two case studies of cost-effective remedial actions for PCB contaminated soils, In: 38th Industrial Waste Conference, Purdue University, West Lafayette, IN, May 11-13, Ann Arbor Science, Ann Arbor, MI, 1983, pp. 369-375.
- J.H. Exner (Ed.), *Detoxification of Hazardous Waste*, Ann Arbor Science Publishers, Ann Arbor, MI, 1982, 362 pp.
- J.W. Fedorko, Protection of the Hosensak 230 kV and Three Mile Island 500 kV capacitor banks, Paper presented at the Pennsylvania Electric Association Fall Conference, Lancaster, PA, September 14, 1982.
- L.L. Fox and N.J. Merrick, Managing polychlorinated biphenyls in the industrial environment - a case history, In: *Toxic and Hazardous Waste: Proc. 15th Mid-Atlantic Industrial Waste Conference*, Lewisburg, PA, June 26-28, Butterworth Publishers, Boston, MA, 1983, pp. 336-344.
- L. Fradkin and S. Barisas, Waste management options for PCBs, In: *Industrial Waste, Proc. 14th Mid-Atlantic Industrial Waste Conference*, College Park, MD, June 27-29, Ann Arbor Science, Ann Arbor, MI, 1982, pp. 398-407.
- M. Gruenfeld, F. Freestone and I. Wilder, EPA's mobile lab and treatment system responds to hazardous spills, *Ind. Water Eng.*, 15(5) (1978) 19-23.
- W.H. Hedley, S.C. Cheng, B.O. Desai, C.S. Smith and H.D. Toy, Alternate treatment of organic solvents and sludges from metal finishing operations: Final Report, NTIS Publication No. PB84-102151, EPA-600/52-83-094, U.S. Environmental Protection Agency, Cincinnati, OH, 1983, 363 pp.
- T.J. Hennings, P.A. Painter, L.L. Scinto and A.M. Takata, Preliminary operations plan and guidelines for the at-sea incineration of liquid PCB wastes, NTIS Publication No. PB83-181834, EPA-600/52-82-968, U.S. Environmental Protection Agency, Research Triangle Park, NC, 1982, 121 pp.
- E. Herbst, I. Scheunert, W. Klein and F. Korte, Fate of PCBs-<sup>14</sup>C in sewage treatment - laboratory experiments with activated sludge, *Chemosphere*, 6 (1977) 725-730.
- L.J. Hetling, T.J. Tofflemire, E.G. Horn, R. Thomas and R.Mt. Pleasant, The Hudson river PCB problem: management alternatives, Paper presented at the New York Academy of Sciences Conference on the Health Effects of Halogenated Aromatic Hydrocarbons, New York, NY, June 24-27, 1978, p. 630.
- A.W. Hornig, Destruction of PCB-contaminated soils with a high-temperature fluid-wall (HTFW) reactor, NTIS PB84-168798, EPA-600/D-84-072, U.S. Environmental Protection Agency, Cincinnati, OH, 1984, 23 pp.
- T.F. Jenkins, D.C. Leggett, L.V. Parker, J.L. Oliphant, C.J. Martel, B.T. Foley and C.J. Deiner, Assessment of the treatability of toxic organics by overland flow, CREL 83-3, U.S. Army Corps of Engineers, Cold Regions Research and Engineering Laboratory, Hanover, NH, 1983.
- J.F. Kitchens, G.L. Anspach, L.B. Mongoba and E.A. Kobylinski, Cleanup of spilled chlorinated organics with the LARC process, In: *Proc. of the 1984 Hazardous Materials Spills Conference, Prevention, Behavior, Control, and Cleanup of Waste Sites*, Nashville, TN, April 9-12, Government Institute, Rockville, MD, 1984, pp. 110-115.
- J.F. Kitchens, W.E. Jones, G.L. Anspach and D.C. Schubert, Light-activated reduction of chemicals for destruction of polychlorinated biphenyls in oil and soil, In: *Symposium on the Detoxification of Hazardous Waste*, Ann Arbor Science, Ann Arbor, MI, 1982, pp. 215-226.

- F.R. Kluge, P.M. Balma, W. Turiansky and R.V. Snow, PJM joint reactive project installation of 230 kV shunt capacitor banks on the public service electric and Gas Company system, Paper presented at the Pennsylvania Electric Association Fall Conference, Lancaster, PA, September 14, 1982.
- L. Kokozska and J. Flood, A guide to EPA-approved PCB disposal methods, *Chem. Eng.*, July 8 (1985) 41-43.
- R.V. Komai, Current PCB research, Presented before the Edison Electric Institute/Envirosphere Company 5th Annual Conference on Environmental Licensing and Regulatory Requirements Affecting the Electric Utility Industry, October 27-29, Washington, DC, 1982, 13 pp.
- R.Y. Komai, D.C. Van der Meer, J. Figler, M.A. Bender, R.P. Kale, M.S. Makar and B.E. Pyatt, Current PCB research, Paper presented at the 5th Conference of the Edison Electric Institute, Envirosphere Environmental Licensing, and the Electric Industry, Washington, DC, October 27-29, 1982.
- A.L. Kopecky, PCB Biodegradation using Sybron Bi-Chem 1006 PB/Hudson River Isolates, Status Report, Sybron Chemicals, Birmingham, NJ, 1985.
- J.P. Lafornera, Cleanup after spills of toxic substances, *Water Pollut. Control. Fed. J.*, 50(4) (1978) 617-627.
- W. Lahey and M. Connor, The case for ocean waste disposal, *Technol. Rev.*, 86(6) (1983) 61-68.
- J.D. Lauber, Burning chemical wastes as fuels in cement kilns, *J. Air Pollut. Control Assoc.*, 32(7) (1982) 771-777.
- J. Ludwigson (Ed.), Proc. of the 1984 Hazardous Material Spills Conference, Prevention, Behavior, Control and Cleanup of Spills and Waste Sites, Nashville, TN, April 9-12, 1984, 445 pp.
- A.D. MaCallum, A dry synthesis of aromatic sulfides: phenylane sulfide resins, *J. Org. Chem.*, 13 (1948) 154-160.
- M.F. McGranaghan and R.F. Gustin, Transient switching studies for EHV shunt capacitor applications on the PJM system, Paper presented at the Pennsylvania Electric Association Fall Conference, Lancaster, PA, September 14, 1982.
- M.F. McGrath, The Fourth Coast, *EPA J.*, 6(5) (1980) 22-24.
- A.E. McIntyre, R. Perry and J.N. Lester, The behavior of polychlorinated biphenyls and organochlorine insecticides in primary mechanical wastewater treatment, *Environ. Pollut. (Series B)*, 21(3) (1981) 223-233.
- K. McManamon, Sunohio's PCBX process, In: Minutes of the Meeting of the Pennsylvania Electric Association, Engineering Section Electrical Equipment Committee, Fall 1982, Lancaster, PA, September 14-15, Pennsylvania Electric Association, Harrisburg, PA, 1982, 9 pp.
- B.W. Mercer, G.W. Dawson, J.A. McNeese and E.G. Baker, Methods/materials matrix of ultimate disposal techniques for spilled hazardous materials, NTIS Publication No. PB85-116853, EPA-600/2-84-170, U.S. Environmental Protection Agency, Cincinnati, OH, 1984, 130 pp.
- N.J. Merrick, L.L. Fox and D.M. Coker, Polychlorinated biphenyl removal by a combined industrial sanitary treatment plants, In: Toxic and Hazardous Waste, Proc. 15th Mid-Atlantic Industrial Waste Conference, Lewisburg, PA, June 26-28, Butterworth, Boston, MA, 1983, pp. 402-415.
- R.C. Millan and N.A. Ostenson, A case study of storage and disposal of PCB contaminated soils, In: Proc. Conference on Hazardous Wastes and Environmental Emergencies: Management, Prevention, Cleanup, and Control, Hazardous Materials Control Research Institute, Silver Spring, MD, 1984, pp. 250-254.
- B. Monick and A. Blake, New industrial wastewater treatment method for removal of multiple contaminants, In: Proc. 70th American Electroplaters Society Annual Technical Conference, Indianapolis, ID, June 27-30, American Electroplaters Society, Winter Park, FL, 1983, 13 pp.
- B.W. Muller, A.R. Brodd and J.P. Leo, Hazardous waste remedial action - Picillo farm, Coventry, Rhode Island; an overview, *J. Hazardous Materials*, 7(2) (1983) 113-129.
- National Institute for Occupational Safety and Health, NIOSH criteria for a recommended stan-

- standard occupational exposure to polychlorinated biphenyls (PCBs), DHEW (NIOSH) Publication No. 77-225, U.S. Department of Health, Education, and Welfare, Washington, DC, 1977, 224 pp.
- New England River Basins Commission, Housatonic river basin overview, NTIS Publication No. PB82-107152, Water Resources Council, Washington, DC, 1980, 209 pp.
- New England River Basins Commission, Housatonic river basin overview, NTIS PB82-107152, Prepared for the Water Resources Council, Washington, DC, 1980, 199 pp.
- H.F. Oeschlaeger, Reactions of ozone with organic compounds, Proc. Ozone Conference, Cincinnati, OH, 1976.
- S.P. Pavlov and W. Horn, PCB removal from the Duwanish river estuary: implications to the management alternative for the Hudson river PCB cleanup, Paper presented at the New York Academy of Sciences Conference on the Health Effects of Halogenated Aromatic Hydrocarbons, New York, NY, June 24-27, 1978, p. 651.
- R.C. Peterson, A.S. Batolomeo, M.H. Corbin and F. Roy, Lehigh electric site superfund PCB cleanup: case history, In: 1983 National Conference on Environmental Engineering, Proc. ASCE Specialty Conference, Boulder, CO, July 6-8, American Society of Chemical Engineers, New York, NY, 1983, pp. 766-774.
- S.A. Peterson and K.K. Randolph (Eds.), Management of Bottom Sediments Containing Toxic Substances: Proc. Second U.S./Japan Experts Meetings, October 1976, Tokyo, Japan, NTIS PB-272684, EPA-600/3-77-083, U.S. Environmental Protection Agency, Corvallis, OR, 1977, 303 pp.
- R. Peterson and E. Milicic, Chemical destruction/detoxification of chlorinated dioxins in contaminated soils, Summary Report (10-84-5-85, sponsored by EPA and Air Force E.56), Galson Research Corporation, E. Syracuse, NY, 1985.
- G-Y. Rhee, Anaerobic biodegradation of PCBs in Hudson river sediments and dredged sediment disposal sites, Proposal submitted to the New York State Department of Environmental Conservation, Albany, NY, 1985, 13 pp.
- R.A. Schact, Pesticides in the Illinois waters of Lake Michigan, NTIS PB-245150, EPA 600/3-74-992, U.S. Environmental Protection Agency, Washington, DC, 1974, 55 pp.
- M.P. Schiaris and G.S. Saylor, Biotransformation of PCB by natural assemblages of freshwater microorganisms, *Environ. Sci. Technol.*, 16 (1982) 367-369.
- R. Scholz and J. Milanowski, EPA Project Summary: Mobile system for extracting spilled hazardous materials from excavated soils, *J. Hazardous Materials*, 9(2) (1984) 241-252.
- T.T. Shen, Estimation of organic compound emissions from waste lagoons, *J. Air Pollut. Control Assoc.*, 32(1) (1982) 79-82.
- R.J. Sloan, K.W. Simpson, R.A. Schroeder and C.R. Barnes, Temporal trends toward stability of Hudson river PCB decontamination, *Bull. Envir. Contam. Toxicol.*, 31 (1983) 377-385.
- Snell Environmental Group, Inc., Rate of biodegradation of toxic organic compounds while in contact with organics which are actively composting: 1982 Final Report, NTIS Publication No. PB84-193150, National Science Foundation, Washington, DC, 1982.
- M.K. Sonksen and J.A. Lease, Evaluation of cement dust stabilization of polychlorinated biphenyl-contaminated sludges, In: Proc. 37th Industrial Waste Conference, Purdue University, West Lafayette, IN, May 11-13, Ann Arbor Science, Ann Arbor, MI, 1983, pp. 405-412.
- T.M. Spittler, Field Measurement of Polychlorinated Biphenyls in Soil and Sediment Using a Portable Gas Chromatograph, ACS Symposium Series, American Chemical Society, Washington, DC, No. 267, 1984, pp. 37-42.
- R. Star, The development of pollution control in Japan. IV. American and Japanese controls of polychlorinated biphenyls (PCBs), *Harv. Environ. Law Rev.*, 1 (1977) 561-567.
- H.J. Streck, Factors affecting the bioavailability of polychlorinated biphenyls (PCBs) in soils, NTIS PB81-209223, Prepared for the Office of Water Research and Technology, Washington, DC, 1980, 109 pp.
- H.J. Streck, J.B. Weber, P.J. Shea, E. Mrozek and M.R. Overcash, Reduction of polychlorinated biphenyl toxicity and uptake of carbon-14 activity by plants through the use of activated carbon, *J. Agric. Food Chem.*, 29(2) (1981) 288-293.

- L.A. Stretz, L.C. Borduin, W.E. Draper, R.A. Koenig and J.S. Vavruska, Controlled air incineration of hazardous chemical waste at the Los Alamos National Laboratory, In: Proc. 1982 Symposium on Waste Management; Waste Isolation in the U.S. and Elsewhere, Technical Programs and Public Communications. Vol. 1: General, Tucson, AZ, March 8-11, Arizona Board of Regents, Tucson, AZ, 1982, pp. 281-299.
- D. Taylor, Managing organics in sludge reuse programs, *BioCycle*, 25(6) (1984) 20-22.
- T.B. Thomason and M. Modell, Supercritical water destruction of aqueous wastes, *Hazard. Waste*, 1(4) (1984) 453-467.
- E.S. Tucker et al., Activated sludge primary biodegradation of PCB's. *Bull. Environ. Contam. Toxicol.*, 14 (1975) 705-713.
- M.T.M. Tulp, R. Schmitz and O. Hutzinger, The bacterial metabolism of 4,4'-dichlorobiphenyl and its suppression by alternative carbon sources, *Chemosphere*, 7(1) (1978) 103-108.
- U.S. Environmental Protection Agency, Identification and listing of hazardous waste, 40 CFR 261, Final regulation, *Federal Register*, 1980.
- U.S. Environmental Protection Agency, Polychlorinated biphenyls (PCBs) in manufacturing, processing, distribution in commerce, and use prohibition, 40 CFR 761, Final regulation, *Federal Register*, 1980.
- U.S. Environmental Protection Agency, Assessment of incinerators as a treatment method for liquid organic hazardous wastes: summary and conclusions, Washington, DC, March 1985, 5 pp.
- U.S. Environmental Protection Agency, and Oil Spill Control Association of America, Control of Hazardous Material Spills; Proc. 1978 National Conference in Control of Hazardous Material Spills, Miami Beach, FL, April 11-13, 1978, 458 pp.
- United States Patent Office, Method and apparatus for treating polychlorinated biphenyl contaminated sludge, U.S. Patent No. 4,402,274, Issued to W.C. Meenan and G.D. Sullivan, September 6, 1983.
- E. v. Löw, Vorkommen und Mikrobieller Umund Abbau von aromatischen Polyzyklen im Boden und in Siedlungsabfällen, *Forum Städte-Hygiene*, 34(5) (1983) 263-267.
- W.C. V. Meyer, A study of thionation reactions for use in destruction of toxic wastes, Initial tests of thionation reaction conditions with priority pollutants, Fairview Industries, Inc., Middleton, WI, July, EPA Contract No. 68-02-3992, 1986.
- Versar, Inc., Assessment of wastewater management, treatment, technology, and associated costs for abatement of PCB concentrations in industrial influents, 1976.
- K.H. Walker, The great lakes cleanup, *Water Sewage Works*, 124(11) (1977) 85.
- G. Weaver, PCB pollution in the New Bedford, Massachusetts area: A Status Report, Massachusetts Office of Coastal Zone Management, Report submitted to the Massachusetts New Bedford PCB Task Force, June 1982, 62 pp.
- M.D. Webber, H.D. Monteith and D.G.M. Corneau, Assessment of heavy metals and PCBs at selected sludge application sites in Ontario, Research Report NO. 109, Research Program for the Abatement of Municipal Pollution under the Provisions of the Canada-Ontario Agreement on Great Lakes Water Quality, Environment Canada, Environmental Protection Service, Ottawa, Ontario, 1981, 27 pp.
- R.H. West and P.G. Hatcher, Polychlorinated biphenyls in sewage sludge and sediments of the New York Bight, *Marine Pollut. B.*, (11) (1980) 126-129.
- F.C. Whitmore and J.D. Barden, A study of PCB destruction efficiency and performance for a coal-fired utility boiler, Vols. 1 and 2, NTIS Publication Nos. PB84-110147 and PB84-110154, EPA-600/52-83-101a and b, U.S. Environmental Protection Agency, Research Triangle Park, NC, 1983, 79 pp.
- D.R. Young and T.C. Heeson, Polychlorinated biphenyls in the near-shore marine ecosystem off San Diego, CA, Report No. SCCWRP-109, Prepared by the Southern California Coastal Water Research Project for the California Regional Water Quality Board, San Diego, CA, NTIS PB-283090, 1977, 21 pp.
- D.R. Young, D. McDermott-Erich and T.C. Heeson, Sediments as sources of DDT and PCB, *Marine Pollut. Bull.*, 8(11) (1977) 254-257.