DuPont Engineering 4200 Camp Ground Road Louisville, KY 40216

## **QUPOND**

#### **DuPont Engineering**

May 10, 2002

Mr. Christopher A. Saari Hydrogeologist Northern Region Remediation and Redevelopment State of Wisconsin Department of Natural Resources Ashland Service Center 2501 Golf Course Road Ashland, Wisconsin 54806

#### DRAINAGE SAMPLING WORK PLAN Former DuPont Barksdale Works Barksdale, Wisconsin

Dear Mr. Saari:

This letter transmits the Drainage Sampling Work Plan for the former E. I. du Pont de Nemours and Company, Inc. (DuPont) Barksdale Works. DuPont has submitted the applicable fees for your review of this work plan to the Wisconsin Department of Natural Resource's Rhinelander, Wisconsin office (see attached letter and copy of payment).

As we have discussed previously, DuPont desires to implement the sampling on May 17, 2002, pending your review of the work plan and the presence surface water flow within the site's drainage features.

If you have any questions or comments regarding work plan, please contact me at (502) 569-2148.

Sincerely,

Fradling A. Nave

Bradley S. Nave DuPont Corporate Remediation Group Project Director

Mr. Cary E. Pooler, P.G., URS Corporation
Mr. Robert J. Raymond, URS Corporation
Mr. Paul Bretting, Bretting Manufacturing, Inc.
Mrs. Amelia Lindsey, RN, Bayfield County Health Department
Mr. Henry Nehls-Lowe, MPH, State of Wisconsin Department of Health and Family Services
Mr. Doug Shultz, Wisconsin Department of Natural Resources, Ashland Service Center

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DuPont Engineering Barley Mill Plaza - Bldg. 27 Lancester Pike & Rte. 141 Wilmington, DE 19805



May 3, 2002

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DuPont Engineering

Janet Kazda

WDNR 107 Sutliff Avenue Rhinelander, W1 54501

Dear Ms. Kazda:

Enclosed is check in the amount of five hundred dollars (\$500.00) to cover the cost of reviewing a sediment/surface water workplan for the former DuPont Barksdale site.

Please forward a receipt to my attention at ---

DuPont Barley Mill Plaza, 27/2278 4417 Lancaster Avenue Wilmington, DE 19805

Thanks,

hoss

Peggy Moss Administrative Secretary

Enclosure

E I du Pont de Nemours and Company

# DRAINAGE SAMPLING PLAN FORMER DUPONT BARKSDALE FACILITY BARKSDALE, WISCONSIN

Date: May 2002

Project No.: 40-D4BA7431.02



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CORPORATE REMEDIATION GROUP An Alliance between DuPont and URS Diamond

- 16. Sr

Barley Mill Plaza, Building 27 Wilmington, Delaware 19805 Drainage Sampling plan former dupont Barksdale facility Barksdale, Wisconsin

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### **1.0 INTRODUCTION**

This sampling plan describes the proposed scope of work for conducting a screening level characterization and evaluation of sediment and surface water at the former Barksdale Works Site (see Figure 1). Currently, there are eight drainage locations at or near the site boundary. Each will be evaluated as part of this sampling program.

The objective of this sampling program is to determine whether site-related constituents are present in sediment and surface water at drainage locations at or near the former manufacturing site boundaries. The results will be used to prioritize future investigations if deemed necessary.

#### 1.1 Background

Site investigation activities were initiated when groundwater samples collected by the Wisconsin Department of Natural Resources (WDNR) in June 1997 from a drinking water well located at a residence adjacent to the site showed detectable concentrations of 2,4-dinitrotoluene (2,4-DNT) and 2,6-DNT. In response, the WDNR and E.I. duPont de Nemours and Company, Inc. (DuPont) met and agreed upon investigative and remedial objectives for the site.

From 1997 to 2000, various groundwater and residential water sampling events occurred to evaluate groundwater and drinking water quality. Also, in 1997, a surface water and sediment study along Boyd Creek was conducted to evaluate the potential for offsite transport of DNT, trinitrotoluene (TNT) and nitroglycerin (NG) at levels of ecological concern. This investigation concluded that DNT isomers were present in the creek's sediments, but that the concentrations of these constituents were not considered to be a potential concern.

During 2001, an extensive onsite investigation program was conducted. This investigation primarily focused on evaluation of groundwater conditions. However, a screening level evaluation of surface soils in production and operation areas also occurred. These evaluations indicated the presence of site-related constituents that could potentially be eroded by storm water run-off.

Following the 2001 investigation, DuPont developed the following overall site objectives.

- Continue to provide drinking water that meets appropriate Wisconsin regulatory standards for all site-related constituents.
- □ Develop a plan for a long-term drinking water supply and implement as appropriate.
- **D** Evaluate the potential for off-site surface issues and take appropriate action.
- □ Identify priority source areas and take appropriate action.

The sampling program presented in this work plan is the first step in addressing the third objective.

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## 2.0 INVESTIGATION AREAS

As mentioned in Section 1.0, there are eight current drainage locations where surface water and sediment are potentially transported beyond the site boundaries. The point at which each of these drainages approximately exits the site is described below and depicted on Figure 1.

- Location 1: Ditch along the northern site boundary that receives drainage from the site's northern area and proceeds along Nolander Road's southern drainage ditch
- Location 2: Culvert along the northern site boundary that empties into a ditch along the south side of Nolander Road
- Location 3: Culvert along the eastern site boundary that receives drainage from the site's northern area
- □ Location 4: Ditch at the front gate on Highway 13 that receives overland flow
- □ Location 5: Culvert along the eastern site boundary that receives drainage from the site's central area
- □ Location 6: Boyd Creek near Highway 13
- □ Location 7: Mission Springs Creek near Highway 13
- □ Location 8: Culvert along the northeastern site boundary (first culvert going south from the intersection of Nolander Road and State Highway 13)

Locations 6 and 7 are most likely to have continuous flow throughout the year.

### 3.0 FIELD ACTIVITIES

All sampling activities will be conducted in accordance with the Quality Assurance Project Plan (QAPP) dated September 2001. Field activities will be coordinated so that sufficient water flow is present in each drainage location for collection of the required number of samples. In addition, to ensure that samples within each drainage feature are comparable (representative of similar conditions), sampling will be conducted at all eight locations during a single sampling event.

#### 3.1 Investigation Rationale and Methodology

The scope of work was designed to provide flexibility in choosing sample locations and matrices due to the uncertainty that is inherent with this type of sampling program. Samples will be collected approximately at the site boundary when safety issues are not a concern. If safety issues are a concern, every attempt will be made to safely collect samples as close to the site boundary as possible.

#### 3.1.1 Sediment Sampling

The method of sediment sample collection will depend on the depth and velocity of flow at the time of sample collection. If the flow is low and direct access to the drainage bed can be safely achieved, a board or other barrier will be inserted to divert fast flowing water, then a decontaminated stainless steel bowl and stainless spoon will be used to collect sediment samples. If flow is fast or deep, a Teflon sampling dipper (pole mounted beaker) will be used to collect the sediment by dragging the drainage bottom. Based upon field conditions, additional equipment may be used. A complete description of actual sample collection activities will presented in the sampling event's summary report (see Section 4 for additional Report information).

Based upon each drainage location's physical characteristics, sediment may not be present or the volume of sediment present is not adequate to collect representative samples in a drainage location. However, soil may be present. Therefore, soil samples may be collected instead.

Once the sample containers are filled, they will be placed immediately into a cooler and preserved to  $< 4^{\circ}$ C with ice.

If soil samples are collected for volatile organic constituents (VOCs), Encore<sup>®</sup> sampling devices will be used.

#### 3.1.2 Surface Water Sampling

A decontaminated Teflon<sup>®</sup> dipper will be used to retrieve surface water samples from the drainage channels (based upon field conditions, additional equipment may be used to retrieve the samples). Aliquots for whole water analyses will be immediately decanted from the dipper reservoir into the sample containers. For VOC samples, every attempt will be made to minimize disturbance of the water. For dissolved constituent analyses,

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the water will be pumped from the dipper reservoir into the containers using a peristaltic pump outfitted with fresh tubing and an unused 45-micron filter. Once the sample containers are filled, they will be placed immediately into a cooler and preserved to  $< 4^{\circ}$ C with ice.

#### 3.2 Sampling Locations

At each location, a minimum of one sediment/soil sample and one surface water sample will be collected. Sediment samples will be collected if there is sufficient volume present. If the drainage bed does not contain sufficient sediment, soil samples will be collected from the drainage bottom instead. Figure 1 shows the proposed sampling locations. Table 2 summarizes sampling information for each sample location/drainage area.

The physical characteristics of each sampling area (total of eight) will be documented after analytical samples are collected. These characteristics, at a minimum, will include the following.

- □ Water flow (depth, width, velocity), turbidity, temperature and conductivity
- □ Sediment, sheen or other non-aqueous features within the flow channel
- Description and photos of surrounding features and the drainage channel within 100-ft upstream of the sampling point: grass area, along side of a road, etc.

In drainages where the flow channel is well defined and average water depth is greater than 3 inches, velocity will be measured using a Marsh-Mcburney ultrasonic doppler velocity meter or equivalent at spacings of no more than 18-inches across the wetted stream cross-section. Where flow is dispersed or depth is less than the required submergence of the meter, velocity may be estimated using a floating object and a stopwatch. Where the flow is slow and confined to a narrow accessible cross-section (culverts), direct measurement using a bucket and stopwatch will be used rather than velocity measurements.

Turbidity, temperature, dissolved oxygen, conductivity and pH will be measured at the mid-depth in the flow channel using a Horiba U-10 Water Quality meter or equivalent.

Sediment and non-aqueous flow components (foam, sheen, detritus etc.) will be identified by plunging a clean bucket into the center of the stream to the bottom and visually describing the contents after the bucket has been allowed to stand and separate.

#### 3.3 Analytical Methods

#### 3.3.1 Field Parameters

For each surface water sample, the following parameters will be evaluated in the field:

- □ pH
- □ Turbidity
- Dissolved Oxygen (DO)

- Specific Conductivity
- □ Temperature

#### 3.3.2 Laboratory Parameters

All samples will be submitted to Severn Trent Laboratories (STL) for analyses. The following analyses will be performed on all samples.

- □ Metals (surface water will be quantified for total and dissolved metals)
- □ Nitroaromatics/nitramines
- □ Semivolatile organic constituents (SVOCs)
- U VOCs

Table 1 presents the specific constituent lists. Since the purpose of this sampling program is characterization, comprehensive analytical lists were selected to allow a thorough evaluation of sediment and surface water quality. In addition, all nitroaromatic/nitramine analyses will be submitted for independent data validation in accordance with guidance from the *National Functional Guidelines for Organic Data Review*, U.S. EPA, 2/94 and 10/99 updates.

#### Chain of Custody

The laboratory will also initiate the chain-of-custody (COC) at the time the sample containers are prepared for shipment to the site and will accompany the empty containers to the site. The following information will be recorded on the COC prior to shipping samples from the site to the laboratory.

- □ Collector's name
- □ Sample collection dates and times
- □ Sample identification numbers
- □ Number of containers for each sample aliquot (if not already indicated)

□ Container size/type (if not already indicated)

- **Type of preservation (including ice)**
- □ Parameters (analytes from each sample aliquot)
- **u** Turn-around requirements
- Special handling instructions
- Destination of samples
- □ Name, date, time and signature of each individual possessing the samples

The COC will be signed by each individual responsible for custody of the sample containers. The original signed COC will accompany the samples to the laboratory and be returned as part of the final data package to DuPont.

#### 3.3.3 QA/QC Samples

The following QA/QC samples will be collected at the specified frequency to evaluate QA/QC issues.

- □ Trip Blanks: a minimum of one trip blank per shipment of VOC analyses (water samples only)
- **□** Equipment Blanks: a minimum of one equipment blank per day
- □ Field Duplicates: a minimum of one field duplicate per 20 field samples of the same matrix
- Matrix Spike/Matrix Spike Duplicate (MS/MSD) samples: a minimum of one MS/MSD per 20 field samples of the same matrix

Trip blanks will be used to detect outside contamination sources that might influence contaminant concentrations reported in field samples both quantitatively and qualitatively. Possible contamination sources include:

- □ Laboratory reagent water
- □ Sample containers
- Cross contamination during sampling or shipment
- Ambient air or contact with analytical instrumentation during field sampling or analysis at the laboratory
- □ Laboratory reagents used in analytical procedures
- □ Instrument carry-over

Equipment blanks will be used to evaluate the completeness of decontamination procedures in preventing cross contamination of samples by sampling equipment. Field duplicate samples will be used to evaluate the cumulative precision due to limitations of the analytical method, sample matrix, and sample collection techniques. MSs/MSDs will be used to evaluate analytical accuracy and precision.

#### 3.4 H&S/Waste Management

All field activities will be conducted as per the site specific Health and Safety Plan (HASP) and Waste Management Plan (WMP).

#### 3.5 Schedule

Every attempt will be made to collect the samples in a timely manner. Below is a tentative schedule showing activities following sample collection.

- □ Sample Collection
- □ 30 days after Sample Collection Date: Analytical Report submitted to DuPont
- 7 days after Analytical Report submitted to DuPont: Validation Report submitted to DuPont

90 days after Validation Report submitted to DuPont: Report summarizing field activities and data evaluation submitted to WDNR

## 4.0 **REPORTS**

A summary report detailing all sampling activities will be prepared and submitted to the Wisconsin Department of Natural Resources (WDNR) for review. The report will present, at a minimum, the following information.

- Description of all field activities, data evaluation results, and conclusions/recommendations.
- □ Tables summarizing all data.
- **D** Photo documentation of the sampling locations and drainage features
- □ Laboratory report.
- □ Maps presenting pertinent data.

## TABLES

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Metals	
Antimony (SW-846 6010B)	Mercury (SW-846 7470/71B)
Arsenic (SW-846 6020)	Nickel (SW-846 6010B)
Barium (SW-846 6010B)	Selenium (SW-846 6020)
Cadmium (SW-846 6010B)	Silver (SW-846 6010B)
Chromium (SW-846 6010B)	Thallium (SW-846 6020)
Cobalt (SW-846 6010B)	Tin (SW-846 6010B)
Copper (SW-846 6010B)	Vanadium (SW-846 6010B)
Lead (SW-846 6020)	Zinc (SW-846 6010B)
Nitroaromatics/Nitramines (SW 846 Method 8321A)	
1,3-Dinitrobenzene	4-Amino-2,6-dinitrotoluene
1,3,5-Trinitrobenzene	4-Nitrotoluene
2-Amino-4,6-dinitrotoluene	HMX <sup>(1)</sup>
2-Nitrotoluene	Nitrobenzene
2,4-Dinitrotoluene	Nitroglycerin
2,4,6-Trinitrotoluene	PETN <sup>(2)</sup>
2,6-Dinitrotoluene	RDX <sup>(3)</sup>
3-Nitrotoluene	Tetryl (**
SVOCs (SW846 Method 8270C)	
1-Naphthylamine	4-Chlorophenyl phenyl ether
1,2-Dichlorobenzene	4-Dimethylaminoazobenzene
1,2,4-Trichlorobenzene	4-Nitroaniline
1,2,4,5-Tetrachlorobenzene	4-Nitrophenol
1,3-Dichlorobenzene	4-Phenylenediamine
1,3-Dinitrobenzene	4,6-Dinitro-2-methylphenol
1,3,5-Trinitrobenzene	5-Nitro-o-toluidine
1,4-Dichlorobenzene	7,12-Dimethylbenz(a)anthracene
1,4-Naphthoguinone	a,a-Dimethylphenethylamine
2-Acetylaminofluorene	Acenaphthene
2-Chloronaphthalene	Acenaphthylene
2-Chlorophenol	Acetophenone
2-Methylnaphthalene	Aniline
2-Methylphenol	Anthracene
2-Naphthylamine	Aramite
2-Nitroaniline	Benzo(a)anthracene
2-Nitrophenol	Benzo(b)fluoranthene
2-Picoline	Benzo(ghi)pervlene
2 3 4 6-Tetrachiorophenol	Benzo(k)fluoranthene
2.4-Dichlorophenol	Benzo(a)ovrene
2,4-Dimethylphenol	Benzyl alcohol
	bis(2-Chloroethoxy)methane
	his(2-Chloroethyl) ether
2,4-Dimuoloidene	his(2-Chloroisonronyl) ether
2,4,5-methorophenol	bis(2-Chlorosopropy) ether
2,4,6-Inchorophenol	Dis(2-Eurymexy) philialate
	Chlorobonziloto
2,6-Dinitrololuene	Chorobenzilate
3-ivieuryichoranthrene	
3-ivietnyiphenoi & 4-ivietnyiphenoi	Dianale Dihanz(a h)anthrasana
3,3Dicniorobenziaine	
	Diethylphthalate
4-Aminopipnenyi	
4-Bromophenyi phenyi ether	Dimetnyi phthalate
4-Chloro-3-methylphenol	Di-n-butyi phthalate
4-Chloroaniline	Di-n-octyl phthalate

Table 1 Analytical List

svoc	s (SW846 Method 8270C) Continued	
	Diphenylamine	N-Nitrosodiphenylamine
	Ethyl methanesulfonate	N-Nitrosomethylethylamine
	Fluoranthene	N-Nitrosomorpholine
	Fluorene	N-Nitrosopiperidine
	Hexachlorobenzene	N-Nitrosopyrrolidine
	Hexachlorobutadiene	O,O,O-Triethyl phosphorothioate
	Hexachlorocyclopentadiene	o-Toluidine
	Hexachloroethane	Parathion
	Hexachloropropene	Pentachlorobenzene
	Indeno(1,2,3-cd)pyrene	Pentachloroethane
	Isodrin	Pentachloronitrobenzene
	Isosafrole	Pentachlorophenol
	Isophorone	Phenacetin
	Methapyrilene	Phenanthrene
	Methyl methanesulfonate	Phenol
	Naphthalene	Phorate
	Nitrobenzene	Pronamide
	Nitroquinoline-I-oxide	Pvrene
	N-Nitrosodiethvlamine	Pvridine
	N-Nitrosodimethylamine	Safrole
	N-Nitrosodi-n-butvlamine	Sulfotepp
	N-Nitrosodi-n-propylamine	Thionazin
/OCs	(SW846 Method 8260B)	
	1.1-Dichloroethane	Chloroform
	1.1-Dichloroethene	Chloromethane
	1 1.1-Trichloroethane	Chloroprene
	1.1.1.2-Tetrachloroethane	Cis-1 2-Dichloroethene
	1.1.2-Trichloroethane	Cis-1.3-Dichloropropene
	1.1.2.2-Tetrachloroethane	Dibromochloromethane
	1.2-Dibromo-3-chloropropane (DBCP)	Dibromomethane
	1.2-Dibromoethane (EDB)	Dichlorodifluoromethane
	1 2-Dichloroethane	Ethyl methacrylate
	1 2-Dichloropropane	Ethylbenzene
	1.2.3-Trichloropropane	lodomethane
	1 4-Dioxane	Isobutyl alcohol
	2-Butanone (MEK)	Methachylopitrile
	2-Hexanone	Methyl methacrylate
	4-Methyl-2-pentanone	Methylene chloride
	Acetone	Propionitrile
	Acetonitrile	Styrene
	Acrolein	Tetrachloroethene
	Acrylonitrile	Toluene
	Allyl chloride	Trans-1 3-Dichloropropene
	Benzene	Trans-1,2-Dichloroethene
	Bromodichloromethane	Trans-1 4-Dichloro-2-butene
	Bromoform	Trichloroethene
	Bromomothane	Trichlorofluoromethane
	Carbon digulfido	Vinul apotato
	Carbon usunue	Vinyi alletate Vinyi alletate
	Chlorobonzono	Vilenes (total)
	Chloroothana	Ayiches (luid)
	Childhoethane	

### Table 1 Analytical List

(1) HMX= Octahydro-1,3,5,7-tetranitro-1,3,5,7-tetrazocine(2) PETN= Pentaerythritol Tetranitrate

(3) RDX= Hexahydro-1,3,5-trinitro-1,3,5-triazine(4) Tetryl= Methyl-2,4,6-trinitrophenylnitramine

Area	Matrix	Sample	Analyses			
		IDs	VOCs	SVOCs	Nitroaromatics	Metals
	Sediment	1SD1	X	X	X	X
	Surface Water	1SW1	X	x X	Х	X <sup>(1)</sup>
2	Sediment	2SD1	Х	X	X	X
	Surface Water	2SW1	X	X	X	X <sup>(1)</sup>
3	Sediment	3SD1	Х	X	Х	X
	Surface Water	3SW1	X	X	X	X <sup>(1)</sup>
4	Sediment	4SD1	X	Х	X	X
	Surface Water	4SW1	X	X	X	X <sup>(1)</sup>
5	Sediment	5SD1	X	X	X	X
	Surface Water	5SW1	Х	Х	Х	$\overline{X}^{(1)}$
6	Sediment	6SD1	Х	X	Х	X
	Surface Water	6SW1	Х	_X	X	X <sup>(1)</sup>
7	Sediment	7SD1	X	X	X	X
	Surface Water	7SW1	Х	X	Х	X <sup>(1)</sup>
8	Sediment	8SD1	Х	X	X	X
	Surface Water	8SW1	Х	X	X	X <sup>(1)</sup>

Table 2Sampling Locations

(1): Total and Dissolved

## FIGURES

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