# Appendix 13.1

Soil and Hydrogeological Investigation (K.T. Cullen & Co. Ltd) & Hydrogeological Assessment for Hammond Lane Metal Company

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# SOIL AND HYDROGEOLOGICAL INVESTIGATION

Greenfield Site, Ringaskiddy, Co. Cork,

FINAL REPORT Janaury 2001

Prepared for:

Project Management, Kilakee House, Belgard Square, Tallaght, Dublin 24.

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# TABLE OF CONTENTS

1	INTRODUCTION	1
2	SITE DESCRIPTION	1
2.1	Physical Features	1
2.2	Land Use	2
2.3	Hydrology	2
	2.3.1 Regional Drainage	2
	2.3.2 Local Drainage	2
2.4	General Geology and Hydrogeology	2
	2.4.1 Bedrock Geology	2
•.	2.4.2 Overburden Geology	3
	2.4.3 Hydrogeology	3
	2.4.4 Aquifer Vulnerability	3
3	FIELD ACTIVITIES	3
3.1	Soil Sampling	4
3.2	Monitoring Well Installation	4
3.3	Groundwater Sampling	5
4	ANALYTICAL RESULTS	6
4.1	Soil Analytical Results	6
	4.1.1 PROs, DROs, and Mineral Oils	6
	4.1.2 BTEX Compounds	6
	4.1.3 Volatile Organic Compounds	6
	4.1.4 Polynuclear Aromatic Hydrocarbons	6
	4.1.5 Metals and Total Phenols	7
	4.1.6 Pesticides	7
	4.1.7 Polychlorinated Biphenyls	7
4.2	Groundwater Analytical Results	8
••-	4.2.1 PROs, DROs, and Mineral Oils	8
	4.2.2 BTEX Compounds	8
	4.2.3 Volatile Organic Compounds	8
	4.2.4 Polynuclear Aromatic Hydrocarbons	8
	4.2.5 Toxic Metals	8
	4.2.6 Pesticides	8



	4.2.7	Polychlorinated Biphenyls			8
	4.2.8	Inorganics			9
5	SUM	MARY OF FINDINGS			9
_	5.1	Physical Observations			9
	5.2	Soil Quality Investigation			9
	5.3	Groundwater Quality Investigation			10
6	CONCLUSIONS				10
	6.1	Soil and Groundwater Quality			10
	6.2	Site Vulnerability			10
	6.4	Future Monitoring		2 - 19 2 - 1	10

### **TABLES**

## Soil Analytical Results

Table 1	DRO, PRO & Mineral Oils
Table 2	BTEX Compounds
Table 3 (a)&(b)	Volatile Organic Compounds
Table 4 (a)&(b)	PAHs - Original and Repeat Analysis
Table 5	Metals & Phenols
Table 6	Pesticides
Table 7 (a)&(b)	Polychlorinated Biphenyls

# Groundwater Analytical Results

Table 8	DRO, PRO & Mineral Oils
Table 9	BTEX Compounds
Table 10 (a)&(b)	Volatile Organic Compounds
Table 11	Polynuclear Aromatic Hydrocarbons
Table 12	Metals
Table 13	Pesticides
Table 14	Polychlorinated Biphenyls
Table 15	Inorganics



#### **FIGURES**

Figure 1 Regional Geology Map

Figure 2 Site Layout

### **APPENDICES**

Appendix A Trial Pit Logs
Appendix B Borehole Logs

Appendix C Letter from Alcontrol Geochem

Appendix D GSI Vulnerability Mapping Guidelines

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Soil and Hydrogeological Investigation

at

Greenfield Site, Ringaskiddy, Co. Cork

#### 1 INTRODUCTION

At the request of Project Management Ltd., K.T. Cullen & Co. Ltd were requested to undertake a full baseline hydrogeological investigation of a greenfield site at Ringaskiddy, Co. Cork.

This hydrogeological investigation involved the excavation of trial pits, installation of monitoring wells, and sampling/analyses of both soil and groundwater.

The investigation was carried out to establish baseline conditions of soil and groundwater beneath the ite, and to determine any going concerns regarding potential contamination in the subsurface. Field data was also collected on the hydrogeological conditions encountered on site.

#### 2 SITE DESCRIPTION

#### 2.1 Physical Features

The site is currently covered with grassland and appears to have been used for agricultural purposes in the past. The east of the site is covered with gorse scrub. It is understood that large amounts of soil was removed from the site in the past for reclamation purposes in the vicinity of the site, resulting in the steep embankment located towards the southern boundary of the site.

#### 2.2 Land Use

The surrounding land is predominantly agricultural but industrial sites are common in the Ringaskiddy area. Existing developments include the Hammond Lane Metal Company (HLM), located in the centre of the proposed site for development. Ispat Metal Processors are located to the north of the site, and Ringaskiddy Port to the northeast of the site.

### 2.3 Hydrology

#### 2.3.1 Regional Drainage

The proposed area for development lies within 50m of the West Channel into Cork Harbour.

#### 2.3.2 Local Drainage

Surface water within the site boundary appears to drain naturally through land drains along the field boundaries, following the natural topography of the landscape, generally towards the north of the site. Drainage is poor close to the road due to recent site activities, resulting in some flooding on site.

#### 2.4 General Geology and Hydrogeology

In considering the impact of the proposed development on the geology and groundwater quality, K.T. Cullen & Co. Ltd. have examined the following factors:

- Rock type and permeability
- Overburden type, thickness and, permeability
- Depth to water table
- Importance of groundwater as a resource
- Groundwater vulnerability

Data has been collated from previous investigations undertaken by this office in the Cork region, from the GSI database for County Cork, and on-site observations.

#### 2.4.1 Bedrock Geology

At this site the bedrock consists of pale green/grey mudstone, and is typical of the Lower Carboniferous Kinsale Formation (Cuskinny Member). It is thought to be between 235 metres and 243 metres thick. This member is typically made up of flaser-bedded sandstones and lenticular-bedded mudstones. It has been described as being composed of relatively thick sometimes conglomeratic sandstone units, alternating with thin sandstone laminated mudstones, massive claystones and heterolithic sediments (Geological Survey of Ireland - Geology of South Cork, 1994)

#### 2.4.2 Overburden Geology

The overburden geology consists of a shallow topsoil layer underlain by soft silty clays with some fine sands and gravels. Depth to bedrock varies across the site, from 1.0 metres below ground level (bgl) at BH-1, to greater than 9.0 metres bgl at BH-2. This thickness variation is a reflection of the undulating pre-glacial topography.

Sands were encountered in TP-1, TP-7, TP-16 and TP-17, and these areas are likely to allow water to be stored and to move through the subsurface. As some of the overburden is less than 1.0 metre in thickness (i.e. BH1), vertical migration of water directly into the bedrock aquifer is likely.

#### 2.4.3 Hydrogeology

The groundwater potential of Irish rocks is typically a function of fissure flow movement and storage, which is controlled by the intensity and development status of fissures, fractures and joints. The rocks are thought to be generally unproductive (i.e. individual well yields of less than 100 m<sup>3</sup>/day and often lower than 40 m<sup>3</sup>day - Geological Survey of Ireland "Geology of South Cork", 1995) although hydrogeological data is limited. This situation could be confirmed at the site by the drilling of a deep water well into the bedrock.

Water strikes in the bedrock were observed between 5 and 12 metres bgl in the overburden, typically occurring beneath the clays and immediately above the clean bedrock in the fractured/weathered zone.

#### 2.4.4 Aquifer Vulnerability

The GSI's Groundwater Protection Scheme Classification ranks the site as having extreme (E) vulnerability due to the limited overburden cover, which is less than 1.0 metres in thickness at some of soil and groundwater survey points. As the bedrock is considered to be a poor but locally productive aquifer (PI), the area can be assigned the rating PI/E under the GSI classification system.

#### 3 FIELD ACTIVITIES

Field activities for the purpose of this hydrogeological investigation were undertaken in November 2000 and consisted of the following stages:

- Desktop Review of Geology and Hydrogeology
- Soil Sampling
- Monitoring Well Installation



- Groundwater Sampling
- Elevation Survey (yet to be undertaken)

#### 3.1 Soil Sampling

A total of ten trial pits (TP-1 to TP-10) were initially excavated across the site in late November 2000. Additional sampling was undertaken in January 2001 (TP-11 to TP-17) and all sampling locations are shown in Figure 2 of this report. These excavations were undertaken to allow representative soil sample collection. Based on visual observations made on site, soil samples from varying layers were taken from each of the seventeen trial pit locations. Samples were sealed in a laboratory-supplied sample container and maintained at a temperature of <4°C in a mobile field laboratory.

The seventeen soil samples were submitted to Geochem Group Laboratories Ltd. and analysed for the following parameters:

- Petrol and Diesel Range Organics, Mineral Oils
- BTEX Compounds
- Volatile Organic Compounds (VOCs)
- Polycyclic Aromatic Hydrocarbons (PAHs)
- Metals and Total Phenols
- Pesticides (OPPs, OCPs, ONPs)
- Polychlorinated Biphenyls (PCBS)

Trial pit sampling logs are included in Appendix A.

#### 3.2 Monitoring Well Installation

Five permanent monitoring boreholes locations (BH-1, BH-2, BH-3, BH-4, and BH-5) were drilled under the continuous supervision of a K.T. Cullen & Co. Ltd. (KTC) Geologist. The well locations are shown in Figure 2 of this report. These locations were selected during the preliminary site walkover, and are based on the topography and geography of the site.

Items of concern noted during the site walkover include the Hammond Lane Metal Co. (HLM), which is located in the centre of the proposed site. Current activities at this site include the preparation of scrap metal, primarily from crushed cars, for reprocessing at the Ispat site located to the north of the property. The metal is crushed and sorted using magnetic techniques. Potential sources of contamination from this site would include hydrocarbon products remaining in the crushed cars.



Four of the monitoring wells are installed in bedrock. BH-2 was drilled to a depth of 8.5 metres bgl in the overburden, but did not encounter bedrock. Drilling and well construction logs are included in Appendix B of this report.

Narrow slotted screen was installed at all well borings locations, with an internal diameter of 0.05 metres. All screens were connected to the surface by PVC risers. A fine gravel pack was installed around each screen in order to filter water entering the well. Each pack was sealed above by a bentonite seal in order to prevent the vertical migration of fluids through the well annulus.

### 3.3 Groundwater Sampling

Following installation, each monitoring well was developed by the evacuation of more than three times the annular volume of the well. Well development grades the gravel pack into more complete contact with the aquifer and allows removal of suspended sediment which may remain following the drilling of the monitoring wells. More importantly, well development ensures that future sampling is representative of the quality of water in the surrounding aquifer.

All five monitoring wells were sampled on November 30<sup>th</sup> 2000, and these samples were subsequently forwarded to Alcontrol/ Geochem Group Laboratories in the U.K. for the following detailed analysis:

- Petrol and Diesel Range Organics, Mineral Oils
- BTEX Compounds
- Volatile Organic Compounds (VOCs)
- Polycyclic Aromatic Hydrocarbons (PAHs)
- Metals
- Pesticides (OPPs, OCPs, ONPs)
- Polychlorinated Biphenyls (PCBS)
- Inorganics

All samples were filled directly from a PVC bailer, preserved at <4°C and shipped to the laboratory in dedicated containers. The number of bottles, their codes and volumes were recorded on Monitoring Well Sampling Logs and on Chain of Custody forms.

#### 4. ANALYTICAL RESULTS

The analytical results for both soil and groundwater are presented in Tables 1 - 15 of this report,

Where relevant, the soil analytical results are discussed below with reference to the Dutch MAC (Maximum Admissible Concentration) thresholds, as standards for soil are not available in Ireland at present.

Groundwater analytical results are compared to the Irish Water Quality Standard for Drinking Water (S.I. No 81 of 1988) and the Dutch MAC Guidelines for groundwater as no other guidelines are currently available. Under the Dutch criteria for both soil and groundwater, the degree of contamination is assessed using the following guidelines:

S-Value

Reference for normal uncontaminated soil/groundwater

I-Value

Threshold for intervention

#### 4.1 Soil Analytical Results

The soil analytical results are presented in Tables 1-7 of this report.

#### 4.1.1 PROs, DROs and Mineral Oils

The analytical results for PROS, DROs and Mineral Oils are presented in Table 1. Detected concentrations for PROs, DROs and Mineral Oils reflect normal background concentrations for these parameters.

#### 4.1.2 BTEX Compounds

Results for the BTEX compounds are included in Table 2. Detected concentrations for these parameters were all below the laboratory detection limit of <0.01 mg/kg at all sampling locations.

#### 4.1.3 Volatile Organic Compounds

The analytical results for the VOCs are presented in Tables 3a and 3b and consist of 59 VOC parameters (EPA List). Detected concentrations for all VOCs were below the laboratory detection limit of 1 µg/kg.

#### 4.1.4 Polycyclic Aromatic Hydrocarbons (PAHs)

The analytical results for PAHs are presented in Tables 4a and 4b. The initial PAH results (sum of 10) included in Table 4a indicated concentrations for this parameter at all sampling locations, ranging between 1530  $\mu$ g/kg to 29282  $\mu$ g/kg across the site. These values exceed the Dutch MAC S-value of



1000 μg/kg for this parameter.

As the site is a greenfield site, and these values were not expected, a second series of trial pits were excavated in January 2001. TP-11 was placed immediately adjacent to TP-10 following a particularly elevated PAH concentration at this location. A further seven trial pits were excavated around the site. The results for this second sampling round are presented in Table 4b.

In the samples from the repeat trial pits, detected concentrations for the sum of 10 PAHs are considerably lower than in the original sampling round, ranging from  $6\mu g/kg$  in TP-11 to  $54\mu g/kg$  again in TP-11. None of the detected concentrations exceed the Dutch MAC S-value for the sum of 10 PAHs.

### 4.1.5 Metals and Total Phenols

The analytical results for Metals and Total Phenols are presented in Table 5 of this report. All metal parameters were detected below their respective Dutch MAC S-Values.

#### 4.1.6 Pesticides

The analytical results for Pesticides are presented in Table 6 of this report. The Geochem suite consists of three separate types of pesticides including Organochloride, Organonitrate and Organophosphate Pesticides, covering a wide range of these parameters. No pesticides were detected in any soil sample above the laboratory detection limit of  $1 \mu g/kg$  (laboratory detection limit).

#### 4.1.7 Polychlorinated Biphenyls (PCBs)

Analytical results for PCBs are included in Table 7a and Table 7b of this report. Detected concentrations for the 7 congeners (total) were elevated above the Dutch MAC S-Value for background conditions (20  $\mu$ g/kg) at TP-2 (0-5.5 metres), and TP-7 (0-2 metres), with levels of 643  $\mu$ g/kg and 98  $\mu$ g/kg respectively.

As the site is a greenfield location, KTC rescheduled this analysis to confirm the Alcontrol/ Geochem analytical results. These results are included in Table 7b. The repeated results give values of 13µgkg in TP-2 and 2µgkg in TP7, using a different extraction method, which is more applicable for greenfield sites. These values are less than the Dutch MAC S-Value.

To confirm the above finding, KTC resampled the site at 7 additional sampling locations (TP-11 to TP-17 inclusive). Results for this analysis are included in Table 7 (c) of this report. Detected concentrations for the repeat analysis were below the laboratory detection limit of  $1 \mu g/kg$ .

### 4.2 Groundwater Analytical Results

Groundwater analytical results for the five monitoring wells sampled, BH-1 to BH-5 inclusive, are presented in Tables 8 to 15 of this report.

#### 4.2.1 PROs. DROs, Mineral Oils

The analytical results for PROs, DROs and Mineral Oils are included in Table 8 of this report. Concentrations were all below the laboratory detection limit of 10µg/l.

#### 4.2.2 BTEX Compounds

Detected concentrations for the BTEX parameters (Table 9) were below the laboratory detection limit of 10 ug/l at all well sampling locations.

### 4.2.3 Volatile Organic Compounds

All VOC parameters analysed for in the five groundwater sampling locations were below the laboratory detection limit of 1  $\mu$ g/l (Table 10a and Table 10b)

### 4.2.4 Polycyclic Aromatic Hydrocarbons (PAHs)

The 16 priority PAH pollutants, for groundwater are presented in Table 11. Two of the PAHs, in particular Flouranthene and Phenanthrene, were slightly above their respective Dutch MAC S-values in BH-2, however these can be found naturally at such low concentrations.

#### 4.2.5 Toxic Metals

Details of the toxic metal results are included in Table 12 of this report. Arsenic was detected in BH-2 'ightly above the laboratory detection limit of 0.05, and appears to be an anomalous result. KTC has discussed this result with Alcontrol Geochem Ltd and has concluded that sample 'noise' may have affected the reading. Detected concentrations of the other eight metals analysed for are below their respective laboratory detection limits.

#### 4.2.6 Pesticides

The analytical results for Pesticides are presented in Table 13 of this report. Pesticide compounds were not detected in any samples above the laboratory detection limit of 1  $\mu$ g/l.

### 4.2.7 Polychlorinated Biphenyls (PCBs)

The analytical results for PCBs are presented in Table 14. PCBs were not detected in any samples above the laboratory detection limit of 1  $\mu$ g/l.

#### 4.2.8 Inorganics

Results for the inorganic parameters are included in Table 15 of this report. Detected concentrations for many of these parameters are indicative of this type of agricultural setting.

However, Ammonia and Nitrite values across the site appear slightly elevated above background concentrations at some of the sampling locations. Ammonia concentrations ranged between 1.0 mg/l and 1.9 mg/l, and Nitrite concentrations reached 0.1 mg/l in BH-3 and BH-4. Elevated concentrations for these parameters suggest slight organic contamination, and may be of an agricultural nature.

#### 5 SUMMARY OF FINDINGS

#### 5.1 Physical Observations

T' initial visual walkover survey showed no physical evidence of contamination across the c. 30 acre site.

The physical examination of the soil and groundwater samples carried out at the Greenfield Site, Ringaskiddy, Co. Cork revealed no physical evidence of contamination, such as chemical odours, iridescence, or other signs of contamination in any of the samples.

Much of the soil at the south and east end of the site has been removed for use in reclamation work in the vicinity of the site to the north. The west of the site has been used for agricultural purposes only, and excluding the Hammond Lane Metal Co. located in the centre of the site, there is no evidence of other developments at the site.

#### 5 Soil Quality Investigation

Initial soil samples taken from the site indicated concentrations for PAHs and PCBs above normal background levels. These results did not reflect on-site observations, and additional samples were taken to establish true conditions on site.

Results from samples taken during the repeat sampling round showed the soil to contain normal concentrations below background levels for these parameters. Following detailed queries regarding the PAH and PCB analysis, Alcontol Geochem has issued an explanation for the anomalous results (Appendix C). It appears that there was a problem with the solvent extraction process during the preparation of the soil samples.

Following detailed soil sampling and repeat analysis, the soil is free of industrial contaminants.



#### 5.3 Groundwater Quality Investigation

Slightly elevated Ammonia and Nitrite concentrations suggest potential organic contamination in the bedrock aquifer. Due to the shallow overburden cover and agricultural activity in this area, these values are not uncommon and are likely to be of agricultural origin.

All other groundwater results reflected normal background conditions for this type of environmental setting.

#### 6 CONCLUSIONS

#### 6.1 Soil and Groundwater Quality

The results of the soil and groundwater sampling suggest that there is no significant soil or groundwater contamination at the Ringaskiddy Greenfield site in Co. Cork.

Repeat sampling for PAHs and PCBs in the soils revealed that previous elevated results were erroneous.

High inorganics in the bedrock aquifer can most likely be attributed to agricultural activities on a site with very little or no overburden cover. It should also be noted that the levels of contamination in the groundwater are only slightly elevated above background.

#### 6.2 Site Vulnerability

Based on visual observations made on site during drilling and soil sampling, the overburden cover is very shallow, in some cases less than 1.0 metres in thickness in parts of the site.

Based on the thickness and type of overburden cover, the aquifer vulnerability for this site is considered extreme (GSI Guidelines for aquifer protection). As the bedrock is considered to be a poor but locally productive aquifer (PI), the area can be assigned the rating PI/E under the GSI classification system. (See Appendix D)

#### 6.3 Future Monitoring

To assess any variations in groundwater during the development of the Ringaskiddy Greenfield Site, monitoring of certain indicator parameters at all groundwater sampling locations is recommended.

Respectively submitted,

K. T. Cullen & Co. Ltd.

**CONOR WALL** 

Senior Environmental Scientist

1/2/01

DATE

MICHAEL CUNNINGHAM

**Industry Division Manager** 

1/2/201

# APPENDIX A

Project No.: 2626

Location: Ringaskiddy, Co. Cork

Date:

29/11/2000

Excavation Method: HyMac

Supervisor: Andrew Skelton

## TRIAL PIT NO.

TP1

eology:

Till

Depth (m): **Description:** 

0-0.1

TOPSOIL comprising medium brown soft damp silty clay with rootlets

0.1-0.4

Greyish brown loose dry gravelly silty CLAY Orange loose dry gravelly silty CLAY

0.4-0.9 0.9 - 4

Greyish brown loose gravelly silty CLAY with some greenish horizons with sands becoming

frequent

4-4.5

Pale green broken MUDSTONE

ominant Matrix: ominant Clasts:

pth to Rock: 4m

Rock Type: Pale green mudstone

atic Water Level: 1m

Water Entry: 1.8m

Total Depth: 4.5m

Comments: Pit collapsing from 2.5m

Sampled at 1.8m



ullen & Co. Ltd.

Project No.: 2626

Location: Ringaskiddy, Co. Cork

Date:

29/11/2000

Excavation Method: HyMac

Supervisor: Andrew Skelton

## TRIAL PIT NO.

TP2

eology: Depth (m):

**Description:** 

0-0.2

TOPSOIL comprising medium brown moist sandy silt with rootlets

0.2-0.5

Orange slightly loose dry sandy clayey SILT

0.5-5.5

Medium brown slightly loose dry gravelly sandy SILT with occasional boulders

edrock 5.5-5.6 Pale green broken MUDSTONE

minant Matrix: minant Clasts:

Depth to Rock: 5.5m

Rock Type: Pale green mudstone

Water Level: 4.5m

Water Entry: 5m

Total Depth: 5.6m

Comments: Sampled from 0-5.5m



en & Co. Ltd.

Project No.: 2626

Location: Ringaskiddy, Co. Cork

Date:

30/11/2000

Excavation Method: HyMac

Supervisor:

**Andrew Skelton** 

## TRIAL PIT NO.

TP3

leology: Depth (m):

**Description:** 

0-0.1

TOPSOIL comprising medium brown soft silty clay with rootlets

0.1-0.6

Orange brown silty CLAY

0.6-1.9 edrock

Weathered fractured green MUDSTONE

ominant Matrix:

ominant Clasts:

7th to Rock: 0.6m

Rock Type: Green mudstone

itic Water Level: -

Water Entry: -

Total Depth: 1.9m

Comments: Sampled 0-1.9m



len & Co. Ltd.

Project No.: 2626

Location: Ringaskiddy, Co. Cork

Date:

30/11/2000

Excavation Method: HyMac

Supervisor: Conor Wall

### TRIAL PIT NO.

TP4

ology: Depth (m): Description:

0-0.1

**TOPSOIL** 

0.1-2.4

Medium brown silty gravelly CLAY

2.4-4

Medium brown clayey SILT with fine sands

minant Matrix:

ninant Clasts:

Depth to Rock: -

.ock Type: -

Water Level: -

Water Entry: Slight entry at 3.4m

Total Depth: 4m

Comments: Sampled 0-3.5m



n & Co. Ltd.

No.: 2626

Location: Ringaskiddy, Co. Cork

Date: 30/11/2000

tion Method: HyMac

Supervisor: Andrew Skelton

# TRIAL PIT NO.

TP5

Depth (m): Description:

0-0.2

**TOPSOIL** 

0.2-1

Medium brown soft clayey SILT with occasional gravels

1-2.8

Medium brown soft clayey SILT with sands and gravels

/latrix:

>lasts:

o Rock: -

k Type: -

r Level: 2m

r Entry: 1m, 2.5m

Depth: 2.8m

ıments: Pit collapsing

Sample 0 - 2.8m

Project No.: 2626

Location: Ringaskiddy, Co. Cork

Date: 30/11/2000

Excavation Method: HyMac

Supervisor: **Andrew Skelton** 

### TRIAL PIT NO.

TP7

eology: Depth (m):

Description:

Medium brown soft clayey SILT with subrounded gravels and frequent subrounded

cobbles

2-3

Medium brown soft clayey fine SAND with subrounded cobbles

edrock 3-4 Broken green mudstone BEDROCK

minant Matrix:

minant Clasts:

Or 'h to Rock: 3m

Rock Type: Green mudstone

c Water Level: 3m

Water Entry: 3m

Total Depth: 4m

Comments: Sampled 0-2m and 3-4m



1 & Co. Ltd.

Project No.: 2626

Location: Ringaskiddy, Co. Cork

Date: 30/11/2000

Excavation Method: HyMac

Supervisor: Andrew Skelton

## TRIAL PIT NO.

TP9

eology: Depth (m):

**Description:** 

0-0.3

Medium brown soft clayey SILT with subrounded gravels and frequent subrounded

cobbles

edrock 0.3 - 1 Pale green broken mudstone BEDROCK

minant Matrix:

minant Clasts:

Depth to Rock: 0.3m

ock Type: Green mudstone

lic Water Level: -

Water Entry: -

Total Depth: 1m

Comments: Sampled 0-1m



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Project No.: 2626

Location: Ringaskiddy, Co. Cork

Date: 30/11/2000

Excavation Method: HyMac

Supervisor: Andrew Skelton

## TRIAL PIT NO.

**TP10** 

eology: Depth (m):

Description:

0-1

edrock 1-1.2

Medium brown gravelly SILT withj frequent subrounded cobbles

Pale green broken mudstone BEDROCK

ominant Matrix:

minant Clasts:

Depth to Rock: 1m

Rock Type: Green mudstone

ic Water Level: -

Water Entry: -

Total Depth: 1.2m

Comments: Sampled 0-1m



en & Co. Ltd.

Project No.: 2626

Location: Ringaskiddy, Co. Cork

Date:

12/1/2001

Excavation Method: HyMac

Supervisor:

**Andrew Skelton** 

## TRIAL PIT NO.

**TP11** 

eology: Depth (m):

**Description:** 

0-3

Medium brown firm dry clayey SILT with frequent angular gravels and frequent angular

cobbles and boulders

ominant Matrix:

minant Clasts:

Depth to Rock: -

Rock Type: -

c Water Level: -

Water Entry: 2.1m

Total Depth: 3m

Comments: Sampled 0-1m

Sampled 1-3m

No odour



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Project No.: 2626

Location: Ringaskiddy, Co. Cork

Date:

12/1/2001

Excavation Method: HyMac

Supervisor: Andrew Skelton

### TRIAL PIT NO.

**TP12** 

eology: Depth (m):

**Description:** 

0-2.6

Medium brown firm dry gravelly clayey SILT with frequent angular cobbles Pale green fissile mudstone BEDROCK

ock 2.6-

pminant Matrix:

minant Clasts:

Depth to Rock: 2.6m

Rock Type: Pale green mudstone bedrock

itic Water Level: -

Water Entry: -

Total Depth: 2.6m

Comments: Sampled 0-1m

Sampled 1-2.6m

No odour



en & Co. Ltd.

Project No.: 2626

Location: Ringaskiddy, Co. Cork

Date:

12/1/2001

Excavation Method: HyMac

**Andrew Skelton** Supervisor:

# TRIAL PIT NO.

**TP13** 

Depth (m): **Description:** eology:

0-0.2

Grey brown silty GRAVEL

0.2-2

Medium brown firm dry sandy clayey SILT with frewuent subrounded cobbles and gravels

2-3.6

Medium brown soft damp silty SAND

ominant Matrix:

minant Clasts:

Depth to Rock: -

Rock Type: -

lic Water Level: -

Water Entry: 3.4m

Total Depth: 3.6m

Comments: Sampled 0-1m

Sampled 1-3.4m

No odour



llen & Co. Ltd.

roject No.: 2626 Location: Ringaskiddy, Co. Cork

Date: 12/1/2001

Excavation Method: HyMac

Supervisor: Andrew Skelton

# TRIAL PIT NO.

**TP14** 

plogy: Depth (m): Description:

0-2.6

Medium brown firm to soft dry clayey gravelly SILT with subangular cobbles

2.6-3.8 N

Medium brown soft damp fine sandy CLAY

minant Matrix:

minant Clasts:

Depth to Rock: -

Rock Type: -

lic Water Level: -

Water Entry: 3m

Total Depth: 3.8m

Comments: Sampled 0-3.8m

No odour



Cullen & Co. Ltd.

Project No.: 2626

Location: Ringaskiddy, Co. Cork

Date:

12/1/2001

Excavation Method: HyMac

Supervisor:

Andrew Skelton

### TRIAL PIT NO.

**TP15** 

Geology: Depth (m): Description:

0-0.3

Medium brown loose dry silty TOPSOIL

0.3-0.5

Orange clayey gravelly SILT

0.5-2

Medium brown loose gravelly clayey SILT

Dominant Matrix:

Dominant Clasts:

Depth to Rock: -

Rock Type: -

latic Water Level: 1m

Water Entry: 1m (field drain)

Total Depth: 2m

Comments: Sampled 0-2m

No odour



illen & Co. Ltd.

Project No.: 2626

Location: Ringaskiddy, Co. Cork

Date:

12/1/2001

Excavation Method: HyMac

Supervisor: Andrew Skelton

# TRIAL PIT NO.

### **TP16**

Geology: Depth (m): Description:

Till 0-0.3 0.3-0.5 Medium brown loose dry clayey SILT Orangey brown loose dry clayey SILT

0.5-1.5

Medium brown soft silty fine SAND

1.5-2.5

Medium brown soft wet fine SAND with gravels

Dominant Matrix:

Dominant Clasts:

Depth to Rock: -

Rock Type: -

tatic Water Level: 1.5m

Water Entry: 2.5m

Total Depth: 2.5m

Comments: Sampled 0-2.5m

No odour



Cullen & Co. Ltd.

ct No.: 2626 Loca

Location: Ringaskiddy, Co. Cork

Date: 12/1/2001

avation Method: HyMac

Supervisor: Andrew Skelton

### TRIAL PIT NO.

**TP17** 

: Depth (m): Description:

0-0.4 Greyish brown soft dry clayey SILT

0.4-1.4 Medium brown to pale brown soft very fine sandy SILT

1.4-2 Medium brown wet silty fine SAND

Matrix:

Clasts:

to Rock: -

or' Type: -

ier Level: 1m

er Entry: 1.5m

al Depth: 2m

o. Ltd.

mments: Sampled 0-2m

No odour

# APPENDIX B

A control of the second of the

### **WELL LOG**

Well Ident

2626/BH1

Description
Monitoring Well

Location

Ringaskiddy, Co. Cork

**Drilling Date** 

All diameters in mm All depths in metres Scale

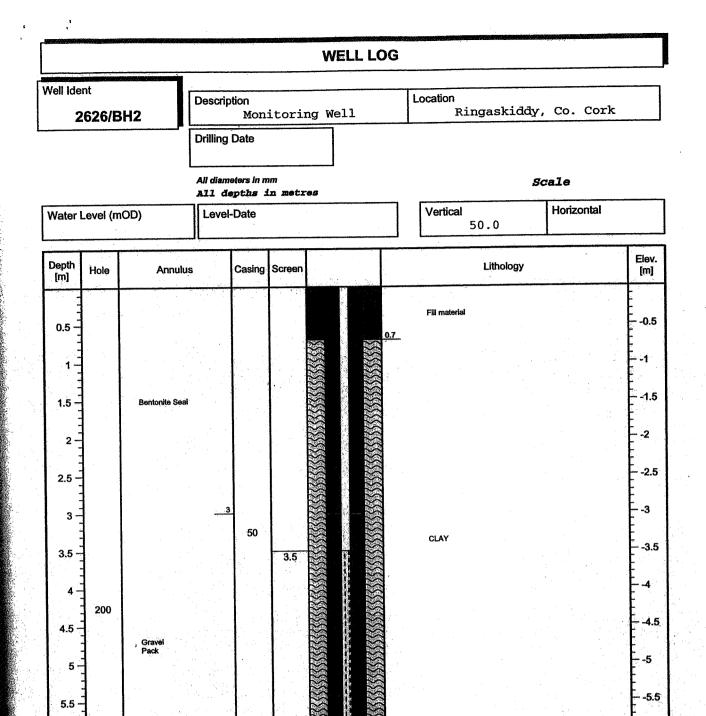
Water Level (mOD)

Level-Date

Vertical 50.0

Horizontal

Depth [m]	Hole	Annulus	Casing	Screen		Lithology	Elev. [m]
0.5		Backfill	l			CLAY	ò.5
1-			-1			늘 이 그는 사람 사람들이 그 사람	1
							1.5
1.5 -							- 1.8
2	8 L	Bentonite Seal					2
			.				Ę
2.5							2.5 -
-			3			Pale grey/green mudstone	E
3-							3 -
3.5 -							3.5
	200		50				Ē
4-	200						-4
					4.		
4.5 −				4.5	3		4.5 -
5-					5	Water Entry	- 5
		Gravel					Ę
5.5 -	1	Pack					-5.5
							Ē,
6 -							6 -
6.5 -	}					Pale grey/green mudstone	- 6.5
0.5							Ē
7-	]						-7
	]						F
7.5 -	7.6		7.6 7.6	7.6	<b>138 138</b> 7	7.6	7.5 -
8 -	]						E _8
8-	}						Ė.
8.5 -	1				1		-8.5
	1						E
9 -	1						9



6.5

8.5

Side wall collapse

6.5

Water Entry

CLAY

8.5

6

6.5

7

7.5

8

8.5

9

8.5

-6

- -6.5

-7.5

-8

-8.5

# **WELL LOG**

Well Ident

2626/BH3

Description
Monitoring Well

Location

Ringaskiddy, Co. Cork

Drilling Date

All depths in metres

Scale

Water Level (mOD)

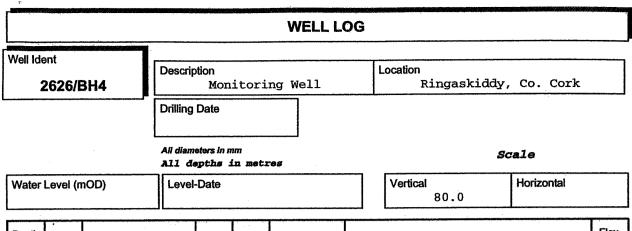
Level-Date

Vertical Horizontal 80.0

Depth Hole	Annulus	Casing Screen		Lithology	Elev. [m]
1 2 3 4 5 6 7 7 10 10 11 11 12 13 14 15 14 11 12 13 14 14 15 14 14 15 15 16 17 17 17 17 17 17 17 17 17 17 17 17 17	Bentonite Seal	<u>5</u> 50	3		1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 14 14

	ļ					WELL LO	OG			
Well Ide	ent 2626/1	3H3	Descri		oring	Well		Location Ringaskid	ldy, Co. Cork	
				Date  eters in mm  epths in					Scale	
Water	Level (n	nOD)	Level	-Date				Vertical 80.0	Horizontal	
Depth [m]	Hole	Annulu	s	Casing S	creen			Lithology		Elev. [m]

Depth [m]	Hole	Annulus	Casing	Screen		Lithology	Elev. [m]
	2030	Gravel Pack <u>15</u>	<del>5</del> 8	15		15_ Pale grey mudstone	
15	•						-15
=							F
16							-16
				*			
17							-17
1							F I
18 -							18
							Ė I
							E I
19					·		-19
							E I
20 -							-20
		4					
			ŀ				E
21 -							-21
		ж • 1					<b>E</b> 1
22 -							-22
22 -							E - 1
-							Ė I
23 -							-23
							E I
			-				-24
24 –							24
							F I
25							-25
							Εl
							E
26 -					4	·	26
							F
27 -							-27
	1						E
=						•	E
28 -							-28
]							E
	1						E I
I ~	1	l	1	L	L		



Depth [m]	Hole	Annulus	Casing	Screen	Lithology	Elev. [m]
1	200	Bentonite Seal  Bentonite Seal  Gravel Pack	50	9	Boulder clay  5  Pale grey mudstone  Water Entry  Pale grey mudstone	-1 -2 -3 -4 -5 -6 -7 -8 -9 -10 -11 -12 -13 -14 -14 -15 -14 -14 -15 -14 -15 -16 -17 -18 -19 -10 -11 -12 -13 -14 -15 -16 -17 -18 -19 -10 -11 -12 -13 -14 -15 -16 -17 -18 -19 -10 -11 -12 -13 -14 -15 -16 -17 -18 -19 -10 -11 -12 -13 -14 -15 -16 -17 -18 -19 -10 -10 -11 -12 -13 -14 -15 -16 -17 -18 -19 -10 -10 -10 -10 -10 -10 -10 -10 -10 -10

**WELL LOG** Well Ident Description Location 2626/BH5 Monitoring Well Ringaskiddy, Co. Cork **Drilling Date** 29.11.2000 All diameters in mm Scale All depths in metres Water Level (mOD) Level-Date Vertical Horizontal 80.0

<u></u>						80.0	
Depth [m]	Hole	Annulus	Casing	Screen		Lithology	Elev. [m]
1 2 3		Backvill	.5		\$3,535,535,535,535,535,535,535,535,535,5	CLAY	7 - 1 - 2 - 3 - 3
4 55		Bentonite Seal	4				-4 -1
5 6 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	200		50			Pale grey mudstone	-5 -6
8		Gravel Pack		7	8.5		-7 8
9 10 10	10		10 10	10	10	Water Entry Pale grey mudstone	10
11 11 11 11 11 11 11 11 11 11 11 11 11							-11
13					-		-13
14							-14 14

# APPENDIX C

Wed, Jan 31, 2001 14:28

#### repeat data

From: Hazel Davidson <a href="https://www.hazel.davidson@geochem.com">https://www.hazel.davidson@geochem.com</a>

To: "'cwall@ktcullen.ie'" <cwall@ktcullen.ie>

Date: Tue, Dec 19, 2000, 18:27

**Subject:** repeat data

#### Dear Conor

With reference to the repeated PAH data, the tests confirm the presence of trace amounts fof PAHs, but the naphthalene levels are significantly reduced. Upon investigation, this was found to be due to an artefact introduced during the solvent extraction process, caused by a particular batch of solvent. This has now been rectified.

The soil samples do appear to be contaminated with PCBs, but the inconsistency in the repeat analyses is probably due to a lack of homogeneity in the wet soil samples, as small inclusions of contaminated material may cause 'hot spots'. We would recommend further analysis of these samples to provide a better overview of the site.

For our own benefit, we are running two of the samples using a semi-volatile full scan, which will enable us to carry out a full library search.

We will endeavour to provide this additional data as soon as possible, and please do not hesitate to contact me if you wish to discuss this further.

Regards

Hazel

For and on behalf of ALcontrol Geochem Chester Street Chester, CH4 8RD United Kingdom

Phone: +44 (0)1244 671121

Fax: +44 (0)1244 683306

website: www.alcontrol.com

Parth Sciences & Environmental Laboratory Services

Marketing Info : mkt@geochem.com

The information in this e-mail is confidential and may also be legally

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Registered Office: Templeborough House, Mill Close, Rotherham S60 1BZ
Registered in England & Wales No. 4057291

# APPENDIX D

# 3. Land Surface Zoning for Groundwater Protection

# 3.1 Information and Mapping Requirements for Land Surface Zoning

The groundwater resources protection zone map is a land-use planning map, and therefore is the most useful map for the decision-making process. It is the ultimate or final map as it is obtained by combining the aquifer and vulnerability maps. The aquifer map boundaries, in turn, are based on the bedrock map boundaries and the aquifer categories are obtained from an assessment of the available hydrogeological data. The vulnerability map is based on the subsoils map, together with an assessment of relevant hydrogeological data, in particular indications of permeability and karstification. This is illustrated in Figure 3.

Similarly, the source protection zone maps result from combining vulnerability and source protection area maps. The source protection areas are based largely on assessments of hyr geological data. This is illustrated in Figure 4.

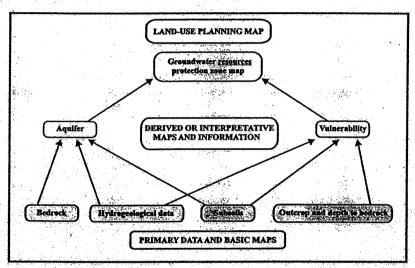


Figure 3. Conceptual framework for production of groundwater resource protection zones, indicating information needs and links

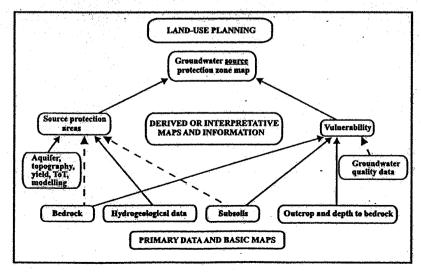


Figure 4. Conceptual framework for production of groundwater source protection zones, indicating information needs and links

# 3.2 Vulnerability Categories

Vulnerability is a term used to represent the intrinsic geological and hydrogeological characteristics that determine the ease with which groundwater may be contaminated by human activities.

The vulnerability of groundwater depends on: (i) the time of travel of infiltrating water (and contaminants); (ii) the relative quantity of contaminants that can reach the groundwater; and (iii) the contaminant attenuation capacity of the geological materials through which the water and contaminants infiltrate. As all groundwater is hydrologically connected to the land surface, it is the effectiveness of this connection that determines the relative vulnerability to contamination. Groundwater that readily and quickly receives water (and contaminants) from the land surface is considered to be more vulnerable than groundwater that receives water (and contaminants) more slowly and in lower quantities. The travel time, attenuation capacity and quantity of contaminants are a function of the following natural geological and hydrogeological attributes of any area:

- (i) the subsoils that overlie the groundwater;
- (ii) the type of recharge whether point or diffuse; and
  - ) the thickness of the unsaturated zone through which the contaminant moves.

In general, little attenuation of contaminants occurs in the bedrock in Ireland because flow is almost wholly via fissures. Consequently, the subsoils (sands, gravels, glacial tills (or boulder clays), peat, lake and alluvial silts and clays), are the single most important natural feature influencing groundwater vulnerability and groundwater contamination prevention. Groundwater is most at risk where the subsoils are absent or thin and, in areas of karstic limestone, where surface streams sink underground at swallow holes.

The geological and hydrogeological characteristics can be examined and mapped, thereby providing a groundwater vulnerability assessment for any area or site. Four groundwater vulnerability categories are used in the scheme - extreme (E), high (H), moderate (M) and low (L). The hydrogeological basis for these categories is summarised in Table 1 and further details can be obtained from the GSI. The ratings are based on pragmatic judgements, experience and available technical and scientific information. However, provided the limitations are appreciated, vulnerability assessments are essential when considering the location of potentially polluting activities. As groundwater is considered to be present everywhere in Ireland, the vulnerability concept is applied to the entire land surface. The ranking of vulnerability does not take into consideration the biologically-active soil zone, as contaminants from point sources are usually discharged below this zone, often at depths of at least 1m. However, the groundwater protection responses take account of the point of discharge for each activity.

Vulnerability maps are an important part of groundwater protection schemes and are an essential element in the decision-making on the location of potentially polluting activities. Firstly, the vulnerability rating for an area indicates, and is a measure of, the likelihood of contamination. Secondly, the vulnerability map helps to ensure that a groundwater protection scheme is not unnecessarily restrictive on human economic activity. Thirdly, the vulnerability map helps in the choice of preventative measures and enables developments, which have a significant potential to contaminate, to be located in areas of lower vulnerability.

In summary, the entire land surface is divided into four vulnerability categories - extreme (E), high (H), moderate (M) and low (L) - based on the geological and hydrogeological factors described above. This subdivision is shown on a groundwater vulnerability map. The map shows the vulnerability of the first groundwater encountered (in either sand/gravel aquifers or in bedrock) to contaminants released at depths of 1-2 m below the ground surface. Where contaminants are released at significantly different depths, there will be a need to determine groundwater vulnerability using site-specific data. The characteristics of individual contaminants are not taken into account.

		Hydrog	geological Condition	15	
Vulnerability Rating	Subsoil Pe	ermeability (Type)	) and Thickness	Unsaturated Zone	Karst Features
	High permeability (sand/gravel)	Moderate permeability (e.g. Sandy subsoil)	Low permeability (e.g. Clayey subsoil, clay, peat)	(Sand/gravel aquifers only)	(<30 m radius)
Extreme (E)	0 - 3.0m	0 - 3.0m	0 - 3.0m	0 - 3.0m	
High (H)	>3.0m	3.0 - 10.0m	3.0 - 5.0m	>3.0m	N/A
Moderate (M)	N/A	>10.0m	5.0 - 10.0m	N/A	N/A
Low (L)	· N/A	N/A	> 10.0m	N/A	N/A

Notes: (1) N/A = not applicable.

(2) Precise permeability values cannot be given at present.

(3) Release point of contaminants is assumed to be 1-2 m below ground surface.

Table 1. Vulnerability Mapping Guidelines

# 1.3 Source Protection Zones

iroundwater sources, particularly public, group scheme and industrial supplies, are of critical nportance in many regions. Consequently, the objective of source protection zones is to rovide protection by placing tighter controls on activities within all or part of the zone of ontribution (ZOC) of the source.

here are two main elements to source protection land surface zoning:

Areas surrounding individual groundwater sources; these are termed source protection areas (SPAs)

Division of the SPAs on the basis of the vulnerability of the underlying groundwater to contamination.

nese elements are integrated to give the source protection zones.

### 3.1 Delineation of Source Protection Areas

vo source protection areas are recommended for delineation:

Inner Protection Area (SI):

Outer Protection Area (SO), encompassing the remainder of the source catchment area or ZOC.

delineating the inner (SI) and outer (SO) protection areas, there are two broad approaches: st, using arbitrary fixed radii, which do not incorporate hydrogeological considerations; and condly, a scientific approach using hydrogeological information and analysis, in particular hydrogeological characteristics of the aquifer, the direction of groundwater flow, the pumping e and the recharge.

nere the hydrogeological information is poor and/or where time and resources are limited, simple zonation approach using the arbitrary fixed radius method is a good first step that tuires little technical expertise. However, it can both over- and under-protect. It usually er-protects on the downgradient side of the source and may under-protect on the upgradient e, particularly in karst areas. It is particularly inappropriate in the case of springs where are is no part of the downgradient side in the ZOC. Also, the lack of a scientific basis reduces defensibility as a method.

There are several hydrogeological methods for delineating SPAs. They vary in complexity, cost and the level of data and hydrogeological analysis required. Four methods, in order of increasing technical sophistication, are used by the GSI:

- (i) calculated fixed radius;
- (ii) analytical methods;
- (iii) hydrogeological mapping; and
- (iv) numerical modelling.

Each method has limitations. Even with relatively good hydrogeological data, the heterogeneity of Irish aquifers will generally prevent the delineation of definitive SPA boundaries. Consequently, the boundaries must be seen as a <u>guide</u> for decision-making, which can be reappraised in the light of new knowledge or changed circumstances.

# 3.3.1.1 Inner Protection Area (SI)

This area is designed to protect against the effects of human activities that might have an immediate effect on the source and, in particular, against microbial pollution. The area is defined by a 100-day time of travel (TOT) from any point below the water table to the source. (The TOT varies significantly between regulatory agencies in different countries. The 100-day

it is chosen for Ireland as a relatively conservative limit to allow for the heterogeneous nature of Irish aquifers and to reduce the risk of pollution from bacteria and viruses, which in some circumstances can live longer than 50 days in groundwater.) In karst areas, it will not usually be feasible to delineate 100-day TOT boundaries, as there are large variations in permeability, high flow velocities and a low level of predictability. In these areas, the total catchment area of the source will frequently be classed as SI.

If it is necessary to use the arbitrary fixed radius method, a distance of 300m is normally used. A semi-circular area is used for springs. The distance may be increased for sources in karst aquifers and reduced in granular aquifers and around low yielding sources.

# 3.3.1.2 Outer Protection Area (SO)

This area covers the remainder of the ZOC (or complete catchment area) of the groundwater source. It is defined as the area needed to support an abstraction from long-term groundwater recharge i.e. the proportion of effective rainfall that infiltrates to the water table. The abstraction rate used in delineating the zone will depend on the views and recommendations of the source owner. A factor of safety can be taken into account whereby the maximum daily abstraction rate is increased (typically by 50%) to allow for possible future increases in abstraction and 1 expansion of the ZOC in dry periods. In order to take account of the heterogeneity of many lrish aquifers and possible errors in estimating the groundwater flow direction, a variation in the flow direction (typically ±10-20°) is frequently included as a safety margin in delineating the ZOC.

A conceptual model of the ZOC and the 100-day TOT boundary is given in Figure 5.

If the arbitrary fixed radius method is used, a distance of 1000m is recommended with, in some instances, variations in karst aquifers and around springs and low-yielding wells.

The boundaries of the SPAs are based on the horizontal flow of water to the source and, in the case particularly of the Inner Protection Area, on the time of travel in the aquifer. Consequently, the vertical movement of a water particle or contaminant from the land surface to the water table is not taken into account. This vertical movement is a critical factor in contaminant attenuation, contaminant flow velocities and in dictating the likelihood of contamination. It can be taken into account by mapping the groundwater vulnerability to contamination.

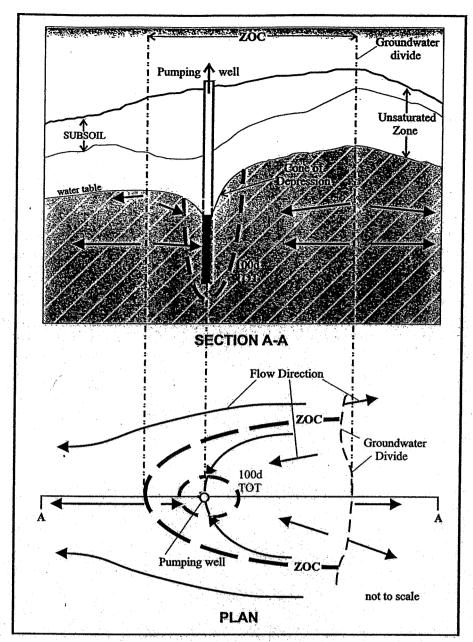


Figure 5. Conceptual Model of the Zone of Contribution (ZOC) at a Pumping Well (adapted from US EPA, 1987)

# 3.3.2 Delineation of Source Protection Zones

The matrix in Table 2 below gives the result of integrating the two elements of land surface zoning (SPAs and vulnerability categories) – a possible total of eight source protection zones. In practice, the source protection zones are obtained by superimposing the vulnerability map on the source protection area map. Each zone is represented by a code e.g. SO/H, which represents an Outer Source Protection area where the groundwater is highly vulnerable to contamination. The recommended map scale is 1:10,560 (or 1:10,000 if available), though a smaller scale may be appropriate for large springs.

VULNERABILITY	SOURCE PROT	<b>TECTION ZONE</b>
RATING	Inner (SI)	Outer (SO)
Extreme (E)	SI/E	SO/E
High (H)	SI/H	SO/H
Moderate (M)	SI/M	SO/M
Low (L)	SI/L	SO/L

Table 2. Matrix of Source Protection Zones

All of the hydrogeological settings represented by the zones may not be present around each groundwater source. The integration of the SPAs and the vulnerability ratings is illustrated in Figure 6.

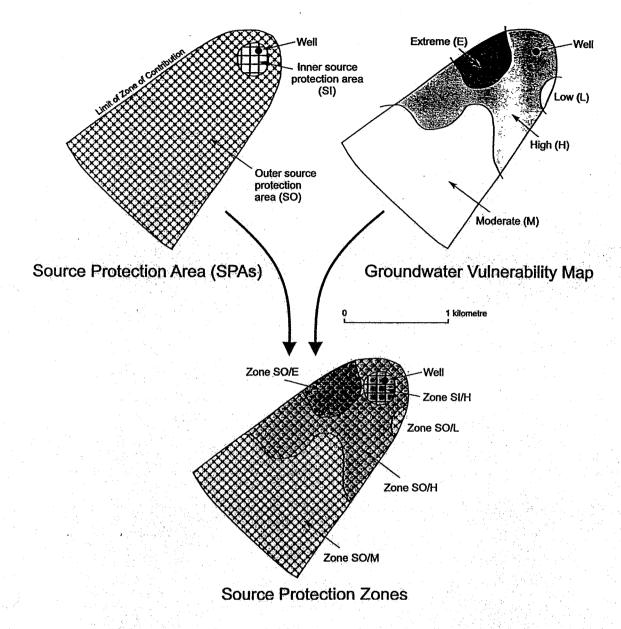


Figure 6. Delineation of source protection zones around a public supply well from the integration of the source protection area map and the vulnerability map.

# 3.4 Resource Protection Zones

For any region, the area outside the SPAs can be subdivided, based on the value of the resource and the hydrogeological characteristics, into eight aquifer categories:

## Regionally Important (R) Aquifers

- (i) Karstified aquifers (Rk)
- (ii) Fissured bedrock aquifers (Rf)
- (iii) Extensive sand/gravel aquifers (Rg)

# Locally Important (L) Aquifers

- (i) Sand/gravel (Lg)
- (ii) Bedrock which is Generally Moderately Productive (Lm)
- (iii) Bedrock which is Moderately Productive only in Local Zones (LI)

# Poor (P) Aquifers

- (i) Bedrock which is Generally Unproductive except for Local Zones (PI)
- (ii) Bedrock which is Generally Unproductive (Pu)

These aquifer categories are shown on an aquifer map, which can be used not only as an element of a groundwater protection scheme but also for groundwater development purposes.

The matrix in Table 3 below gives the result of integrating the two regional elements of land surface zoning (vulnerability categories and resource protection areas) — a possible total of 24 resource protection zones. In practice this is achieved by superimposing the vulnerability map on the aquifer map. Each zone is represented by a code e.g. Rf/M, which represents areas of regionally important fissured aquifers where the groundwater is moderately vulnerable to contamination. In land surface zoning for groundwater protection purposes, regionally important sand/gravel (Rg) and fissured aquifers (Rf) are zoned together, as are locally important sand/gravel (Lg) and bedrock which is moderately productive (Lm). All of the hydrogeological settings represented by the zones may not be present in each local authority area.

		RESOU	RCE PRO	TECTION	ZONES	
VULNERABILITY, RATING		Important ers (R)		mportant		quifers P)
	Rk	Rf/Rg	Lm/Lg	L1	P1	Pu
Extreme (E)	Rk/E	Rf/E	Lm/E	L1/E	P1/E	
High (H)	Rk/H	Rf/H	Lm/H	L1/H	P1/H	Pu/E
Moderate (M)	Rk/M	Rf/M				Pu/H
Low (L)			Lm/M	L1/M	P1/M	Pu/M
Lion (Li)	Rk/L	Rf/L	Lm/L	L1/L	P1/L	Pu/L

Table 3. Matrix of Resource Protection Zones

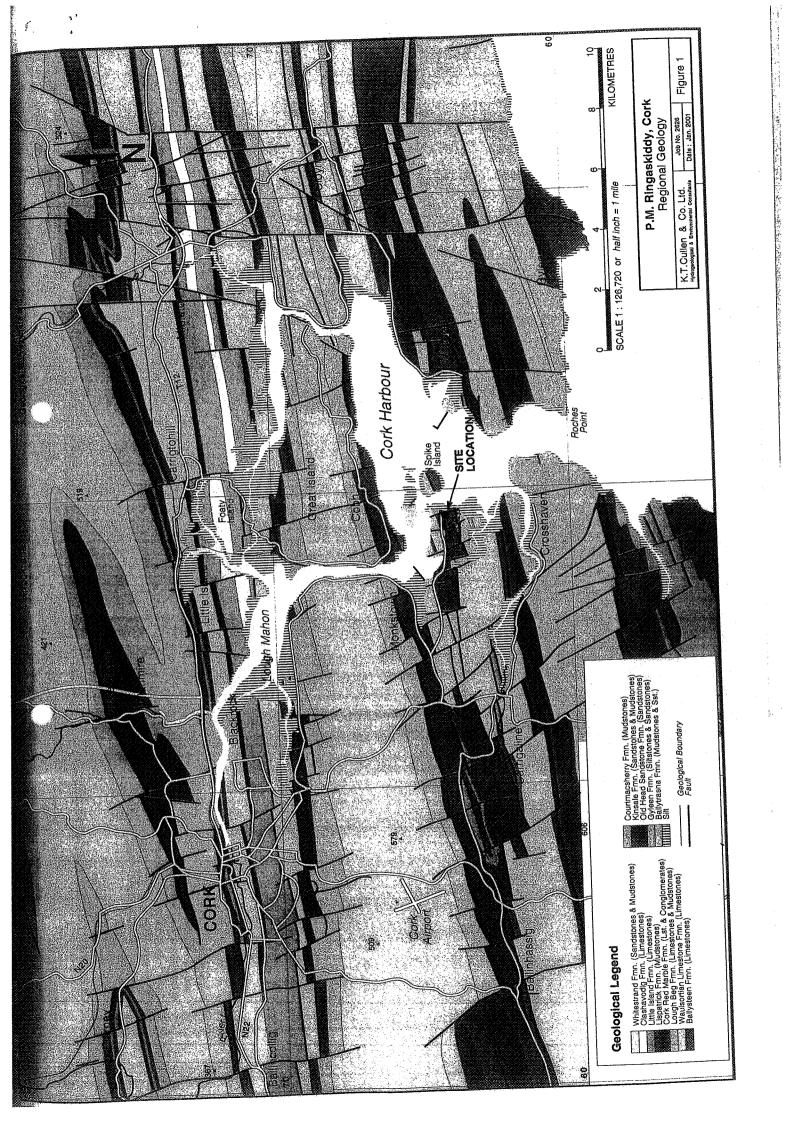
# 3.5 Flexibility, Limitations and Uncertainty

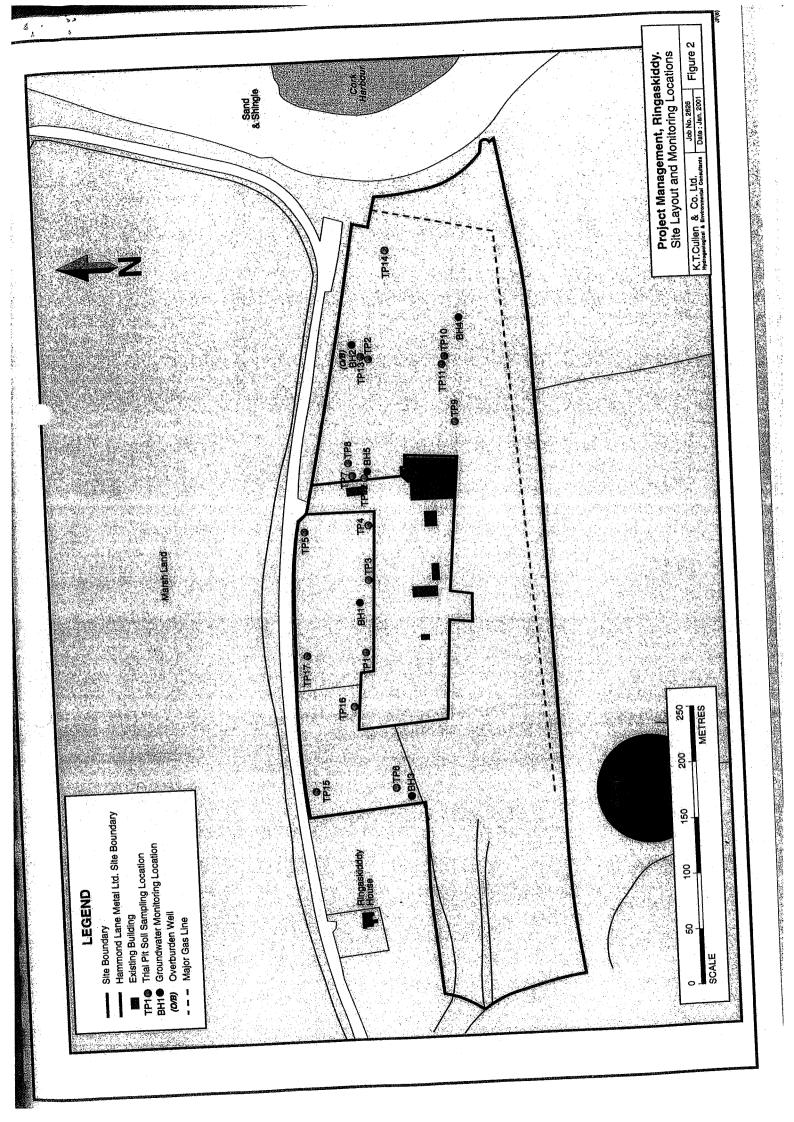
The land surface zoning is only as good as the information which is used in its compilation (geological mapping, hydrogeological assessment, etc.) and these are subject to revision as new information is produced. Therefore a scheme must be flexible and allow for regular revision.

Uncertainty is an inherent element in drawing geological boundaries and there is a degree of generalisation because of the map scales used. Therefore the scheme is not intended to give sufficient information for site-specific decisions. Also, where site specific data received by a regulatory body in the future are at variance with the maps, this does not undermine a scheme, but rather provides an opportunity to improve it.

FIGURES

and the second s





# **TABLES**

Table 1. Soil Analytical Results - PRO, DRO, Mineral Oil - PM, Ringaskiddy (Dec 2000)

								- WILL	10 July	Dutch MACs	MACs
		COL	TP3	TP 4	TPS	TiP 6	TP7	117	0.1	S-Value	I-Value
Location	14	33.0	0.1.5	0-3	0.3	0-3.2	7.0	Title C	mo/kp	mg/kg	mg/kg
Depth (m)	L.O mo/ko	me/kg	mg/kg	mg/kg	mg/kg	mg/kg	mg/kg	Swam	0.0		
					ų	7.6		•	78	1	
Discal Bange Organics	09	43	\$	 83	<del></del>	ī				05	2000
Diesei Mango Cigares	•	ç	g	6	13	∞	<b>8</b> 2	•	×o	3	
Mineral Oil	×,	CT.	}	č	ç	5	Z0:02	<0.01	<0.01	•	ı
Petrol Range Organics	<0.01	<0.01	<0.01	TO:00	70.9×	3					

K.T.Cullen & Co. Ltd Hydrogeological and Environmental Consultants

mg/kg - milligrams per kilogram
Dutch MACs - Dutch Maximum Admissible Concentration guidelines
S-Value= Target Value
I-Value= Intervention Value
Results are underlined where the Dutch S-MAC for Mineral Oil is exceeded.
"-" = Dutch MAC not available
"<" = Less than

Table 2. Soil Analytical Results - BTEX - PM Ringaskiddy (Dec 2000)

ΣL	S-Value 1-Value 0.05 1 0.05 130 0.05 50 0.05 25	K.T.Cullen & Co. Ltd Hydrogeological and Environmental Consultants
TP 10		K.T.Cullen & Co. Ltd Hydrogeological and E
TOT.	3-4:1 mg/kg <0.01 <0.01 <0.01	
	0.2 mg/kg 0.01 0.01 0.01	
	17P 6 0 - 3.2 mg/kg <0.01 <0.01 <0.01 <0.01	
	11P 5 0 - 3 11P 5 0 - 3 0 - 3 0 0 01 <0.01 <0.01	
	TP 4 0 - 3 mg/kg <0.01 <0.01	70.0>
		<0:01
	40.01 40.01 40.01	<0.01
		<0.01
	Location   TP 1   L8	
	Benzene Toluene Ethylbenzene	Xylene

Legendi.

mg/kg - milligrams per kilogram

mg/kg - milligrams per kilogram

Dutch MACs - Dutch Maximum Admissible Concentration guidelines

S.Value= Target Value

I.Value= Intervention Value

Results are underlined where the Dutch S-MAC for Mineral Oil is exceeded.

Results are underlined where the Dutch S-MAC for Mineral Oil is exceeded.

"." = Dutch MAC not available

"." = Less than

Trace Organics (VOCs)		TP1	TP3	TP4	TP5
	Depth (m)	1.8	0 - 1.5	0 - 3.0	0 - 3.0
	Units		4 ( 5		
Dichlorofluoromethane	ug/kg	<1	<1	<1	<1
Chloromethane	μg/kg	<1	<1	<1	<1
Vinylchloride	µg/kg	<1	<1	<1	<1
Bromomethane	µg/kg	<1	<1	<1	<1
Chloroethane	μg/kg	<1	<1	<1	<1
Trichlorofluoromethane	μg/kg	<1	<1	<1	<1
trans-1,2-Dichloroethene	μg/kg	<1	<1	<1	<1
Dichloromethane	μg/kg	<1	<1	<1	<1
1.1 Dichloroethene	μg/kg	<1	<1	<1	<1
1.1 Dichloroethane	μg/kg	<b>∢</b> 1	<1	<1	<1
cis-1,2-Dichloroethene	μg/kg	<1	<1	<1	<1
Bromochloromethane	µg/kg	<1	<1	<1	<1
Chloroform	μg/kg	<1	<1	<1	<1
2,2-Dichloropropane	µg/kg	<1	<1	<1	<1
1,2-Dichloroethane	ug/kg	<1	<1	<1	<1
1.1.1-Trichloroethane	ug/kg	<1	<1	<1	<1
1.1-Dichloropropene	μg/kg	<1	<1	<1	<1
Benzene	μg/kg	<1	<1	<1	<1
Carbontetrachloride	µg/kg	<1	<1	<1	<1_
Dibromomethane	µg/kg	<1	<1	<1	<1
1.2-Dichloropropane	μg/kg	<1	<1	<1	<1
Bromodichloromethane	μg/kg	<1	<1	<1	<1
Trichloroethene	μg/kg	<1	<1	<1	<1
cis-1,3-Dichloropropene	μg/kg	<1	<1	<1	<1
trans-1,3-Dichloropropene	μg/kg	<1	<1	<1	<1
1,1,2-Trichloroethane	μg/kg	<1	<1	<1	<1
Toluene	μg/kg	<1	<1	<1	<1
1,3-Dichloropropane	μg/kg	<1	<1<1	<1	<1
Dibromochloromethane	μg/kg	<1	<1	<1	<1
1.2-Dibromoethane	μg/kg	<1	<1	<1	<1
Tetrachloroethene	μg/kg	<1	<1	<1	<1<1_
1,1,1,2 -Tetrachloroethane	μg/kg	<1	<1	<1	<1
Chlorobenzene	μg/kg	<1	<1	<1	<1
Ethylbenzene	μg/kg	<1	<1	<1	<1

Dutch	MACs
S-Value	I-Value
-	-
-	_
_	100
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-	_
-	20,000
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-	<del>,</del> -
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-	-
-	4,000
-	-
-	-
. 50	1,000
<del>-</del>	-
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_	-
. 1	60,000
-	
-	-
50	130,000
-	+
-	1
10	4,000
	50 006
50	50,000

### LEGEND

µg/kg: micrograms per kilogram
MAC: Maximum Admissible Concentration

Dutch S-Value: Target Value
Dutch I-Value: Intervention Value -': MAC Guideline Not Available

<= Below current laboratory detection limit

10012		Lbi	TP3	1.14	CIT	THE PARTY OF THE P
Trace Organics (VUCS)		10	0-15	0-30	0.3.0	S-Value
	Depth (m)	1.0	2			
	Units					62
	tre/ke	⊽	7	⊽	⊽	3
o/m Xylenes	110/1/0	V	⊽	⊽	⊽	<u> </u>
Sromoform	94/60	V	▽	⊽	⊽	100
tyrene	PAE/AS		1	\	V	
2. Totrachloroethane	Lg/kg	√	7	/		
	TIE/KE	⊽	⊽	⊽	Ÿ	
0 - Aylene	110/kp	⊽	⊽	⊽	⊽	1
2,5-1racmoropropage	110/10	⊽	⊽	⊽	⊽	,
sopropylbenzene	4.00		V	⊽	⊽	•
Втоторепzепе	SV.ST	715		V	⊽	•
2-Chlorotoluene	HEIKE	7				
Promylhenzene	Lg/kg	⊽	7	7		
Chlorotolisane	LIES/Kg	⊽	⊽	7	7	
C. I. T. C. Charleson	Ug/kg	⊽	⊽	⊽	₹	_
A-1 I mem Anonyme	1/o/kg	⊽	7	⊽	7	•
-Isopropyitoniene	isa/ka	V	⊽	⊽	▽	•
3,5-1 rimethylbenzene	10.00 P		V	V	⊽	10
2.Dichlorobenzene	Sw.Sm	7 5	   	V	V	10
(.4-Dichlorobenzene	LEJ/KE	7				<u> </u>
secButylbenzene	Lg/kg	₹	<b>₹</b>	<b>小</b>	1	ŀ
art-Rutvlhenzene	LIG/Kg	7	⊽	Ž	7	١
2 Dinklanahanzene	ug/kg	⊽	⊽	⊽	7	3
	uo/ke	⊽	⊽	⊽	⊽	•
n-Butyinenzene	110/kg	V	⊽	⊽	7	
2-Dibromo-3-Cinoropi opane	27/23		V	V	⊽	10
1,2,4-Trichlorobenzene	HEAT P			7	⊽	
Naphthalene	HEJ.K.B	7	  -  -		K	
2 3-frichlorobenzene	Lg/kg	⊽	7	7		
	180/69	⊽	⊽	<b>∀</b>	7	

LEGEND.

µg/kg: micrograms per kilogram

µg/kg: micrograms per kilogram

MAC: Maximum Admissible Concentration

Dutch S-Value: Target Value

Dutch I-Value: Intervention Value

-'; MAC Guideline Not Available

< = Below current laboratory detection limit

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Soil Analytical Results	
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Table 4a.	
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		TP1	TP2	TP3	TP4	TPS	TP6	T.b.	TP7	TP10		
Parameters	Depth (m)	1.8	0 - 5.5	0-1.5	6-3	0-3	0-3.2	0-2	3-4.1	0-1	Dutch MA	<b>Dutch MAC Values</b>
	Units										S-Value	I-Value
Acenaphthene	hg/kg	24		95	14	32	⊽	⊽	₽	55		
Acenaphthylene	µg/kg	⊽	7	⊽	⊽	⊽	⊽	⊽	⊽	340		
Benzo(B)fluoranthene	lug/kg	30	[49	2.7	7	27	26	ī×	31	1142		-
Dibenz(AH)anthracene	µg/kg	⊽	₽	.<1	⊽	⊽	⊽	⊽	Ī	⊽		
Fluorene	Lg/kg	▼ _	29	4	∀1	⊽	⊽	₹	  -	250		
Pyrene	Lg/kg	244	308	202	⊽	⊽	⊽	⊽	95	6321		
PAHs included in 'PAH (Sum of 10)' Dutch S and IMAC val	m of 10)' Dutch	S and I MA		tes for PAHs in soil								
Anthracene	ng/kg	25	25	25	2	14	17	1	14	862	•	
Benzo(a)anthracene	Lig/kg	113	183	147	1		66	⊽	121	1081		
Benzo(a)pyrene	µg/kg	1.2	131	14	7	31	14	⊽	27	245	•	
Benzo(ghi)perylene	hg/kg	7	⊽	⊽	ľ>	⊽	⊽	⊽	⊽	⊽		1
Benzo(k)flouranthene	µg/kg	29	27.7	22	7	31	22	⊽	30	6/6	•	-
Chrysene	µg/kg	65	295	6	<b>7</b>	83	7.4	⊽	6	3064		
Fluoranthene	l µg/kg	261	281	216	<b>I&gt;</b>	7	⊽	⊽	100	5804	,	-
Indeno(123-cd)pyrene	l µg/kg	7	[>		<b>[&gt;</b>	⊽	⊽	⊽	⊽	⊽	•	·
Naphthalene	l µg/kg	21.12	2153	2731	1932	1190	2291	2162	1821	10024		-
Phenanthrene	lug/kg	149	152	203	86	112	119	95	14	7221		٠
PAH (Sum of 10)	ug/kg	2768	3499	3457	2042	1530	2636	2271	2356	29282	1000	40000
PAH (Total)	l µg/kg	3066	4010	3741	2049	1588	2659	2264	2480	37388	,	

ug/kg: micrograms per kilogram
MAC: Maximum admissable concentration
S-level: Dutch guideline for normal uncontaminated soil
I-Level: Dutch guideline for Intervention
Results awaiting confirmation
"-": MAC not available
<= below laboratory detection limit

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Table 4b Soil Ansiving Results	ï
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		TP11	TP11	TP12	TP12	TP13	TP13	TP14	TP15	TP16	TP17		
Parameters	Depth (m)	0 <del>.</del> 1	1-3	0-1	1-2.6	0-1	1-3.4	0-3.8	0-2	0-2.5	0-2	Dutch M.	Dutch MAC Values
	Inits											S-Value	I-Value
Acensahthene	119/kg		V	99	4	45	45	2	4	2	26		٠
Acenanhthylene	119/kg		V	37	⊽	I>	13	⊽	V	ľ	2	•	•
Renzo(R)finoranthene	Hø/kg	V	v	₽	⊽	ĪV	٧	ĪŸ	V	⊽	Į,	•	•
Dibenzi A Hanthracene	119/kg	2	V	▽	⊽	⊽	⊽	⊽	⊽	⊽	[>	•	•
Hilosepe	119//89		V	5	⊽	⊽	1	⊽	⊽		_	•	,
Pyrana	110/kg	>	V	6	V	V	2	⊽	F	1	4	•	•
DAHs included in 'PAH (Sum of 10)' Dutch Sand I WAC values for	um of 100' Dutch	S and I MA		PAHs in soil									
Anthracene	Horke	2		2	⊽	⊽	3	4	⊽	⊽	2	•	•
Benzo(a)anthracene	Uo/kg		⊽	⊽	⊽	₽	3	⊽	⊽	<b> </b>	1	٠	·
Benzo(a)nvrene	110/kg	V	V	v	⊽	⊽	7	⊽	⊽	V	⊽	1	1
Benzo(chi)nervlene	110/160	2	V	V	⊽	V		⊽	⊽	⊽	V		1
Denzo (Inflamentanthane	Halen Halen		V	V	V	V	ĪV	V	⊽	V	v		,
Chrysene	llø/kø	9	V	2	V	⊽	3	⊽	⊽	-	4	1	•
Fluoranthene	11g/kg	82		91	4	2	∞	_	2	9	13	,	ŧ
Indenof [23-cd)pyrene	ue/kg	2	⊽	V	⊽	⊽	1	<b> </b>	₽	<		•	•
Naphthalene	ug/kg	2	4	*	2	3	œ	⊽		⊽	2	•	•
Phenanthrene	ug/kg	<u>«</u>	4	20	7	5	61	4	4	7	23	-	
PAH (Sum of 10)	ug/kg	54	9	87	13	1.0	46	6	7	4	45	900	40000
PAH (Total)	ug/kg	2	9	651	U	55	107	=	22		200		•

MAC: Maximum admissable concentration
S-level: Dutch guideline for normal uncontaminated soil
I-Level: Dutch guideline for Intervention
"-": MAC not available
<= below laboratory detection limit

Sample	Depth (m)	Arsenic	Cadmium	Chromium ms/ke	Copper mg/kg	Mercury mg/kg	Nickel mg/kg	Lead mg/kg	Selenium mg/kg	Zinc mg/kg	ng/kg
TDI	8-	11	V	4	15	⊽	26	48	⊽	83	<0.01
TP7	0.55	14	, v	15	61	⊽	24	12	⊽	95	<0.01
TP3	0-1.5	16	⊽	20	17	⊽	32	£1 ^	⊽	92	90.04
TP4	0-3	17	⊽	18	24	⊽	25	6	⊽	76	<0.01
TP5	0-3	13	⊽	91	18	⊽	20	01	⊽	28	<0.01
TP6	0 - 3.2	91	⊽	15	21	⊽	29	17	⊽	126	<0.01
14L	0.2	41	⊽	18	16	7	19	[3	7	62	<0.01
TP-7	3-4.1	15	⊽	15	18	⊽	23	10	~	62	<0.01
TP-10	0-1	15	⊽	43	25	⊽	23	10	⊽	76	<0.01
Outsh MAC C Values	90	20	9,8	001	36	0.3	35	85	ť	140	0.05
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Dutch MAC I Values

mg/kg. milligrams per kilogram

MAC: Dutch Standard Maximum Admissible Concentration

S Value: Dutch Guidline for normal uncontaminated soil

I Value: Dutch Guideline for Intervention

".": MAC Guideline not available

n.a. = not analysed

"<" = below detection limit

Pesticide	Units	TP 1	Z4I	TP3	TP-4	TP 5	TP 6	TP 7	TP7	TP 10	Dutch Values	Values
	Depth (m)	1.8	0.5.5	0-1.5	0-3	0-3	0-3.2	0.2	3 - 4.1	0 - 1	S- Value	I Value
Dichlorvos	ug/kg	<1	[ <b> </b>		>	<1	<1	<1	<1	[>	•	٠
Mevinphos	µg/kg	<ا		⊽	<1	<1	<b> </b>	<1	[×	⊽	•	1
Phorate	µg/kg	<1	⊳	₽	<b> </b>	<1	<1	<1	<1	<b> </b>	1	
Alpha-BHC	ug/kg	ľ>	I>	<1	[>	<b>~</b> 1	I>	₽	[∨	⊽	2.5	٠
Beta-BHC	µg/kg	⊽	I>	⊽	<b>-</b> 1>	7	⊽	⊽	!∨	V	1	ı
Gamma-BHC	µg/kg		>	>	<1	√1		<b>~</b> 1	<b> </b>	⊽	0.05	
Diazinon	µg/kg	٦	l>	7	<1	⊽	[>		√1		•	
Disulfoton	l µg/kg	7	I>	⊽	7		⊽	7	⊽	⊽	1	٠
Delta-BHC	ug/kg	7	⊽	>	√1	⊽	⊽	⊽	⊽	⊽	٠	,
Methyl Parathion	ug/kg		1>	-   	<1 <1	√	[∨	, .  >	⊽	īV	•	
Heptachlor	hg/kg	⊽	1>	<1	٧	⊽	⊽	1>	[>	⊽	•	
Fenitrothion	mg/kg	7	P	<1	[>	⊽	⊽	∀		⊽	•	1
Aldrin	ng/kg	[>	⊽	₽	[>	₽	⊽	⊽	⊽	⊽	2.5	•
Malathion	ug/kg	7	!>	 	ŀ	<b> </b>	⊽	7	⊽	⊽	•	
Parathion	lug/kg	۷	1>		<1	<br </td <td>7</td> <td><b>√</b>1</td> <td>⊽</td> <td>⊽</td> <td>•</td> <td>٠</td>	7	<b>√</b> 1	⊽	⊽	•	٠
Heptachlor Epoxide	µg/kg	کا	₽	>	<b>~</b> 1	<b>~</b>	7		7	7		٠
Endosulfan I	µg/kg		<b>!&gt;</b>	₽	<1	<1	⊽	⊽	⊽	⊽	•	•
Dieldrin	ug/kg	₽	7>		<1	<1	<1	 		⊽	0.5	١
4,4-DDE	µg/kg	<b>~</b> 1	⊽	<ا	۲	>	⊽	</td <td>   -</td> <td>⊽</td> <td>2.5</td> <td>4000</td>	  -	⊽	2.5	4000
Endrin Ketone	µg/kg		⊳	<1	⊽	7	⊽	<b>'</b>	<b> </b>   ∨	⊽	•	1
Endosulfan II	µg/kg	⊽	I>	7	<1		[>	۱۷	₽	⊽	١	٠
4,4-DDD	µg/kg		⊽	7	<1	<1	<b>~</b> 1	<1	<b> </b>	ī	2.5	4000
Ethion	ug/kg	<b>~</b>	 	⊳	[>	<b> </b>	[>	~ 1	<ا	⊽.	•	
Endrin	l µg/kg	۲	I>	₽	Ī	⊽	ī	⊽	⊽	Į,		1
Endosulfan Sulphate	l ug/kg	⊽	₽	⊽	<1	<1	<b>~</b> 1	₽	[>	⊽	1	1
4,4-DDT	µg/kg	<b>~</b> 1	.[>		<1	<1	[>	. [>	אן >	⊽	2.5	4000
Methoxychlor	µg/kg	⊽	⊽	⊽	<b> </b>	<1	<b>~</b> 1	<b> </b>	<b>~</b> 1	[V		
Azinphos Methyl	µg/kg	⊽	⊽	⊽	V	<1	</td <td></td> <td>ľ</td> <td>7</td> <td>1</td> <td></td>		ľ	7	1	

Legend

ug/kg: micrograms per kilogram
MAC: Maximum Admissable Concentration
S-level: Dutch guideline for normal uncontaminated soil
I-Level: Dutch guideline for Intervention
: MAC not available
< = below laboratory detection limit

	Г	_	Γ-			_	_	_			_
	C Values	I-Vales			•			ı			1000
	Dutch MA	S-Value I-Vales	1	,	•	1	1				20
	TP10	0 - 1		V	⊽	⊽	⊽	⊽	⊽	V	⊽
	TP7	3-4.1		<1	ï	⊽	⊽	⊽	⊽	⊽	⊽
	TP7	0-2		⊽	⊽	23	⊽	23	27	25	88
	TP 6	0 - 3.2		⊽	⊽	⊽	⊽	⊽	[ <b>&gt;</b>	<1	⊽
•	TP 5	0-3		⊽	⊽	₽	⊽	⊽	√	⊽	₽
	TP 4	0-3		⊽		  >	- 1>	⊽	₽	⊽	  >
	TP 3	0-1.5		>	1>	I>	<b>!&gt;</b>	1>	7	ī <b>∨</b>	[>
	TP 2	0.5.5		₽	<u> </u>	20	14	182	214	183	643
	TP 1	1.8		₽	₽	⊽	ŀ	⊽		   	- 1>
		Depth (m)	Units	ug/kg	ng/kg	ng/kg	ng/kg	ng/kg	ug/kg	ng/kg	ug/kg
	Parameters			PCB Congener 28	PCB Congener 52	PCB Congener 101	PCB Congener 118	PCB Congener 153	PCB Congener 138	PCB Congener 180	PCB total

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ug/kg: micrograms per kilogram
MAC: Maximum admissable concentration
S-level: Dutch guideline for normal uncontaminated soil
I-Level: Dutch guideline for Intervention
-: MAC not available
<= below laboratory detection limit

Parameters		TP 2	TP7	Dittch M	AC Volume
	Depth (m)	0-5.5	0-2	S.Value	T Volos
	Units			anne.	SIRAT
PCB Congener 28	ug/kg	-	12		•
PCB Congener 52	ug/kg	V			
PCB Congener 101	Le/kg	V	i V		
PCB Congener 118	ile/ke	V		1	
PCB Congener 153	ug/kg	3	-		
	LIP/Kg	4			
PCB Congener 180	Lg/kg	5	Ī		•
PCB total	ng/kg	13	6	١	1000

Lg/kg: micrograms per kilogram
MAC: Maximum admissable concentration
S-level: Dutch guideline for normal uncontaminated soil
I-Level: Dutch guideline for Intervention
-: MAC not available
<=below laboratory detection limit

- ---

Location	图1	BH 2	BH3	BH 4	BHS		Dutch	MACs
Units	l/g/l	l/g/l	Lg/l	/an	/vii	, ,	S-Value I-Valu	I-Valı
					mg/.	-,-	mg/1	Lg/I
Diesel Range Organics	<10	<10	<10	√10	<10			
Mineral Oil	5	<u> </u>	,					1
	) /	217	010	~ V	<10		20	909
Petrol Range Organics	010	V V V	<10	7	ç			
			}	27	OT>	1	ı	

I-Value hg/l

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ugl - micrograms per Litre
Dutch MACs - Dutch Maximum Admissible Concentration guidelines
S-Value= Target Value
I-Value= Intervention Value
"-" = Dutch MAC not available
"<" = Less than

	Location	BIE 1	BH 2	BH3	BH 4	BHS		Dutch MACs	Cs
	Units	l/grl	Lg/l	/gri	Vän	Van	S-Value µg/l		I-Value µg∕l
Benzene		<10	<10	<10	<10	<10	0.20		30
Toluene		<10	<10	<10	<10	<10	0.20		0001
Ethylbenzene		<10	<10	<10	<10	<10	0.20		150
Xylene		o10	<10	<10	<10	<10	0.20		70
							-		

Legend:

μg/l - micrograms per Litre Dutch MACs - Dutch Maximum Admissible Concentration guidelines

K.T.Cullen & Co. Ltd Hydrogeological and Environmental Consultants

S-Value= Target Value I-Value= Intervention Value

Results are underlined where the Dutch S-MAC for Mineral Oil is exceeded.
"-" = Dutch MAC not available
"<" = Less than

Trace Organics (VOCs)	Location	BH 1	BH 2	BH 3	BH 4	BH 5
	Units					
Dichlorofluoromethane	μg/l	<1	<1	<1	<1	<1
Chloromethane	μg/l	<1	<1	<1	<1	<1
Vinylchloride	μg/l	<1	<1	<1	<1	<1
Bromomethane	μg/l	<1	<1	<1	<1	<1
Chloroethane	μg/l	<1	<1	<1	<1	<1
Trichlorofluoromethane	µg/l	<1	<1	<1	<1	<1
trans-1,2-Dichloroethene	μg/l	<1	<1	<1	<l< td=""><td>&lt;1</td></l<>	<1
Dichloromethane	μg/Ι	<1	<1	<1	<1	<1
1,1 Dichloroethene	μg/l	<1	<1	<1	<1	<1
1,1 Dichloroethane	μg/l	<b>&lt;</b> 1	<1	<1	<1	<1
cis-1,2-Dichloroethene	μg/l	<1	<1	<1	<l< td=""><td>&lt;1</td></l<>	<1
Bromochloromethane	μg/1	<1	<1	<1	<1	<1
Chloroform	µg/I	<1	<1	<1	<1	<1
2,2-Dichloropropane	μg/l	<1	<1	<1	<1	<1
1,2-Dichloroethane	μg/l	<1	<1	<1	<1	<1
1,1,1-Trichloroethane	μg/l	<1	<1	<1	<1	<1
1,1-Dichloropropene	μg/l	<1	<1	<1	<1	<1
Benzene	μg/l	<1	<1	<1	<1	<1
Carbontetrachloride	μg/l	<1	<1	<1	<1	<1
Dibromomethane	μg/l	<1	<1	<1	<1	<1
1,2-Dichloropropane	μg/l	<1	<1	<1	<1	<1
Bromodichloromethane	μg/l	<1	<1	<1	<1	<1
Trichloroethene	μg/l	<1	<1	<1	<1	<1
cis-1,3-Dichloropropene	μg/l	<1	. <1	<1	<1	<1
trans-1,3-Dichloropropene	μg/l	<1	<1	<1	<1	<1
1,1,2-Trichloroethane	μg/l	<1	<1	<1	<1	<1
Toluene	μg/l	<1	<1	<1	<1	<1
1,3-Dichloropropane	μg/l	<1	<1	<1	<1	<1
Dibromochloromethane	μg/l	<1	<1	<1	<1	<1
1,2-Dibromoethane	μg/l	<1	<1	<1	<1	<1
Tetrachloroethene	μg/l	<1	<1	<1	<1	<1
1,1,1,2 -Tetrachloroethane	μg/l	<1	<l< td=""><td>&lt;1</td><td>&lt;1</td><td>&lt;1</td></l<>	<1	<1	<1
Chlorobenzene	μg/l	<1	<b>⊲1</b>	<1	<1	<1
Ethylbenzene	µg/l	<1	<b>V</b> I	<1	<1	<1

Dutch	MACs
S-Value	I-Value
-	-
-	-
-	-
-	-
-	-
-	-
-	-
-	<b>-</b> 3
<u> </u>	• 4
-	•
-	-
0.01	50
0.01	400
- '	-
	•
0.01	50
-	
0.01	50
	-
-	
-	_
0.20	- 30
0.20	30
-	•
<del></del>	
_	•
_	-
_ ::	

## **LEGEND**

μg/l: micrograms per litre
MAC: Maximum Admissible Concentration
Dutch S-Value: Target Value

**Dutch I-Value: Intervention Value** -': MAC Guideline Not Available

<= Below current laboratory detection limit

Trace Organics (VOCs)	Location	BH 1	BH 2	BH 3	BH 4	BH 5
Trace Organies (1005)	Units					
p/m Xylenes	μg/l	<1	<1	<1	<1	<1
Bromoform	µg/l	<1	<1	<1	<1	<1
Styrene	μg/l	<1	<1	<1	<1	<1
1,1,2,2-Tetrachloroethane	μg/l	<1	<1	<1	<1	<1
o - Xylene	μg/i	<1	<1	<1	<1	<1
1,2,3-Trichloropropane	μg/l	<1	<1	<1	<1	<1
Isopropylbenzene	μg/l	<1	<1	<1	ব	<1
Bromobenzene	μg/l	<1	<1	<1	<1	<1
2-Chlorotoluene	μg/1	<1	<1	<1	<1	<1
Propylbenzene	μg/l	<1	<1	<1	<1	<1
4-Chlorotoluene	μg/I	<1.	<1	<1	<1	<1
1,2,4-Trimethylbenzene	μg/l	<1	<1	<1	<1	<1
<b>A ™sopropyltoluene</b>	μg/l	<1	<1	<1	_<1	<1
5-Trimethylbenzene	με/Ι	<b>~1</b>	<1	<1	<1	<1
1.2-Dichlorobenzene	μg/l	<1	<1	<1	<1	<1
1,4-Dichlorobenzene	μg/l	<1	<1	<1	<1	<1
sec-Butylbenzene	μg/l	<1	<1	<1	<1	<1
tert-Butylbenzene	µg/l	<1	<1	<1	<1	<1
1.3-Dichlorobenzene	μg/l	<1	<1	<1	<1	<1
n-Butylbenzene	μg/l	<1	<1	<1	<1	<1
1,2-Dibromo-3-Chloropropane	μg/l	<1	<1	<1	<1	<1
1,2,4-Trichlorobenzene	µg/l	<1	<1	<1	<1	<1
Naphthalene	μg/l	<1	<1	<1	<1	<1
1,2,3-trichlorobenzene	μg/l	<1	<1	<1	<1	<1
Hexachlorobutadiene	μg/l	<1	<1	<1	<1	<1

Dutch	MACs
S-Value	I-Value
0-Value	
	-
-	-
-	-
_	-
_	_
0.2	150
_	-
_	, -
	<b>-</b>
- · ·	-
-	-
	-
0.2	70
	-
0.5	300
-	
0.01	40
0.2	1000
	-
-	
-	
	-
-	0.7

### **LEGEND**

µg/l: micrograms per litre
MAC: Maximum Admissible Concentration

Dutch S-Value: Target Value Dutch I-Value: Intervention Value -: MAC Guideline Not Available

< = Below current laboratory detection limit

				,				
		BH 1	PH2	BH 3	BH 4	BHS		
Parameters					6		Dutch MAC Value	C Value
	Units						S-Value	I-Val
Acenaphthene	l/gu	131	<10	135	<10	<10	•	
Acenaphthylene	l/gu	28	21	285	<10	25	-1	•
Benzo(B)fluoranthene	l/gu	<10	<10	<10	<10	<10	•	
Dibenz(AH)anthracene	l/gu	<10 <10	<10	<10	<10	<10	•	
Fluorene	l/gu	<10	<10	<10	<10	<10	1	•
Pyrene	l/gu	<10	19	<b>0</b> 1>	<10	<10	,	•
PAHs included in 'PAH (Sur	Sum of 10) Dutch S and I MA	Sand I MA	C values for PAHs in soil	PAHs in soil				
Anthracene	//gu	20	17	19	11	<10	20	200
Benzo(a)anthracene	l/gu	<10	<10	<10	<10	<10	2	200
Benzo(a)pyrene	l/gu	<10 <10	<10	<10	<10	<10	1	20
Benzo(ghi)perylene	l/gu	<10	<10	<10	<10	<10	0.2	50
Benzo(k)flouranthene	l/gu	<10	<10	<10	<10	<10	0.2	20
Chrysene	l/gu	<10	<10	<10	<10	<10	2	50
Fluoranthene	l/ga	<10	26	<10	<10	<10	5	100
Indeno(123-cd)pyrene	∕8u	<10	<10	<10	<10	<10	9.0	20
Naphthalene	l/gu	35	19	36	65	25	100	)00/
Phenanthrene	l/gu	13	43	11	14	18	20	005

ng/l: nanograms per litre
MAC: Maximum admissable concentration
S-level: Dutch guideline for normal uncontaminated groundwater
L-Level: Dutch guideline for Intervention
"-": MAC not available
<= below laboratory detection limit

Commission	Arconio	Codminm	Chromium	Copper	Mercury	Nickel	Lead	Selenium	Zinc
Sample	mg/l	mg/l	mg/l	mg/l	mg/l	mg/l	mg/l	mg/l	mg/l
BH 1	<0.05	<0.05	<0.05	<0.05	<0.05	<0,05	<0.05	<0.1	<0.05
BH 2	90.0	<0.05	<0.05	<0.05	<0.05	<0.05	<0.05	<0.1	<0.05
ВН 3	<0.05	<0.05	<0.05	<0.05	<0.05	<0.05	<0.05	<0.1	<0.05
BH 4	<0.05	<0.05	<0.05	<0.05	<0.05	<0.05	<0.05	<0.1	<0.05
BH 5	<0.05	<0.05	<0.05	<0.05	<0.05	<0.05	<0.05	<0.1	<0.05
		Z							
Dutch MAC S Values	0.01	0.0004	0.001	0:03	0.005	0,015	0.015		0.065
Dutch MAC I Values	90.0	900'0	0.03	0.075	0.003	0.075	0.075		0.80

mg/l: milligrams per Litre

MAC: Dutch Standard Maximum Admissible Concentration

S Value: Dutch Guidline for normal uncontaminated soil

I Value: Dutch Guideline for Intervention

".": MAC Guideline not available

n.a. = not analysed

"<" = below detection limit

Pesticide Location  Dichlorvos Lug/l Mevinphos Lug/l Mevinphos Lug/l Alpha-BHC Lug/l Beta-BHC Lug/l Gamma-BHC Lug/l Diazinon Lug/l Diazinon Lug/l Diazinon Lug/l Diazinon Lug/l Herrachlor Lug/l Diazinon Lug/l Diazinon Lug/l Diazinon Lug/l Delta-BHC Lug/l	Ocation Units Units UNITS US/I US/I US/I US/I US/I US/I US/I US/		##	MH V V V V V V V V V V V V V V V V V V V	BH 4	BHS	Dutch	Dutch Values
IC athion		<b>30000000</b>	<b>7777777</b>	7777777	v			T Wolus
4C athion	222222	<b>33377777</b>	<b>7777777</b>	<b>777777</b>	V		S- Value	T Value
IC athion	555555	<b>7 7 7 7 7 7 7 7</b>	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	<b>7777</b>		<b> </b>		•
IC athion	555555	<b>VVVVV</b>	<b>V</b> VVVV	7777	>	7	1	-
IC athion	5,5,5,5,5	<b>777777</b>	\[ \sqrt{\sq}}}}}}}\sqrt{\sqrt{\sqrt{\sqrt{\sqrt{\sq}}}}}}}\signtifien\sintitite{\sqrt{\sqrt{\sqrt{\sqrt{\sqrt{\sqrt{\sqrt{\sqrt{\sqrt{\sqrt{\sqrt{\sq}}}}}}}}}\sqrt{\sqrt{\sqrt{\sqrt{\sqrt{\sqrt{\sqrt{\sq}}}}}}}}}}}\signtifien\sqrt{\sqrt{\sqrt{\sqrt{\sqrt{\sq}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}}	V V V		7		•
4C athion		<b>77777</b>	<b>7777</b>	⊽⊽,	⊽	7	•	•
4C athion		ママママ	⊽⊽⊽	⊽ ₹	Į>	[>	-	
ion		⊽⊽⊽	⊽⊽	1	⊽	⊽	200	
athion	1/3	₽₽₽	- √	7	<1	₹		-
athion	/8	 		-   	<b>~</b> 1	  -		
		<1	⊽	⊽	7	7	•	•
	1/8		< <u> </u>	-   	<b>1</b> >	⊽	•	•
	1/S11	-	<b>1&gt;</b>	!>	[>	⊽	•	•
Fenitrothion   µg/	g/l	-  -	<1	<b>-</b> 1>	<1	< <u>1</u>	•	
	1/8	  >	<1	<1	<1	<b> </b>		-
Malathion   µg/l	[ J/B	<b> </b>	<1	<1	<1	<b>~</b> 1	•	*
	] [/8	<1	<1	<1	<1	7	-	1
Heptachlor Epoxide   µg/1	1/8	>	<b>1</b> >	<1		<b>V</b>	•	-
an I	1/8	<1	<1	<1	</td <td>[&gt;</td> <td>•</td> <td>•</td>	[>	•	•
Dieldrin µg/	1/3	<1	<1	<1	<1	<b>!</b>	70	1
4,4-DDE   μg	[/S	<	<b>!</b> >	\   	<1	7	•	t
Endrin Ketone   µg/l	[/S	-  -	<1	[>	>	√1		•
an II	[/5	<1	-1>	<1	<1	\   	•	1
(4,4-DDD	1/8	⊽	7	[∨	⊽	\ \	•	1
Ethion   µg/l	[] []	[>	<1	-\  -	</td <td>⊽</td> <td>•</td> <td>ı</td>	⊽	•	ı
Endrin Lg	/6	<1	\r	[>	V	✓		1
Endosulfan Sulphate   µg/l	J/8	حا	<b>~</b> 1	7	ī	 	•	•
4,4-DDT	/8	\   \	√1	\   	<b>~</b>	>		1
	g/l	. <u>.</u>	⊽		⊽			1
Azinphos Methyl   µg	l/grl	∇	⊽	∨	⊽	⊽	•	,

Legend

µg/l: micrograms per litre

MAC: Maximum Admissable Concentration
S-level: Dutch guideline for normal uncontaminated soil
I-Level: Dutch guideline for Intervention
-: MAC not available
< = below laboratory detection limit

Table 14. Groundwater Analytical Results - PCBs - PM, Ringaskiddy (Dec 2000)

	Parameters	Location	TP1	TP 2	TP 3	TP 4	TPS	Dutch MA	E
Units								TAT TOATO	זנ
ner 28 ner 52 ner 101 ner 118 ner 153 ter 138		Units						o-vaine	I-vales
ner 52 ner 101 ner 118 ner 153 ner 153 ner 138	뎔	l/g/l	⊽	Ī▽	V	1	1	•	
ner 101 ner 118 ner 153 ner 138 ner 180	ē	[/ā1]	V	V	7		7 5	•	ı.
ner 118 ner 153 ner 138 ner 180	her	[/ell		1	7  7	7	7	•	•
ner 113 ner 138 ner 180		192	7	7	7	\ <u>\</u>	⊽		
ner 153 ner 138 ner 180	힠	ug/1	<b>∀</b>	⊽	⊽	⊽	⊽		
ner 138 ner 180	161	l/gn	⊽'	₹	V	12	1		
or 180	ner	l/an	⊽	V	7				1
	lی	1/6/1	V		   	7	7		•
	1	1/2/1	 		7	7	7		1
	1000	12871	7	₹	⊽	▽	⊽	•	

ug/kg: micrograms per Litre
MAC: Maximum admissable concentration
S-level: Dutch guideline for normal uncontaminated groundwater
I-Level: Dutch guideline for Intervention
-: MAC not available
< = below laboratory detection limit

Г																								_
	POTABLE WATER M.A.C.			0-9	C - 0	1500	n.a.	п.а.	<0.05	<0.05	200	200	20	20	•	150	12	50	0.1	250	250	0.3	0.0005	
	BH 5			8 4	o.;	825	238	180	<0.05	<0.05	89	<0.05	<u>-</u>	0.38	0	93	2.6	3.7	<0.05	132	8	1.0	<0.01	
		-			۰.																			_
	BH 4			0 7	0.0	708	154	250	<0.05	<0.05	46	<0.05	· ∞	0.05	0	29	1.8	29.4	0.1	49	56	1.9	<0.01	
i																		-				<u> </u>		-
	внз			7	0./	892	203	270	<0.05	<0.05	99	<0.05	∞	<0.05	0	89	3.9	7.4	0.1	94	71	1.3	<0.01	
	BH 2			0	, V	407	328	250	<0.05	0.05	06	<0.05	22	0.05	C	56	7.0	3.0	<0.05	145	105	1.2	<0.01	
		L	····										· · ·			-,	<u></u>	<u> </u>				-		-
	BH 1			t	ν,	1002	342	100	C	<0.05	75	<0.05	y	0.05	6.05 50.05	33	3.1	11.3	<0.05	230	24	1.4	<0.01	
	UNIT			•	units	uS/cm	CaCO3 mg/l	CaCO3 ma/l	mo/l	L/SIII	mo/l	me/l	Ш6/1	me/l	mo/l	Na me/l	K me/l	NO3 mg/l	NO2 mg/l	Cl mg/l	SO4 mg/l	NH4 mg/l	mg/l	
	PARAMETERS			j	Ha	Conductivity	Total Hardness	Total Alkalinity	Aluminium	Boron	Coloinm	Tron	Mamerium	Manganese	Dhoenhorous	Sodium	Potassium	Nitrate	Strits	Chloride	Sulphate	Ammonia	Total Phenols	_

LEGEND

M.A.C. = Maximum Admissible Concentration under Drinking Water Regulation S.I. No. 81 of 1988
"-": MAC not available
<= Less Than
mg/l = milligram per litre

Rutland Street

Cork



# HYDROGEOLOGICAL ASSESSMENT HAMMOND LANE METAL COMPANY RINGASKIDDY CORK

# Prepared For: -

Hammond Lane Metal Company, Ringaskiddy, County Cork.

# Prepared By: -

O' Callaghan Moran & Associates, Granary House, Rutland Street, Cork.

23<sup>rd</sup> August 2011

# TABLE OF CONTENTS

# **PAGE**

1	INT	FRODUCTION	1
2	SIT	TE DESCRIPTION	2
	2.1	SITE LOCATION	
	2.2	SITE HISTORY	
	2.3	SITE LAYOUT	
	2.4	GEOLOGY AND HYDROGEOLOGY	
	2.4.1		
	2.4.2		
	2.5	Hydrogeology	7
	2.5.1	l Aquifer Classification	7
	2.5.2	2 Aquifer Vulnerability	10
	2.5.3		
	2.5.4	4 Groundwater Flow Direction	10
	2.5.5	Neighbouring Wells and Karst Features	10
	2.5.6	5 Designated Areas	10
	2.6	CONCEPTUAL MODEL	11
3	GRO	OUNDWATER INVESTIGATION	14
	3.1	WELL INSTALLATION	
	3.2	GROUNDWATER SAMPLING	
	3.3	GROUNDWATER ANALYSIS	
	3.4	DATA INTERPRETATION	
4		NCLUSIONS AND RECOMMENDATIONS	
-	4.1	Conclusions	
		RECOMMENDATIONS	
			11
		D	
<b>A</b> .	PPEND	OCM 2010 Report	
4	PPEND	DIX 2 Borehole Logs	
<b>4</b> ]	PPEND	DIX 3 - OCM Sampling Protocol	
<b>A</b> .	PPEND	DIX 4 - Laboratory Results	

# 1 INTRODUCTION

Hammond Lane Metal Company commissioned O'Callaghan Moran & Associates (OCM) to carry out a hydrogeological assessment of their scrap metal processing facility in Ringaskiddy. The facility operates under Waste Permit No WFP-CK-10-0077-02 issued by Cork County Council in 2010.

The assessment was requested by the planning authority as part of a planning application to expand the facility operations. The objective is to determine if there are any impacts on groundwater quality associated with the past use of the site.

An initial assessment of the subsoils within the site boundary, undertaken in 1997 by Mayer Environmental, did not identify the presence of any significant impacts on the subsoils. In 2010 OCM completed a further assessment of the subsoils, which confirmed that there was no evidence of any impact on the soils associated with site operations. The 2010 OCM Report is included in Appendix 1.

The 1997 and 2010 investigations were confined to the investigation of the soils and did not include an assessment of groundwater quality beneath the site. As there were no on-site monitoring wells, the hydrogeological assessment involved the installation and monitoring of four groundwater wells.

This report describes the well installation and groundwater monitoring programme. It also presents an update of site operations and revises the environmental risk assessment, based on the additional monitoring data.

# 2 SITE DESCRIPTION

# 2.1 Site Location

The site location and layout is shown on the Figure 2.1. The site encompasses approximately 1 hectare (ha) and is benched into the hillside to the south of the main Ringaskiddy to Haulbowline Island road.

Immediately to the north, the ground slopes to a low lying field which is the site of the proposed incinerator and is currently used for tillage. The Naval Maritime College is across the public road to the north and further north is Cork Harbour. The bridge crossing to Haulbowline Island is to the northeast. The lands to the west comprise open scrub land, overgrown with furze. There is an ESB Sub-Station adjacent to the northeastern site boundary.

# 2.2 Site History

Hammond Land Metal Company developed the facility in 1989 on a greenfield site previously used for agricultural purposes and the site has always been used as a metal processing facility since that time.

# 2.3 Site Layout

The site is accessed from the public road to the north. There is a weighbridge on the access road, with a site office to the east. The office houses administration, canteen, stores and toilets facilities. Sanitary wastewater discharges to a septic tank and percolation area located to the north of the building. The system was installed when the site was initially developed and Hammond Lane Metal Company informed OCM that it has always operated satisfactorily.

The access road slopes to the south towards the main yard area, which is located at a higher bench level cut into the hillside. The road is paved with tarmac, while the main operational area on site is concrete paved. The concrete is in generally good repair, however there are cracks and damaged areas along the northeast section of the access road. This damaged will be repaired as part of the proposed expansion of facility operations.

Storm water run-off from the site is collected in an ACCO type drain covered with a steel grid that runs along the northern site boundary. OCM inspected drain and the the northern boundary with the adjoining agricultural land and did not identify any evidence of the overflow of oil contaminated run-off from the drain.

In 2010, OCM noted a section of the drain in the western part of the site was partially covered with scrap metal over spilling from the site and that some debris has also fallen into the drain. These materials were since removed and the drain cleared.

The drain connects to a settlement tank located to the east of access road and south of the offices. The settlement tank overflows to an oil interceptor located the west of the site access road. The water from the interceptor discharges to the local authority storm water sewer.

The oil interceptor is routinely inspected and the discharge is monitored. OCM understand that the monitoring has established that the discharge complies with the emission limit values set in the Waste Permit. The oil accumulating in the interceptor is routinely skimmed of and used to lubricate the on-site shredder.

The settlement tank is de-sludged every six weeks and the contents are recycled over the scrap metal stock pile.

A steel frame and metal clad garage is located at the southern site boundary. Some oil staining was observed in 2010 and 2011 on the concrete floor, however the floor appears to be in good condition. There is a 1000 litre (l) polyethylene waste oil storage tank sitting in a steel bund, with two other smaller (205 l) waste oil tanks in an adjoining bund. Oil spill clean-up materials (Oil Dry) are stored inside the building and used to clean-up any minor spills that occur. The waste oil is collected as by ENVA a permitted waste contractor, and removed from the site for treatment.

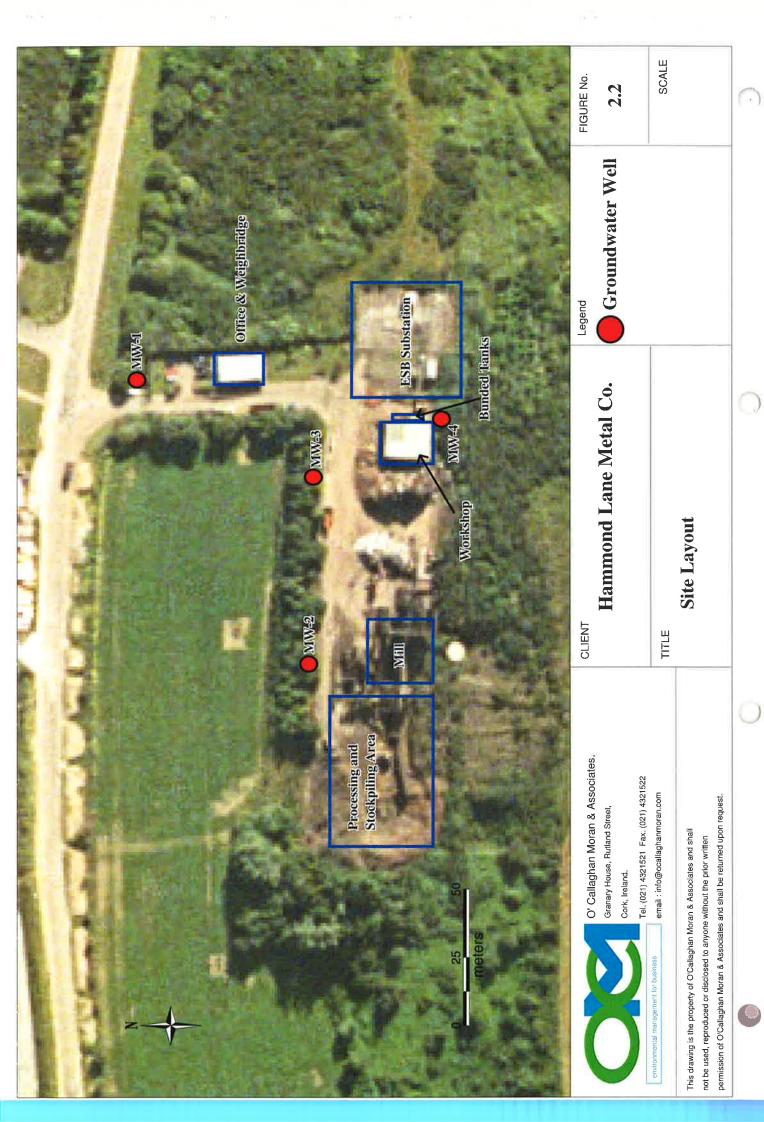
There are two fuel oil tanks in a concrete bund to the east of the garage. The first is a steel rectangular tank c1500l and the second is a polyethylene cylindrical tank of similar capacity. Both tanks appear to be in good repair. The bund is roofed and enclosed on three sides. At the time of the inspection in 2011, OCM noted the presence of a small amount of water in the bund, most likely from rainfall entering the open side to the enclosure. There were four 2051 steel waste oil drums located on a drum storage pallet in this area.

OCM did not observe any evidence of leaks at any of the bunds though some staining is present along the fill port of the steel rectangular tank.

Any oil spill within the garage that was not contained by site staff could escape onto the paved yard and ultimately be collected in the surface drain located in the drain along the northern boundary.

To the west of the garage is the incoming scrap metal stockpile, where materials are stored pending processing in the nearby shredder unit. The shredded metal is stockpiled to the west of the shredder.





# 2.4 Geology and Hydrogeology

Information on the local and regional geology and hydrogeology was derived from a desk study, which included Geological Survey of Ireland (GSI) geology databases; Teagasc Soil Maps for the region; in-house databases prepared by OCM and the well installation programme undertaken by OCM in July 2011. The latter is described in Section 3.

# 2.4.1 Soils and Subsoil

The subsoil distribution is shown on Figure 2.3. The site is underlain by 0.3 – 0.75m of gravelly fill partially comprising in-situ weathered and broken shale and sandstone bedrock. The Teagasc Soil Maps indicate the in-situ soils in the northern section of the site comprise Devonian Sandstone Till (**TDSs**). The southern section of the site is described as being underlain by non calcareous rock (**RckNCa**). This classification was confirmed by the 2011 investigation.

The depth of subsoils ranges from 2.7m in the north of the site to 0.3m near the southern boundary. Rock is exposed along the southern site boundary and also along the northwest site boundary, where the site is benched into the bedrock.

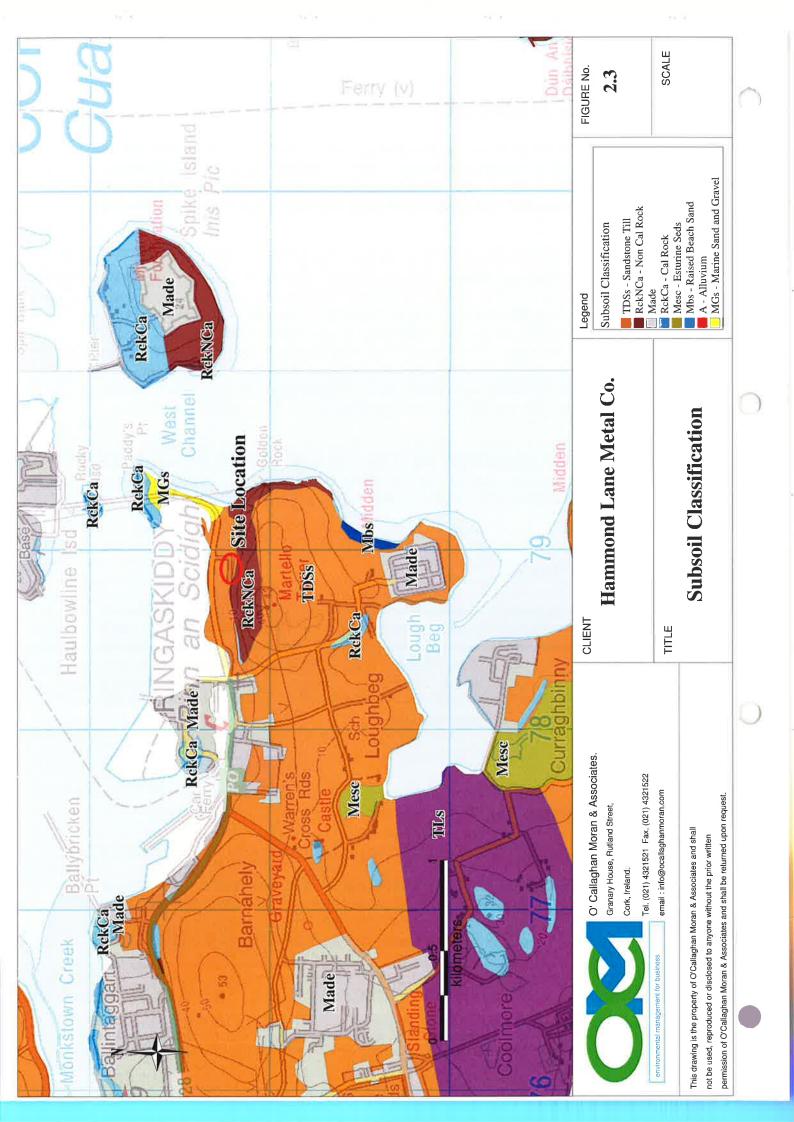
# 2.4.2 Bedrock

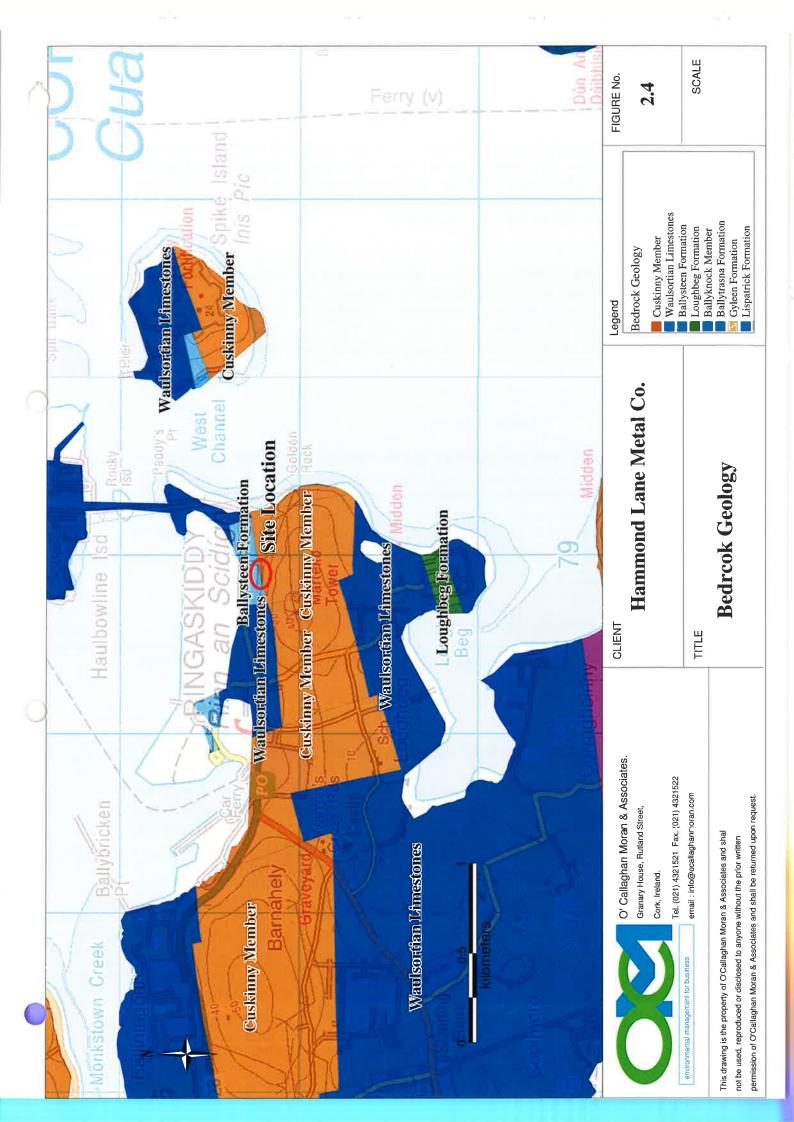
The bedrock geology is illustrated on Figure 2.4. The GSI maps indicated that northern section of the site is underlain by the Ballysteen formation, which comprises dark muddy limestone and shale, while the southern section of the site is underlain by the Cuskinny Member of the Kinsale formation, which comprises Flaser-bedded sandstone & mudstone. However, based on the bedrock encountered during the monitoring well installation, it appears that the entire site is underlain by the Cuskinny Member.

# 2.5 Hydrogeology

# 2.5.1 Aquifer Classification

The GSI has developed a classification system for aquifers based on the value of the resource and the hydrogeological characteristics. The bedrock aquifer beneath the site is characterised by the GSI as a Locally Important Aquifer (Ll), which is moderately productive in local zones. The aquifer classification is illustrated on Figure 2.5.





# 2.5.2 Aquifer Vulnerability

Vulnerability is defined by the GSI as the intrinsic geological and hydrogeological characteristics that determine the ease with which groundwater may be contaminated by human activities. Vulnerability categories range from Extreme (**E**) to High (**H**) to Moderate (**M**) to Low (**L**) and are dependant on the nature and thickness of subsoils above the water table.

The GSI vulnerability map indicates that the rating at the site ranges from extreme (**E**) in the northern part of the site to x-extreme (**X**) in the south. The site investigation confirmed this classification. The depth to bedrock in the northern section of the site at MW-1 was 2.7m, while in the south of the site (MW-4) the depth was 0.3m. The aquifer vulnerability is illustrated on Figure 2.6.

# 2.5.3 Aquifer Characteristics

The Cuskinny shale and sandstone and the Ballysteen Limestone formations are characterised by the GSI as a Locally Important Aquifers that are productive only in local zones (Ll) and are essentially poor aquifers. Typically groundwater yields from these aquifers are very low and in this area and they are not deemed to be a significant resource given that much of the land to the north of the site is reclaimed from the estuary and the likely impacts of saline intrusion.

# 2.5.4 Groundwater Flow Direction

Groundwater flow follows the local topography and is from the south to the north toward Cork Harbour.

# 2.5.5 Neighbouring Wells and Karst Features

The closest recorded wells are approximately 1km to the south in the Lough Beg Hovione facility. These wells are in a separate hydrologic catchment and range in depth from 7m to 21m. There is no information of what the wells are used for. There are no karst features recorded in the vicinity of the site.

# 2.5.6 Designated Areas

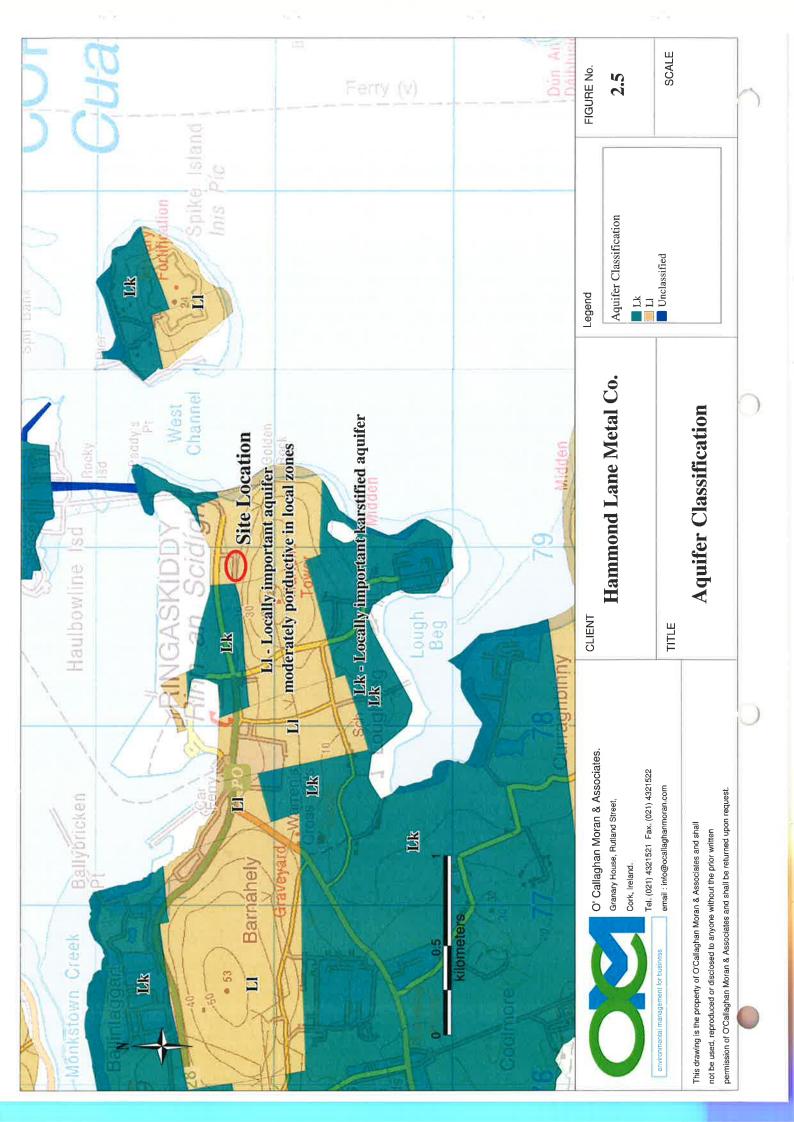
The nearest designated area is a Special Protected Area (SPA) located approximately 600m to the south in Lough Beg. Lough Beg is also a Proposed Natural Heritage Area (PNHA). The closest Special Area of Conservation (SAC) is approximately 6km to the north, in Great Island Channel.

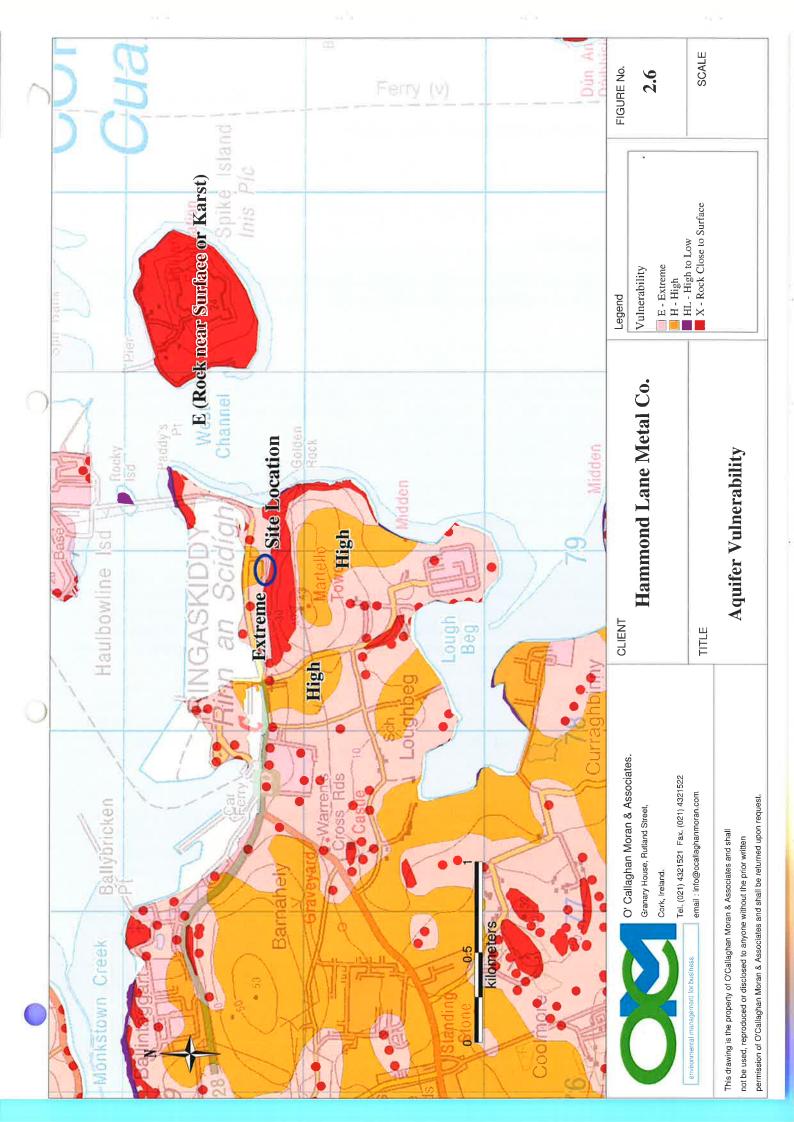
# 2.6 Conceptual Model

There are no surface water streams or springs in the vicinity of the site. The site is entirely covered with impermeable concrete paved yards and buildings, which means that rainfall recharge does not occur within the site boundaries.

Groundwater flow locally is expected to be from the high ground south of the site toward the harbour to the north. The bedrock aquifer comprises low permeability sandstone and shale. Groundwater flowing beneath the site is expected to discharge either to the low reclaimed land to the north and/or the estuary further to the north. The lands to the north of the public road and east of the Navy College have been reclaimed by progressively infilling with dredge spoil and construction demolition debris over many years.

Groundwater moving to the north from the poor aquifer beneath the site is likely to connect to the brackish groundwater near the estuary rather than moving deeper into the bedrock.





# 3 GROUNDWATER INVESTIGATION

# 3.1 Well Installation

Four monitoring wells (MW-1 to MW-4) were installed at the site between the 21<sup>st</sup> and 22<sup>nd</sup> July 2011. The well locations are shown on Figure 3.1. The well locations were selected by OCM.

MW-1 is located along the northern site boundary and immediately down hydraulic gradient of the oil interceptor and the septic tank. MW-2 and MW-3 are directly down hydraulic gradient of the main operational area, while MW-4 is along the southern site boundary and upgradient of the operational areas.

The wells were installed by Ground Investigations Ireland using a rotary percussion drill rig under the supervision of an OCM hydrogeologist. The borehole logs and well construction details are presented in Appendix 2.

The boreholes were drilled at 150mm diameter and cased to the top of bedrock. Groundwater strikes were not recorded in the subsoils. Bedrock was encountered in MW-1 at 2.7m, MW-2 at 2m, MW-3 at 1.5m and in MW-4 at 0.3m. Groundwater was encountered in MW-1 at 3.56m, in MW-2 at 9.5m, in MW-3 at 9.6m and in MW-4 at 10.2m. The total depths of the boreholes ranged from 10m in MW-1 to 14m in MW-3.

The monitoring wells were constructed using uPVC 50 mm diameter standpipe. A slotted section of standpipe was installed in the water bearing section of the bedrock in each borehole. The remainder of the well piping above the slotted section comprised solid 50mm uPVC pipe.

The annulus surrounding the slotted section in each well was back filled with washed pea gravel to act as a filter. Above the pea gravel the annulus surrounding the solid well pipe was back filled with bentonite to act as a seal to prevent the infiltration of surface water into the water bearing section of the well. Each borehole was finished with an upright steel well head set in concrete.

# 3.2 Groundwater Sampling

Groundwater samples were collected from the wells on the 27<sup>th</sup> July 2011 in accordance with OCM sampling protocols, a copy of which is included in Appendix 3. pH, temperature and electrical conductivity were measured in the field and the results are presented in Table 3.1.

Table 3.1	2.66	8.43	6.97	Not Known
Well ID	MW-1	MW-2	MW-3	MW-4
Sample Date	27 <sup>th</sup> July 2011			
Water Level (mBTOC)	3.14	8.45	7.15	8.87
Stick Up (m)	0.37	0.25	0.43	0.30
Water Level (mBGL)	2.77	8.2	6.72	8.57
pH (pH Units)	6.96	7.26	7.55	7.47
Electrical Conductivity (µS/cm)	789	969	897	776
Temperature (°C)	13.1	13.2	13.3	13.1
Water level - mOD	0.11	.23	0.25	????

After completion of groundwater level measurements, the wells were purged to remove the stagnant water from the well pipe and the surrounding gravel packs. Purging is necessary to ensure that the groundwater parameters measured are representative of the formation and not the stagnant water in the monitoring well or surrounding gravel filter.

The samples were placed in laboratory prepared containers and stored in coolers prior to shipment to Jones Environmental Forensics in the UK.

# 3.3 Groundwater Analysis

The samples were analysed for a range of parameters based on the use of the site as a metal processing facility. The parameters included heavy metals (lead, nickel, copper, zinc, arsenic, antimony, cadmium, chromium and mercury), phenols, Petrol Range Organics, Diesel Range Organics, Benzene, Toluene, Ethylbenzene, and Xylene (BTEX), polycyclic aromatic hydrocarbons (PAH). The methodologies used by the laboratory were ISO/CEN approved or equivalent and the method detection limits (MDL) were all below relevant limits and comparative guidance values.

The laboratory reports are included in Appendix 4 and the results are summarised in Tables 3.2 to 3.4. The table includes Interim Guideline Values (IGV) published by the EPA. The IGVs are not statutory, but were developed to assist in the assessment of impacts on groundwater quality in the context of the implementation of the EU Water Framework Directive. The guidelines are based on, but are more conservative than the Drinking Water quality standards. The table also includes for comparative purposes the Groundwater Threshold Values (GTV) set out in the European Communities Environmental Objectives (Groundwater) Regulations (S.I. 9 of 2010).

With the exception of low levels of nickel in MW-2 (at the MDL) and zinc in MW-2, 3 and 4, heavy metals were not detected. The levels of nickel and zinc are below the IGV. Petroleum hydrocarbons were not detected. With the exception of naphthalene, PAH were not detected. While naphthalene was detected in all the samples, the levels were significantly lower than the IGV. There is no GTV for naphthalene.

# 3.4 Data Interpretation

The analytical results confirm that site activities have not impacted on the quality of the groundwater beneath the site. The data also indicates that the septic tank is not having any impact on the shallow groundwater immediately down hydraulic gradient at MW-1.

Δ±Ω	415	0.0375	1.5	0.0075	Ų.	0.00375	0.015	0.01875		0.00075
ICV	10.4	0.03	0.03	0.01	13	0.005	0.02	0.01	0.1	0.001
MW-4	21/07/2011	<0.0015	<0.007	<0.0025	<0.002	<0.0015	<0.002	<0.005	0.015	<0.001
MW-3	21/07/2011	<0.0015	<0.007	<0.0025	<0.002	<0.0015	<0.002	<0.005	0.012	<0.001
MW-2	21/07/2011	<0.0015	<0.007	<0.0025	<0.002	<0.0015	0.002	<0.005	0.012	<0.001
MW-1	21/07/2011	<0.0015	<0.007	<0.0025	<0.002	<0.0015	<0.002	<0.005	<0.003	<0.001
Imite	Cilits	l/gm	mg/l	mg/l	mg/l	l/gm	mg/l	l/gm	mg/l	mg/l
Parameter		Chromium	Copper	Arsenic	Antimony	Cadmium	Nickel	Lead	Zinc	Mercury

 Table 3.3
 Hydrocarbons

í														
AJJ	415	•			16	į	40	Į,	N.E.	0.75	3	r	a.	Ę
$\Delta DI$	10.4	10		10	0.0005	10	10	10	30		10	10	10	10
MW-4	21/07/2011	<10		<10	<0.15	<100	<100	<100	<5	<5	<>	<5	\$	\$
MW-3	21/07/2011	<10		<10	<0.15	<100	<100	<100	<5	<5	<5	<5	<5	\$
MW-2	21/07/2011	<10		<10	<0.15	<100	<100	<100	<5	\$	<5	<5	<5	\$
MW-1	21/07/2011	<10		<10	<0.15	<100	<100	<100	<5	\$	<5	\$	<5	\$
Tinite	CIIIIS	l/gn		l/gn	mg/l	l/gn	l/gn	l/gn	ug/l	l/gn	l/gn	l/gn	l/gn	l/gn
Donomotor	rarameter	EPH (C8-C40)	Mineral Oil	(Calculation)	Total Phenols	GRO (C4-C8)	GRO (C8-C12)	GRO (C4-12)	MTBE	Benzene	Toluene	Ethyl benzene	m/p-Xylene	o-Xylene

Table 3.4 PAHs

AJO	410	2	0)		9		*	*	)	*	*	42	-	.9	0.0075		0.075	,	1.
ΛĐΙ	101	1						1				0.5	0.01	0.05		0.05			
MW-4	21/07/2011	0.020	<0.013	<0.013	<0.014	<0.011	<0.013	<0.012	<0.013	<0.015	<0.011	<0.018	<0.016	<0.011	<0.01	<0.011	<0.195	<0.01	<0.01
MW-3	21/07/2011	0.030	<0.013	<0.013	<0.014	<0.011	<0.013	<0.012	<0.013	<0.015	<0.011	<0.018	<0.016	<0.011	<0.01	<0.011	<0.195	<0.01	<0.01
MW-2	21/07/2011	0.020	<0.013	<0.013	<0.014	<0.011	<0.013	<0.012	<0.013	<0.015	<0.011	<0.018	<0.016	<0.011	<0.01	<0.011	<0.195	<0.01	<0.01
MW-1	21/07/2011	0.020	<0.013	<0.013	<0.014	<0.011	<0.013	<0.012	<0.013	<0.015	<0.011	<0.018	<0.016	<0.011	<0.01	<0.011	<0.195	<0.01	<0.01
Ilnite	Sillis	l/gn	l/gn	l/gn	l/gn	l/gn	l/gn	l/gn	l/gn	l/gn	l/gn	ug/l	ug/l	ug/l	l/gn	l/gn	l/gn	l/gn	ug/l
Doromotor	1 al ameter	Naphthalene	Acenaphthylene	Acenaphthene	Fluorene	Phenanthrene	Anthracene	Fluoranthene	Pyrene	Benz(a)anthracene	Chrysene	Benzo(bk)fluoranthene	Benzo(a)pyrene	Indeno(123cd)pyrene	Dibenzo(ah)anthracene	Benzo(ghi)perylene	PAH 16 Total	Benzo(b)fluoranthene	Benzo(k)fluoranthene

# 4 CONCLUSIONS AND RECOMMENDATIONS

# 4.1 Conclusions

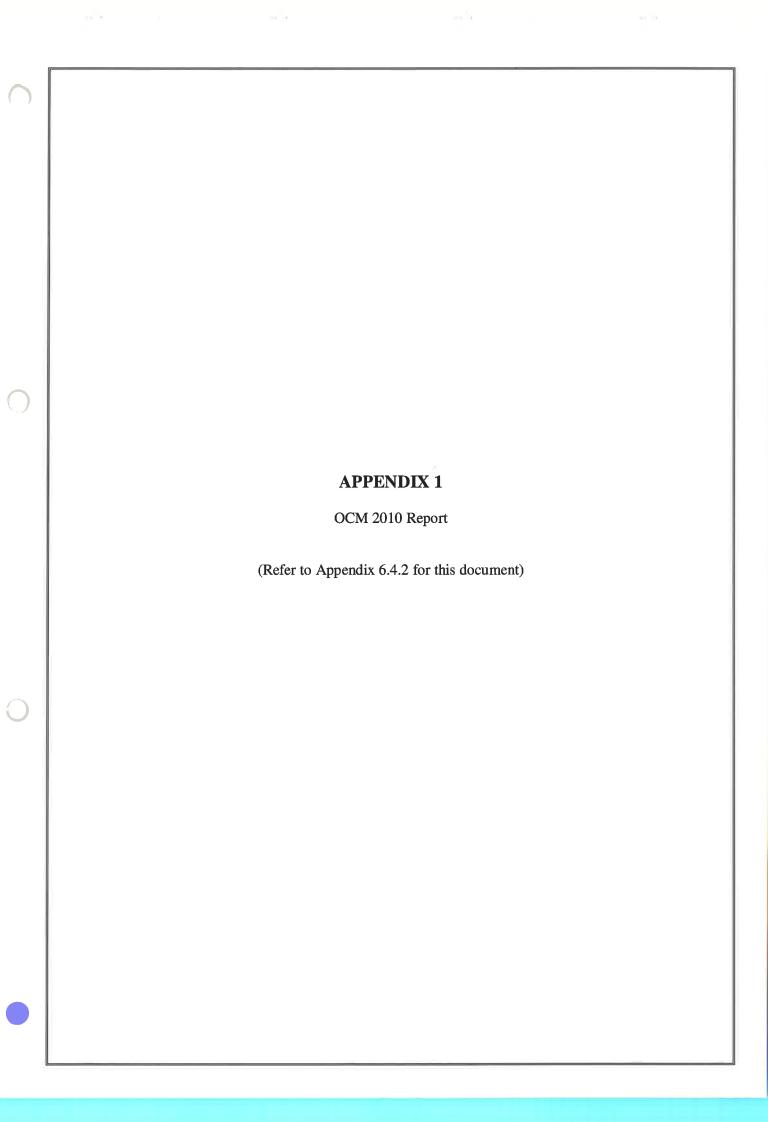
The site is entirely covered with impermeable hardstanding and buildings, which prevents infiltration of rainfall to the subsoils. The surface water drainage system, including the settlement tank and interceptor appear to be functioning properly.

The bedrock aquifer beneath the site is characterised by the GSI as a Locally Important Aquifer (Ll), which is moderately productive only in local zones. Based on the site investigation data, the aquifer vulnerability is extreme. There are no public or private groundwater wells used for potable supply within 2km of the site.

The groundwater quality monitoring has established that the groundwater beneath the site has not been impacted by either the historical or current use of the site. The findings support the conclusions of the 2010 Environmental Site Assessment that the site activities are not impacting on the subsoil or groundwater quality beneath or down gradient of the facility.

# 4.2 Recommendations

The groundwater monitoring should wells be clearly identified and protected by providing buffer areas around the wells where material cannot be placed on the ground. This is to prevent damage to the wells during day to day site activities.





# **APPENDIX 2**

Borehole Logs

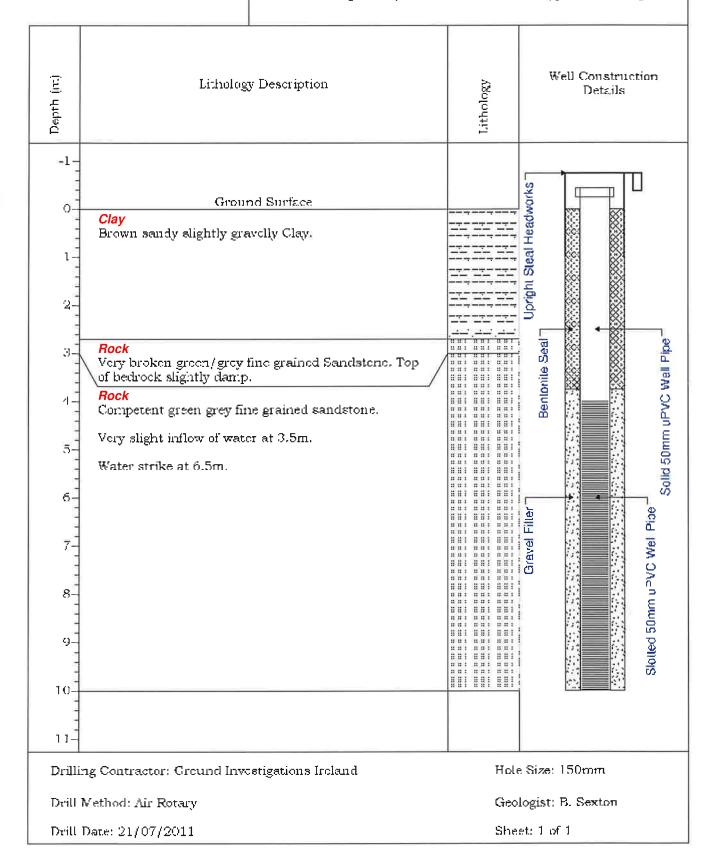


Project: 11 099 10

Borchold Depth: 10m

Client: Hammond Lane Metal Co. SWL (m): 2.66m

Location: Ringaskiddy, Cork





Project: 11 099 10

Borchold Depth: 12.5m

Client: Hammond Lane Metal Co. SWL (m): 8.43m.

Location: Ringaskiddy, Cork

Depth (nr.)	Lithology Description	Lithology	Well Construction Details
-1- 0- 1-	Ground Surface  Fill  Fill material comprising brown sandy slightly gravelly Clay.		Steal Headworks
3-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1	Rock Very broken green/grey fine grained Sandstone.  Rock Competent green grey fine grained sandstone.		Bentonite Seal
6- 7- 8-	<b>Rock</b> Very fine grained brown to red sandstone.		Gravel Filter
10- 11- 12-	Rock Grey fine grained sandstone.  Water inflow at 9.5m.		Skilled 50m
13- 14-	ng Court gatory Crowned Investigations Isoland	U_I	e Size: 150mm
	ng Contractor: Ground Invostigations Ireland Method: Air Rotary		logist: B. Sexton
	Date: 21/07/2011		et: 1 of 1

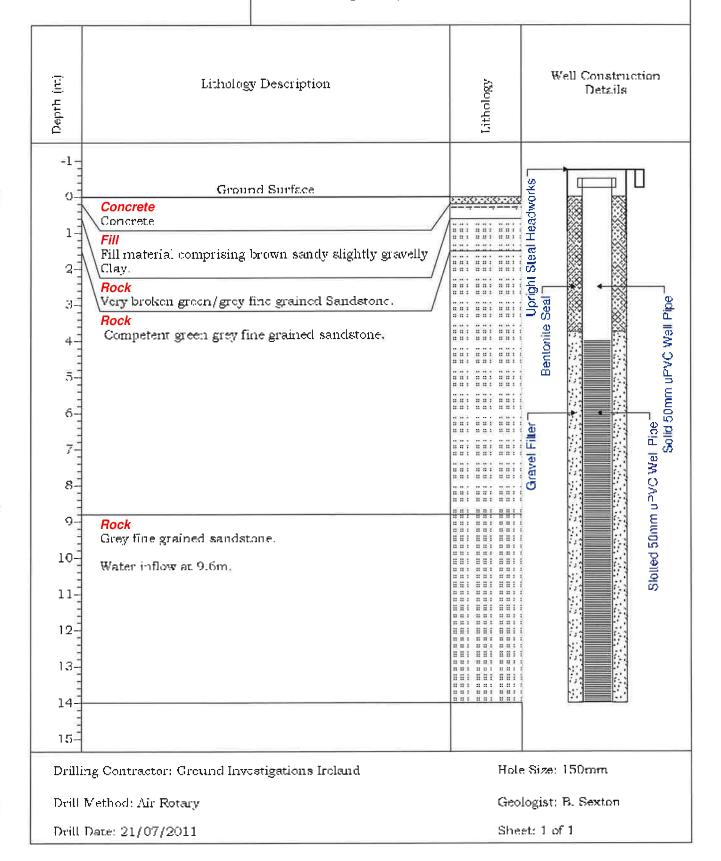


Project: 11 099 10

Borchold Dopth: 14m

Client: Hammond Lane Metal Co., SWL (m): 6,97m.

Location: Ringaskiddy, Cork



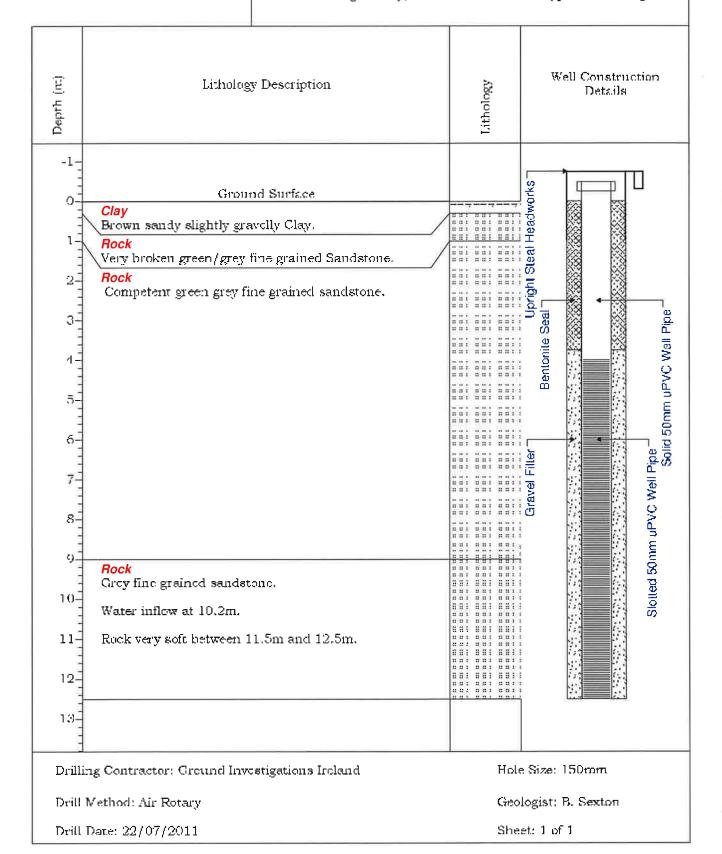


Project: 11 099 10

Borchold Depth: 12.5m

Client: Hammond Lane Metal Co. SWL (m):

Location: Ringaskiddy, Cork



# APPENDIX 3

**OCM Sampling Protocol** 



# STANDARD OPERATING PROCEDURE

## **GROUNDWATER SAMPLING**

The primary objective of groundwater sampling is to evaluate whether the potential contaminant sources at a site have impacted the quality of the groundwater in the underlying aquifer. The additional objective is to measure hydraulic gradient, or slope, of the water table in the shallow aquifer in an effort to evaluate the direction of groundwater flow.

The purpose of this procedure is to ensure that representative samples of groundwater are collected and documented using consistent methods to ensure sample integrity.

## 1.0 SAMPLING PROCEDURES

# 1.1 Well Operating and Purging Procedures

All groundwater sampling will be conducted after the installed and developed wells have been allowed to equilibrate for at least 2 to 3 days. A Field Data Sheet for Well Sampling will be completed for each well.

Groundwater sampling teams will use to following procedure for approaching, opening, purging and sampling all wells unless directed otherwise by the workplan.

- 1) Prior to placing any equipment into the well, decontaminate the sampling equipment according to standard decontamination protocol.
- 2) Approach the well with a working FID/PID, a well key, and a depth-to-water meter.
- 3) Unlock and open the well cap just enough to insert the probe of the PID/FID. Take and record a reading. A decision to upgrade PPE may be necessary based on the FID/PID readings in the breathing zone.
- 4) Where practical, the surface water column will be visually examined for the presence of hydrocarbons, if present or suspected, the thickness of the hydrocarbon layer will be measured using either an oil/water interface probe or transparent bailer prior to taking the depth-to-water measurement.
- 5) Insert the water level probe into the well and measure and record the static water level to the nearest 0.01 m with respect to the established survey point on top of the well casing.

C:\SOP\Gwater.Doc

- 6) Decontaminate the water level probe with DDI water (Do not rinse with any solvents unless product was encountered).
- 7) Calculate and record the minimum volume of water to be purged according to the following conversion factors: -

1 well volume	=	water column in metres x litres/linear metre
2 inch casing 4 inch casing	=	2.0 LPM 8.1 LPM
6 inch casing	=	18.2 LPM
8 inch casing	=	32.4 LPM

- 8) Purge the well of at least 3 casing volumes by pumping using a peristaltic pump with flow controller or bailing with a decontaminated submersible pump or PVC bailer equipped with a bottom filling check valve (if the purge volume is low, generally less than 100 litres, the sampling team might find it more efficient to purge with a bailer than a pump). Use a graduated bucket to track the amount of water removed from the well. The determination of purging and sampling will depend on parameters being analysed. Periodically determine the pH, temperature and specific conductance of the purged water. Continue purging until the well has been completely evacuated or until the pH and specific conductance measurements have stabilised for at least one well volume. Wells that become dewatered prior to producing three casing volumes will be sampled as soon as practical once they recover sufficiently.
- 9) Dispose of purge water collected in the graduated bucket by dumping onto the ground at a distance of 50 to 60 metres from the vicinity of the well. If the water is known or suspected to be significantly contaminated, it may be necessary to store the purge water in a secure container, such as a drum, pending proper disposal.
- 10) Be aware and record any unusual occurrence during purging such as cascading (a shallow water entry zone that trickles into the borehole).

# 1.2 Field Parameter Measurement

Measurements of field parameters of pH, temperature and electrical conductivity are collected and organic vapour screening is conducted while the well is purged. To facilitate the collection of basic field parameters, the field team needs to: -

- Purge three well volumes of water from the well and measure field parameters for each well volume removed.
- Collection of water samples should take place after stabilisation of the following parameters: -
  - Temperature +/- 1°C
  - pH (meter or paper) +/- 0.2 units Dissolved Oxygen +/-0.1 mg/l
  - Specific conductivity +/- 5%

- If the aforementioned parameters do not stabilise within three purge volumes, the well will be purged up to a maximum of six borehole volumes unless two consecutive sets of stabilised parameters are obtained.
- Note any observations in the field logbook.

# 1.3 <u>Collection of Water Samples</u>

All samples or chemical analysis will be placed in laboratory prepared bottles. The types of sample containers and preservative required for each type of analysis are described in the workplan. Where product layers are present a procedure and rational for the collection of such layers should be outlined in the site specific work plan. If required, preservatives will be placed in the sample containers prior to collecting the samples.

The following procedure will be used to sample a well: -

- 1) After the well has been purged and allowed to recover, sample the well using a properly decontaminated or dedicated disposable bailer. Gently lower the bailer into the water column. Allow the bailer to sink and fill with a minimum of surface disturbance.
- 2) Slowly raise the bailer out of the well. Do not allow the bailer line to contact the ground, either by coiling it on a clean plastic sheet or by looping it from arm to arm as the line is extracted from the well.
- 3) Samples will be collected for VOCs analysis immediately after purging is complete and before other samples are collected. Pour the samples slowly into the laboratory prepared 40 ml glass vial. Overfill each vial slightly to eliminate air bubbles, a convex meniscus should be present at the top of the vial. Ensure that the Teflon liner of the septum cap is facing inward and that no bubbles are entrapped. After capping securely, turn bottle upside-down, tap it against your other hand, and observe sample water for bubbles. If bubbles are observed, remove the cap, overfill the vial and reseal. Repeat this step for each vial until the samples with no bubbles are obtained.
- 4) Place a label on the container and enter the following information: -

Client/Site Name
Date Collected
Time Collected
Analysis
Preservative
Sample Identification Number

- 5) Record pertinent information in the field logbook and on the Field Data Sheet for Well Sampling. Complete chain-of-custody form.
- 6) Place custody seals on the container caps. As soon as possible, place sample containers in a cooler with bagged ice and maintain at 4°C until extraction. Surround the bottles with vermiculite.

- 7) Obtain the semi-volatile compound/pesticides/PCBs sample(s) by transferring the water to a laboratory prepared 1000 ml amber glass bottle with Teflon-lined cap. Fill the bottle to the bottom of the neck and follow steps 4, 5 and 6 above.
- 8) Dissolved metals (if necessary) requires the team to filter the sample water through a .45 micron filter. The water is collected in a 1 litre, unpreserved, plastic or glass bottle with HNO<sub>3</sub> preservative. Filtering must be done within 15 minutes of sample collection.
- 9) Obtain the total metals sample by directly transferring the water from the bailer into a laboratory prepared 1000 ml plastic or glass bottle with HNO<sub>3</sub> preservative.
- 10) Be sure the pH of the metals sampled is less than 2 by pouring off an aliquot in a clean jar and testing for pH using litmus paper. Dispose of this water and rinse the jar.
- 11) Collect and prepare Field QA/QC samples in accordance with separate SOP.
- 12) Be sure to record all data required on the Field Data Sheet or Well Sampling and appropriate entries into the field logbook.
- 13) Secure the well cap and replace the locking cover.
- 14) Decontaminate all sampling equipment according to procedure.
- 15) Decontaminate submersible pumps as follows: -

Scrub pump and cord in a tub of Liquinox/or similar and potable water Pump at least 80 litres of soapy water through pump Rinse with potable water Pump at least 80 litres of rinse water through the pump Rinse with D1 water before lowering pump into the next well.

END.

# **APPENDIX 4** Laboratory Results

 $C: \verb|\|11|099\_HammondLaneMetaiCo.|\|10\_Ringaskiddy, Cork|0991001.Doc$ 

August 2011 (SM/BS)



# Jones Environmental Laboratory

Unit 3 Deeside Point

Zone 3

Deeside Industrial Park

Deeside

CH5 2UA

Tel: +44 (0) 1244 833780 Fax: +44 (0) 1244 833781



No.4225

Attention:

Granary House

Rutland Street

Cork Ireland

O'Callaghan Moran & Associates

Date :

Barry Sexton
4th August, 2011

Your reference :

11-099-10

Our reference :

Test Report 11/5533 Batch 1

Location:

HAMMOND LANE RINGASKIDDY

Date samples received:

28th July, 2011

Status:

Final report

Issue:

- 1

Four samples were received for analysis on 28th July, 2011. Please find attached our Test Report which should be read with notes at the end of the report and should include all sections if reproduced. Interpretations and opinions are outside the scope of any accreditation, and all results relate only to samples supplied.

All analysis is carried out on as received samples and reported on a dry weight basis unless stated otherwise. Results are not surrogate corrected.

gelja

J W Farrell- Jones CChem FRSC Chartered Chemist

# Jones Environmental Laboratory

Client Name:

O'Callaghan Moran & Associates

11-099-10

Reference: Location:

HAMMOND LANE RINGASKIDDY

Contact:

Barry Sexton

Liquids/products: V=40ml vial, G=glass bottle, P=plastic bottle

Report : Liquid

100 1 1 AL	44/5500					11.11.00	7 7-6	N-OLL LIN	LINIO			
JE Job No.:	11/5533		_	_		H=H <sub>2</sub> SO <sub>4</sub> ,	Z=ZnAc, N=	NaOH, HN=	:HN0₃	i		
J E Sample No,	1-5	6-10	11-15	16-20								
Sample ID	MW-1	MW-2	E-WM	MW-4								
Depth									i i	Bloom an	e attached r	otoo for all
COC No / misc											iations and a	
					- 1							
Containers	VHPG	VHPG	VHPG	VHPG								
Sample Date	21/07/2011	21/07/2011	21/07/2011	21/07/2011								
Sample Type	Ground Water	Ground Water	Ground Water	Ground Water								
Batch Number	1	1	1	1								Method
Date of Receipt	29/07/2011	28/07/2011		28/07/2011	- 1					LOD	Units	No.
Dissolved Antimony *	<2	<2	<2	<2		-	_			<2	ug/l	TM30/PM14
Dissolved Artenic *	<2.5	<2.5	<2,5	<2,5						<2,5	ug/l	TM30/PM14
Dissolved Cadmium *	<0.5	<0.5	<0.5	<0,5						<0.5	ug/l	TM30/PM14
Total Dissolved Chromium *	<1.5	<1.5	<1.5	<1,5						<1.5	ug/l	TM30/PM14
Dissolved Copper#	<7	<7	<7	<7						<7	ug/l	TM30/PM14
Dissolved Lead "	<5	<5	<5	<5						<5	ug/l	TM30/PM14
Dissolved Mercury#	<1	<1	<1	<1						<1	ug/l	TM30/PM14
Dissolved Nickel*	<2	2	<2	<2						<2	ug/l	TM30/PM14
Dissolved Zinc "	<3	12	12	15						<3	ug/l	TM30/PM14
PAH MS												
Naphthalene "	0.020	0.020	0.030	0.020						<0.014	ug/l	TM4/PM30
Acenaphthylene *	<0.013	<0.013	<0.013	<0.013						<0.013	ug/l	TM4/PM30
Acenaphthene "	<0.013	<0.013	<0.013	<0,013						<0.013	ug/l	TM4/PM30
Fluorene *	<0.014	<0.014	<0.014	<0,014						<0.014	ug/l	TM4/PM30
Phenanthrene *	<0.011	<0,011	<0.011	<0,011						<0.011	ug/l	TM4/PM30
Anthracene *	<0.013	<0.013	<0.013	<0,013						<0.013	ug/l	TM4/PM30
Fluoranthene*	<0.012	<0.012	<0,012	<0,012						<0.012	ug/l	TM4/PM30
Pyrene "	<0.013	<0.013	<0.013	<0,013						<0.013	ug/l	TM4/PM30
Benz(a)anthracene "	<0.015	<0.015	<0.015	<0,015						<0.015	ug/l	TM4/PM30
Chrysene *	<0.011	<0.011	<0.011	<0.011						<0.011	ug/l	TM4/PM30
Benzo(bk)fluoranthene *	<0.018	<0.018	<0.018	<0.018						<0.018	ug/l	TM4/PM30
Benzo(a)pyrene *	<0.016	<0.016	<0.016	<0.016						<0.016	ug/l	TM4/PM30
Indeno(123cd)pyrene "	<0.011	<0.011	<0.011	<0.011						<0.011	ug/l	TM4/PM30
Dibenzo(ah)anthracene *	<0.01	<0.01	<0.01	<0,01						<0.01	ug/l	TM4/PM30
Benzo(ghi)perylene *	<0.011	<0,011	<0.011	<0.011						<0.011	ug/l	TM4/PM30
PAH 16 Total"	<0,195	<0.195	<0.195	<0.195						<0.195	ug/l	TM4/PM30
Benzo(b)fluoranthene	<0.01	<0.01	<0.01	<0.01						<0.01	ug/l	TM4/PM30
Benzo(k)fluoranthene	<0,01	<0.01	<0.01	<0.01						<0.01	ug/l	TM4/PM30
PAH Surrogale % Recovery	121	99	134	124						<0	%	TM4/PM30
EPH (C8-C40) *	<10	<10	<10	<10						<10	ug/l	TM5/PM30
Mineral Oil (Calculation)	<10	<10	<10	<10						<10	ug/l	TM5/PM30
Total Phenols HPLC	<0.15	<0.15	<0.15	<0.15						<0.15	mg/l	TM26/PM0
GRO (C4-C8) *	<100	<100	<100	<100						<100	ug/l	TM36/PM12
GRO (C8-C12) *	<100	<100	<100	<100						<100	ug/l	TM36/PM12
GRO (C4-12) "	<100	<100	<100	<100						<100	ug/l	TM36/PM12
мтве "	<5	<5	<5	<5						<5	ug/l	TM36/PM12
Benzene "	<5	<5	<5	<5						<5	ug/l	TM36/PM12
Toluene <sup>#</sup>	<5	<5	<5	<5						<5	ug/l	TM36/PM12
Ethyl benzene "	<5	<5	<5	<5						<5	ug/l	TM36/PM12
m/p-Xylene *	<5	<5	<5	<5						<5	ug/l	TM36/PM12
o-Xylene *	<5	<5	<5	<5						<5	ug/l	TM36/PM12

# NOTES TO ACCOMPANY ALL SCHEDULES AND REPORTS

#### SOILS

Please note we are only MCERTS accredited for sand, loam and clay and any other matrix is outside our scope of accreditation.

Where an MCERTS report has been requested, you will be notified within 48 hours of any samples that have been identified as being outside our MCERTS scope. As validation has been performed on clay, sand and loam, only samples that are predominantly these matrices, or combinations of them will be within our MCERTS scope. If samples are not one of a combination of the above matrices they will not be marked as MCERTS accredited.

It is assumed that you have taken representative samples on site and require analysis on a representative subsample. Stones will generally be included unless we are requested to remove them.

All samples will be discarded one month after the date of reporting, unless we are instructed to the contrary. If we are instructed to keep samples, a storage charge of £1 (1.5 Euros) per sample per month will be applied until we are asked to dispose of them.

If you have not already done so, please send us a purchase order if this is required by your company.

Where appropriate please make sure that our detection limits are suitable for your needs, if they are not, please notify us immediately.

All analysis is reported on a dry weight basis unless stated otherwise. Results are not surrogate corrected. Samples are dried at 35°C unless otherwise stated. Moisture content for CEN Leachate tests are dried at 105°C

#### WATERS

Please note we are not a Drinking Water Inspectorate (DWI) Approved Laboratory . It is important that detection limits are carefully considered when requesting water analysis.

UKAS accreditation applies to surface water and groundwater and one other matrix which is analysis specific, any other liquids are outside our scope of accreditation

As surface waters require different sample preparation to groundwaters the laboratory must be informed of the water type when submitting samples. All samples are treated as groundwaters and analysis performed on settled samples unless we are instructed otherwise.

## **DEVIATING SAMPLES**

Samples must be received in a condition appropriate to the requested analyses. All samples should be submitted to the laboratory in suitable containers with sufficient ice packs to sustain an appropriate temperature for the requested analysis. If this is not the case you will be informed and any analysis that may be compromised highlighted on your schedule/ report by the use of a symbol.

The use of any of the following symbols indicates that the sample was deviating and the test result may be unreliable:

\$	Sample temperature on receipt considered inappropriate for analysis requested.	
^	Samples exceeding recommended holding times.	
&	Samples received in inappropriate containers (e,g. volatile samples not submitted in VOC jars/vials).	
~	No sampling date given, unable to confirm if samples are with acceptable holding times.	

# **SURROGATES**

Surrogate compounds are added during the preparation process to monitor recovery of analytes. However low recovery in soils is often due to peat, clay or other organic rich matrices. For waters this can be due to oxidants, surfactants, organic rich sediments or remediation fluids. Acceptable limits for most organic methods are 70 - 130% and for VOCs are 50 - 150%. When surrogate recoveries are outside the performance criteria but the associated AQC passes this is assumed to be due to matrix effect. Results are not surrogate corrected.

#### **AQCs**

Where AQC's fall outside UKAS/MCERTS criteria analysis is repeated if possible.

#### NOTE

Data is only accredited when all the requirements of our Quality System have been met. In certain circumstances where the requirements have not been met, the laboratory may issue the data in its final report if it believes that the validity of the data has not been conpromised but will remove the accreditation. Please do not hesitate to contact the laboratory if further details are required of the circumstances which have led to the removal of accreditation.

# ABBREVIATIONS and ACRONYMS USED

#	UKAS accredited,
М	MCERTS accredited.
NAD	No Asbestos Detected.
NFD	No Fibres Detected
ND	None Detected (usually refers to VOC and/SVOC TICs).
SS	Calibrated against a single substance.
3.€	Analysis subcontracted to a Jones Environmental approved laboratory.
W	Results expressed on as received basis.
+	Accreditation has been removed from this result, if appropriate, see 'Note' on previous page.
++	Result outside calibration range, results should be considered as indicative only and are not accredited.
sv	Surrogate recovery outside performance criteria. This may be due to a matrix effect.
DR	Dilution required.

# Jones Environmental Laboratory

# Method Code Appendix

_		appropriate)			only)	Analysis done on As Received (AR) or Air Dried (AD)	expressed on Dry/Wet basis
TM4 1	6 PAH by GC-MS, modified USEPA 8270	PM30	Magnetic stirrer extraction				
	6 PAH by GC-MS, modified USEPA 8270	PM30	Magnetic stirrer extraction	Yes			
TM5 E	PH by GC-FID, modified USEPA 8015	PM30	Magnetic stirrer extraction			AR	
TM5 E	PH by GC-FID, modified USEPA 8015	PM30	Magnetic stirrer extraction	Yes		AR	
TM26 P	Phenols by HPLC	PM0	No Preparation				
ТМ30	vietals by ICP-OES	PM14	Metals by ICP (Waters)	Yes			
TM36 G	GRO by Headspace GC-FID	PM12	GRO GC-FID	Yes			
					-		