

# Middle River, Maryland

July 2004

Prepared for:

Lockheed Martin Corporation

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# Section 1

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Qn\_behalf\_of Lockheed\_Martin Corporation (I MC) Tetra Tech has prepared this Technical

entitled Supplemental Guidance on Performing Risk Assessments in Remedial Investigation, Feasibility Studies (RI/FS) Conducted by Potential Responsible Parties (PRPs), July 2, 1991. This Technical Memorandum is being submitted in lieu of a risk assessment work plan.

### 1.1 GUIDANCE DOCUMENTS

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	The defense of the second state of the second
	Risk Assessment Guidance for Superfund, Part A, Vol. 1: Human Health Evaluation Manual, USEPA, 1989;
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	Risk Assessment Guidance for Superfund (RAGS).Vol. I - Human Health Evaluation Manual (USEPA, 1989).

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### 1.2 PURPOSE AND OBJECTIVES OF THE BHRA

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could come into contact with the chemicals of potential concern (COPCs) in the soil and groundwater at the Site. Another objective is to propose the cleanup goals for each chemical of concern (COC) in the soil and groundwater in order to reduce or mitigate any unacceptable levels of health risks.

### 1.3 SCOPE OF THE BASELINE HEALTH RISK ASSESSMENT

The data from soil and groundwater investigations will be collectively used to perform the site-specific human health risk assessment. This includes, (a) the soil data collected during the subsurface investigations conducted from 2000 through 2002, and (b) the groundwater data collected within the

The BHRA will include the following elements:

- Identification of Chemicals of Potential Concern (COPCs),
- Exposure Assessment,
- Toxicity Assessment and

amonte with the ultimate goal of obtaining approval from the MDE on the proposed approved.

#### 1.4 ORGANIZATION OF THE TECHNICAL MEMORANDUM

Section 2.0 presents the background information on the Site. The physical, geological,

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This section also discusses the screening evaluation that will be conducted in order to focus the BHRA on the COPCs that fail the screening evaluation.

Section 4.0 presents the conceptual site model (CSM) that provides the framework of the exposure assessment. Section 4 also describes the different factors that will be considered in evaluating how, and to what extent, potential exposures could occur. These

incorporated in estimating the chemical dose to each exposed individual.

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FIGURE 2-1 E LOCATION MAP	MIDDLE RIVER
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### 2.4.2 Lockheed Martin Corporation's Investigations

In March 1999, Lockheed Martin collected groundwater monitoring well data to obtain updated

cadmium) were present above the Maximum Contaminant Levels (MCLs) for drinking water..

in groundwater. Each of the four AOCs listed in Section 2.4.1 was investigated through a

and petroleum hydrocarbon areas.

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lateral extent of chemical occurrence in the near-surface groundwater at the Site. A limited number of deep wells were installed to evaluate the vertical extent of VOCs and metals in the groundwater. The results of the lateral investigations indicated that the potential source areas are the Taxiway Tango median area, the drum area, and the petroleum hydrocarbon area – see Section 2.4.1. The primary contaminants were identified to be TCE, vinyl chloride, and cis-1, 2-DCE. The groundwater modeling suggested that VOCs in groundwater appear to be migrating from west to east toward Frog Mortar Creek (Tetra Tech, 2002). The general extent of VOCs in groundwater was delineated to the north and south, but not to the east and west.

Additional multi lower monthance wells was a set a second state of the second state of

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	present at the site continuumy to three primary groundwater pithtes. Dased on the concentration	
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	plumes, in particular with respect to plume migration toward Frog Mortar Creek. The distribution	
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### Section 3

### DATA EVALUATION

The data from the previous investigations will be reviewent to ensure that the number and quality of

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	งละและเบก process พระ 26 แหม่นนอน #1 และ อภาณ ายุมาน.	
	Sizes the investigation areas or ADCs are based on the supported sources of chemical relates. the	
	specific receptors. Therefore, the data set for the BHRA may vary from the data set used for site characterization. The notential human health risks will be evaluated by assuming two exposure	· .
ī	A $b$	
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	consist of buried drums and debris (MFS 1994) Since it is unlikely that there would be surface	
-	hatau ana faat haa ta a aandarwaa dauth af dit faat kan	

Any popstituent that is detected at least once will be included in the screenion\_risk evaluation. A

the cleanup standard for an industrial site (EPA Region III), then the chemical will be identified as a COPC that will be evaluated further in the risk assessment.

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### 3.2 SCREENING EVALUATION

In the screening evaluation, the highest concentration\* of each detected chemical will be compared

average site concentration, is also consistent with the conservative nature of the screening evaluation.

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### Section 4 EXPOSURE ASSESSMENT

The evaneurs accessment identifies and describes notantially accessed human recentors. develops

concentration).

### 4.1 CONCEPTUAL SITE MODEL

Figure 4-1, presents the concentual site model (CSM) that will be used as the framework for

.... --... we arreate them open expected plantage are complete or meetiples, and

incomplete pathways are excluded from the BHRA.

### 4.1.1 Potential Exposure Pathways

An exposure nathway is the mechanism by which a human recentor is exposed to chemicals from a

If any one of the four elements is missing, the exposure pathway is considered incomplete. Only

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current potential exposure pathways are those that exist as a result of the current extent of

existing ponds will not be evaluated in the BHRA. This is not the case for Frog Mortar Creek. Since

### 4.1.2 Current and Future Receptors

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The current and future land use are anticipated to be similar, thus, the current and future, receptors are the on-Site workers. Since a potential Site visitor would have more limited exposures than the on-Site worker, the visitor scenario will be evaluated if the potential risk to the

<u> </u>		POTENTIAL RECEPTORS	
- <u> </u>	Surface Soil Increasing Ruroff		
	(perticulates)		
-	Operations         Dispersion         Soil         Inhalation           Subsurface Soil         (narticulates)         (narticulates)	$\bigcirc \bigcirc \bigcirc$	
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Table 4-1 Summary of Exposure Parameters Martin State Airport

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	Ingestion Rate (mg/day)	50	(1)	480	·
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				<u>, 11</u>	
I ~1.					
	Target Risk	Specific	(4)		
	Target Hazard Index	1.00E-06	(1)	1.00E-06	
	Notes:				
	(1) EPA 1989 Risk Assessment Guidance for Su	nerfund - Volum	a 1 Human Usalih Evalu	istica Manual	
	4.2.3 Ingestion Algorithm				
	_				
-	The accustion for a destation the second second		······································		
	IngestionDose = <u>Cs x IR x EF</u>	F x ED x CF			
	BW x				
	277 X	<i>~</i> 11 <i>7</i>			
	where:				
	Induction Doop		:	/	
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k					
	EF	=	exposure freque	ency (days/year)	
	ED	÷	exposure durati		
	BW	Ξ			
			body weight (kg		
	AT	=	averaging time		
	CF	=	unit conversion	factor (10 <sup>-6</sup> kg/mg)	

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### 4.2.4 Inhalation Algorithm

The equation for calculating intake through inhalation of dust from Site soil is as follows:

Inhalation Dose =	EPCax InhR x ET x EF x ED
	BWxAT

where:

Inhalation Dose	Ж	inhalation dose (mg/kg-day)
InhR	=	inhalation rate (m³/day or m³/hr)
EPCa	=	exposure point concentration in air (mg/m <sup>3</sup> )
		particulates
	=	concentration in soil x (1/PEF)
where	e:	
		PEF = particulate emission factor (m <sup>3</sup> /kg),
ED	=	exonsure duration (vears)
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		· · · · · · · · · · · · · · · · · · ·
BW	=	body weight (kg)
AT	=	averaging time (days)

### 4.2.5 Dermal Algorithms

The equation for calculating intake through dermal contact with soil is as follows:

 $Dermal Dose = \frac{Cs \ x \ SSA \ x \ ABS \ x \ AF \ x \ EF \ x \ ED \ x \ CF}{BW \ x \ AT}$ 

where:

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Dermal Dose	=	dermal dose (mg/kg-day)
Cs	Ħ	exposure point concentration in soil (mg/kg)
AF	≈	soil to skin adherence factor (mg/cm <sup>2</sup> ),
SSA	=	exposed skin surface area (cm²/day)
ABS	=	absorption fraction of chemical from soil
EF	=	exposure frequency (days/year)
<u>r</u> n	-	onpoonio animiori granoj
CF	=	unit conversion (10 <sup>-6</sup> kg/mg)
BW	Ŧ	body weight (kg)
AT	=	averaging time (days)

# Section 5 TOXICITY ASSESSMENT

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report.

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## Section 6 **RISK CHARACTERIZATION**

	This section describes how the calculated exposure doses will be integrated with the toxisity arteria
b	The theoretical excess lifetime cancer risk is an estimate of the increased risk of an individual
	Excess Cancer Risk = Exposure Dose x Slope Factor
	Lifetime daily intakes, using an averaging time of 70 years, effectively prorate the total cumulative dose over a lifetime. This approach is based on the assumption that a high dose of carcinogens
	received over a short period of time_at_any_ane_is_equivalent to a correspondingly low dose received
1	
<b></b>	
۱ <del>۰,</del>	The potential for adverse effects on human health other than cancer will be evaluated by comparing
	available. Utherwise the reference dose based on chronic exposures will be applied. This
¥ \	
	Where: HQ = Hazard Quotient (unitless)
-	
-	
	A HQ of slightly greater than 1 is not necessarily an indication that adverse effects will occur. The
	bezard index in the sum of the LINe for each of the channels count in the second
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than 1.0, it is believed that no threshold health effects will occur. An HI of slightly greater than 1,

traviouar is not nanaceprily on indication that hadth offerts will assure . Summing WOs parace all

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The estimated cancer and noncancer risks will be presented in the BHRA report. The uncertainties associated with each component of the risk assessment will be discussed.

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