



**US Army Corps
of Engineers®**
Engineer Research and
Development Center

ERDC
INNOVATIVE SOLUTIONS
for a safer, better world

Environmental Science and Technology Certification Program

Cost and Performance Report of Incremental Sampling Methodology for Soil Containing Metallic Residues

Project ER-0918

Jay L. Clausen, Thomas Georgian, and Anthony Bednar

September 2013



The US Army Engineer Research and Development Center (ERDC) solves the nation's toughest engineering and environmental challenges. ERDC develops innovative solutions in civil and military engineering, geospatial sciences, water resources, and environmental sciences for the Army, the Department of Defense, civilian agencies, and our nation's public good. Find out more at www.erdcl.usace.army.mil.

To search for other technical reports published by ERDC, visit the ERDC online library at <http://acwc.sdp.sirsi.net/client/default>.

Cost and Performance Report of Incremental Sampling Methodology for Soil Containing Metallic Residues

Project ER-0918

Jay L. Clausen

*Cold Regions Research and Engineering Laboratory (CRREL)
US Army Engineer Research and Development Center
72 Lyme Road
Hanover, NH 03755-1290*

Thomas Georgian

*Environmental and Munitions Center of Expertise (EM CX)
US Army Corps of Engineers
1616 Capitol Avenue
Omaha, NE 68102*

Anthony Bednar

*Environmental Laboratory (EL)
US Army Engineer Research and Development Center
Waterways Experiment Station
3909 Halls Ferry Road
Vicksburg, MS 39180-6199*

Final Report

Approved for public release; distribution is unlimited.

Prepared for Environmental Security Technology Certification Program (ESTCP)
Arlington, VA 22203

Under Project ER-0918

Abstract

Objectives of this project were to demonstrate improved data quality for metal constituents in surface soils on military training ranges and to develop a methodology that would result in the same or lower cost. The demonstration was conducted at two inactive small-arms ranges at Fort Eustis, VA, and Kimama Training Site (TS), ID, and at one active small-arms range at Fort Wainwright, AK. The samples included 63 Incremental Sampling Methodology (ISM) and 50 conventional grab from Fort Wainwright, 18 ISM and 30 grab from Kimama TS, and 27 ISM and 33 grab from Fort Eustis. The variability in metal concentrations as measured with replicate samples and evaluated using percent relative standard deviation (RSD) were less than 10% for all metals using ISM. In contrast, RSDs were often greater than 50% for conventional replicate grab samples. Calculated mean ISM metal concentrations were statistically greater than the mean for conventional grab samples.

DISCLAIMER: The contents of this report are not to be used for advertising, publication, or promotional purposes. Citation of trade names does not constitute an official endorsement or approval of the use of such commercial products. All product names and trademarks cited are the property of their respective owners. The findings of this report are not to be construed as an official Department of the Army position unless so designated by other authorized documents.

DESTROY THIS REPORT WHEN NO LONGER NEEDED. DO NOT RETURN IT TO THE ORIGINATOR.

Contents

Abstract	iv
Illustrations	vii
Preface	ix
Nomenclature	xi
Unit Conversion Factors	xiii
Executive Summary	xiv
1 Introduction	1
1.1 Background	2
1.2 Objective of the demonstration	4
1.3 Regulatory drivers	5
2 Technology	7
2.1 Technology description	7
2.2 Advantages and limitations of the technology	11
3 Performance Objectives	15
4 Site Description	18
4.1 Site location and history	18
4.1.1 Kimama Training Site	18
4.1.2 Fort Eustis	19
4.1.3 Fort Wainwright	20
4.2 Site geology and hydrogeology	21
4.2.1 Kimama Training Site	21
4.2.2 Fort Eustis	22
4.2.3 Fort Wainwright	22
4.3 Contaminant distribution	22
4.3.1 Kimama Training Site	22
4.3.2 Fort Eustis	23
4.3.3 Fort Wainwright	23
5 Test Design	24
5.1 Conceptual experimental design	24
5.2 Baseline characterization	29
5.3 Treatability or laboratory study results	30
5.4 Field testing	30
5.5 Sampling methods	33
5.5.1 Grab samples	33

5.5.2 Incremental sampling methodology samples	33
5.6 Sampling results	35
5.6.1 Kimama Training Site	35
5.6.2 Fort Eustis.....	36
5.6.3 Fort Wainwright.....	37
6 Performance Assessment	39
6.1 Sample reproducibility with Incremental Sample Methodology	39
6.2 Bias evaluation.....	42
6.3 Performance comparison	43
7 Cost Assessment.....	46
7.1 Cost model.....	46
7.2 Cost drivers.....	49
7.3 Cost analysis	50
8 Implementation Issues	53
References	55
Appendix A: Points of Contact.....	61
Report Documentation Page	

Illustrations

Figures

1	Example of multi-increment sampling using a systematic random sampling design for collecting two separate 100-increment samples	4
2	Location of the Kimama Training Site, Idaho, and location of Training Area 3.....	19
3	Map showing location of Fort Eustis, VA, and the 1000-in. Rifle Range.....	20
4	Map of Fort Wainwright, AK, and the Range 16 Record Range.....	21
5	Generic example of a typical small-arms firing range	24
6	The northernmost small-arms range berm face located in Training Area 3 of Kimama TS.....	25
7	The 1000-in. small-arms range berm face at Fort Eustis	25
8	The small-arms firing Range 16 Record berms at Fort Wainwright.....	26
9	Location of berms sampled using ISM and grab techniques at the Range 16 Record Range at Fort Wainwright.....	27
10	Grab sample grid layout for Fort Wainwright	27
11	Grab sample grid layout for Kimama TS berm face.....	28
12	Grab sample grid layout for Fort Eustis berm face	28
13	Arial view of grab sample grid locations and DU boundaries for the 1000-in. Range berm face at Fort Eustis.....	28
14	Arial view of grab sample grid locations and DU boundaries for the NW berm face at Kimama TS	29
15	Grab surface soil results for lead, copper, antimony, and zinc (mg/kg) from the Kimama Training Site small-arms range berm face with Incremental Sampling Methodology comparisons.....	35
16	Analysis of variance for lead at Fort Wainwright	40
17	Analysis of variance for zinc at Fort Wainwright	41

Tables

1	Incremental sampling methodology for metallic residues.....	3
2	Salient differences between Method 3050B and proposed Method 3050C.....	9
3	Chronological summary of multi-increment sampling	10
4	Comparison of the advantages and disadvantages of the Incremental Sample Methodology.....	12
5	Performance objectives for ER-0918.....	16
6	Gantt chart for field demonstration activities.....	31
7	Comparison of Grab versus ISM for this demonstration.....	34

8	Comparison of relative standard deviations for the analytes of interest (copper, lead, antimony, zinc) for ISM and conventional grab samples at three demonstration sites	42
9	Comparison of labor costs between ISM and conventional grab sampling on a per sample and total cost basis based on demonstrations at Kimama Training Site, Fort Eustis, and Fort Wainwright.....	45
10	Comparison of labor hours or costs by cost element between ISM and conventional grab sampling on a per sample and total cost basis based on demonstrations at Kimama Training Site, Fort Eustis, and Fort Wainwright	47
11	Comparison of costs for ISM and conventional grab sampling on a per sample and total cost basis based on demonstrations at Kimama Training Site, Fort Eustis, and Fort Wainwright	48

Preface

Funding for this demonstration was provided by the Environmental Security Technology Certification Program (ESTCP) under Project ER-0918: *Demonstration of the Attributes of Multi-Increment Sampling and Proper Sample Processing Protocols for the Characterization of Metals on DoD Facilities* with Dr. Andrea Leeson as Program Manager for Environmental Restoration.

This report was prepared by Jay Clausen (Biogeochemical Sciences Branch, Dr. Terrence Sobecki, Chief) at the US Army Corps of Engineers (USACE) Engineer Research and Development Center (ERDC), Cold Regions Research and Engineering Laboratory (CRREL) in Hanover, New Hampshire (ERDC-CRREL); Anthony Bednar at the ERDC Environmental Laboratory (EL); and Thomas Georgian, USACE Environmental and Munitions Center of Expertise (EM CX). Other team members included Nancy Perron, Gordon Gooch, Nathan Mulherin, Arthur Gelvin, Stephanie Saari, and Marc Beede with ERDC-CRREL; Andrew Bray, Patricia Tuminello, William Jones, and Shawna Tazik with ERDC-EL; Larry Penfold with Test America; and Diane Anderson with APPL Laboratories.

COL Jeffrey R. Eckstein was the Commander of ERDC, and Dr. Jeffery P. Holland was the Director.

Special thanks go to Dr. Andrea Leeson, Dr. Jeffrey Marquess, and all the ESTCP panel members and technical advisors for the advice and technical review that they have provided.

Thanks also go to Bonnie Packer, Robert Halla, and Anne Woods (all with the National Guard Bureau) for arranging access to the Kimama Training Site and to Travis McCuin at the USACE Baltimore District for arranging access to Fort Eustis. Additionally, we appreciate the site access coordination provided by Arba Williamson and Amber Michel (Fort Eustis) and COL Kyle Carpenter and MAJ Andrew White (Idaho National Guard for the Kimama Training Site).

This report is dedicated to Alan Hewitt, our colleague who conceived of this project. He is admired for his enthusiasm and energy for scientific research, and his contributions will endure.

Nomenclature

BRAC	Base Realignment and Closure
CEC	Cation Exchange Capacity
CMIST	CRREL Multi-Increment Sampling Tool
CRREL	Cold Regions Research and Engineering Laboratory
CSM	Conceptual Site Model
DoD	United States Department of Defense
DQO	Data Quality Objective
DU	Decision Unit
EL	Environmental Laboratory
EM CX	USACE Environmental and Munitions Center of Expertise
EOD	Explosive Ordnance Disposal
ERA	Environmental Research Associates
ERDC	Engineer Research Development Center
ESTCP	Environmental Science Technology Certification Program
FS	Feasibility Study
FUDS	Formerly Used Defense Sites
IQR	Interquartile Range
ISM	Incremental Sample Methodology
ITRC	Interstate Technology Regulatory Council
LCS	Laboratory Control Sample
MB	Method Blank

MMRP	Military Munitions Response Program
NG	Nitroglycerin
ORAP	Operational Range Assessment
QA/QC	Quality Assurance/Quality Control
RI	Remedial Investigation
RSD	[Percent] Relative Standard Deviation
SI	Site Investigation
SU	Sampling Unit
TAL	Target Analyte List
TOC	Total Organic Carbon
TS	Training Site
USACE	United States Army Corps of Engineers
USEPA	United States Environmental Protection Agency
UXO	Unexploded Ordnance

Unit Conversion Factors

Multiply	By	To Obtain
feet	0.3048	meters
inches	0.0254	meters

Executive Summary

Background

Over the last decade, the environmental community has recognized that releases of energetic constituents into the environment as a result of military training occurs in an extremely heterogeneous pattern. Conventional soil sampling and sample preparation methodologies are inadequate to address the level of contaminant heterogeneity observed. Recently, the environmental and regulatory communities have raised questions regarding whether the issues observed for the deposition of energetic constituents also substantively apply to other constituents, such as metals, semi-volatile organic compounds, and polychlorinated biphenyls.

Objectives of the demonstration

This report was completed as a partial fulfillment of the obligations for Environmental Science Technology Certification Program Demonstration Project ER-0918. The primary objective of this project was to develop a sampling and laboratory analysis method for metals in surface soils on military training ranges that demonstrably produces higher quality data at lower costs than conventional sampling and analysis methods.

Demonstration results

Incremental Sampling Methodology (ISM), including both field and laboratory protocol development, was conducted at an active small-arms range at Camp Ethan Allen, VT, as reported in *Evaluation of Sampling and Sample Preparation Modifications for Soil Containing Metal Residues* (Clausen et al. 2012b). In addition, a demonstration was conducted at three additional small-arms ranges as reported in *Demonstration of Incremental Sampling Methodology for Soil Containing Metallic Residues* (Clausen et al. 2013a). The inactive ranges assessed included the 1000-in. Rifle Range at Fort Eustis, VA, and the Northern Area 3 of the Kimama Training Site (TS), ID. Both of these ranges are Military Munitions Response Program (MMRP) sites. A demonstration was also conducted at the active Range 16 Record Range located within the Small Arms Complex at Fort Wainwright, AK.

The demonstration followed initial protocol development at Camp Ethan Allen and included collecting 63 ISM surface soil samples and 50 conventional grab/discrete samples at Fort Wainwright; 18 ISM and 30 grab samples from Kimama TS; and 27 ISM and 33 grab samples at Fort Eustis. ISM involves changes both to the field sampling approach and to laboratory sample preparation procedures. Each incremental sample was prepared (in the field) by combining a set of multiple increments (of roughly equal soil mass) that were collected using systematic random sampling over the same Sampling Unit (SU).

The performance criteria used to determine whether ISM provided technically defensible data were (1) reproducible results for surface soil samples containing metal particles, (2) improved performance of ISM as compared with conventional grab sampling techniques, and (3) ease of ISM implementability. Comparisons of ISM with the conventional grab sampling methodology demonstrated, in general, that ISM provided results more representative and reproducible for all three demonstration sites, consistent with our initial study using the results from Camp Ethan Allen.

Distributional heterogeneity was addressed by collecting at least 30–100 increments over the entire Decision Unit. However, multi-increment field sampling alone is insufficient to overcome the distributional and compositional heterogeneity in the soil samples. Modifications to laboratory sample preparation procedures using United States Environmental Protection Agency (USEPA) Method 3050B are also necessary to reduce variability owing to sample heterogeneity, and *Incremental Sampling Methodology (ISM) for Metallic Residues* (Clausen et al. 2013b) outlines a proposed protocol. The proposed changes for metals adopted many of the recommendations for energetics outlined in USEPA Method 8330B, such as air drying, milling, larger acid volumes to soil digestion ratios, larger digestion masses, and subsampling. Two types of milling equipment (e.g., ball mill and puck mill) yielded satisfactory results. As the digestion procedures in USEPA Method 3050B resulted in poor antimony and tungsten recoveries, Clausen et al. (2013a) developed and discussed alternative digestion methods for these metals.

In general, the demonstration results met the targeted performance criteria using ISM. However, there were instances where the performance criteria were not met (e.g., copper). In these situations, the results indicate that the ISM approach used did not adequately deal with the extreme con-

taminant heterogeneity. Consequently, in some situations, an iterative approach may be necessary whereby the ISM process is modified to meet the performance objectives (e.g., increasing the milling interval, increasing the number of increments collected, increasing the digestion mass, etc.).

Implementation Issues

In addition to the published *Incremental Sampling Methodology (ISM) for Metallic Residues* (Clausen et al. 2013b) protocol, the authors of this report are currently working with the USEPA to modify Method 3050B by incorporating the recommended changes identified from this project into a proposed Method 3050C. These changes include modifying the sample preparation methods and incorporating an Appendix outlining the multi-increment field sampling approach, similar to what was done for USEPA Method 8330B. In the interim, *Technical and Regulatory Guidance: Incremental Sampling Methodology* (ITRC 2012) is a good reference for understanding and implementing ISM. Protocols specific to ISM sampling of sites with metallic residues can be found in *Incremental Sampling Methodology (ISM) for Metallic Residues* (Clausen et al. 2013b).

There are no known limitations to the application of ISM as the equipment used for ISM is the same as that for conventional grab sampling. Implementation costs for ISM are lower than for conventional grab sampling because fewer samples are collected, prepared, and analyzed (e.g., fewer supplies are consumed, less time is required to survey and select sample locations, and there are fewer samples to label and to ship to the laboratory for preparation and analysis). As multiple increments need to be collected to prepare a single sample in the field, the collection time for a single incremental sample is greater than that for a single grab sample. However, as a relatively large number of grab samples are typically needed to provide data of comparable quality as a few incremental samples, ISM does not typically increase the total sample collection time.

Once the SU corners are surveyed, there is no need to survey individual sample increment locations. In contrast, each grab sample requires surveying each location. ISM samples typically have a larger mass than conventional grab samples, resulting in greater per sample shipping costs and sample preparation fees. Fewer ISM samples are analyzed than grab samples. The greatest cost saving is incurred at the laboratory preparation step, owing to fewer samples requiring preparation and analysis. Again, some additional costs are incurred with the addition of the milling and

subsampling step. However, the increased costs are more than offset by the fewer number of ISM samples. Although per-sample (unit) costs are higher for ISM, the total cost of soil sampling and analysis for ISM will generally be less than that for conventional grab sampling.

Cost savings are difficult to quantify because there is no standard procedure for determining the number of soil samples needed to characterize a study area. Conventional grab sampling designs are frequently judgmental in nature, entailing subjective criteria for selecting sampling locations and the number of soil samples. However, based on a review of current practices, case studies, and the results of the demonstration at the three sites, ISM can result in a cost savings of 30%–60% relative to conventional grab sampling methods.

1 Introduction

Since the publication of United States Environmental Protection Agency (USEPA) Method 8330B (USEPA 2006) for explosives, there have been efforts to develop incremental sampling methodologies (ISM) for other analytes, particularly metals. However, there are no published procedures for the laboratory processing of incremental samples for analytes other than energetic compounds. Sample collection and laboratory processing procedures for ISM depend on the nature of the analytes of interest. The laboratory procedures of Method 8330B, which were developed specifically for explosives and propellants, generally need to be modified for other analytes. For example, the drying, sieving, and milling procedures for soil samples described in Method 8330B are inappropriate for volatile organic compounds. Depending on the types of analytes of interest, milling can bias analytical results because of analyte losses or the addition of spurious contaminants. However, because milling increases precision, the larger improvements in precision may outweigh the magnitude of the biases.

After the release of USEPA Method 8330B, a growing concern within the United States Department of Defense (DoD) and in Federal and State agencies has been that similar protocols should be adopted for the characterization of metals on training ranges and at other locations. A variety of metals are used in military munitions (Clausen et al. 2012a, 2010, 2007; Clausen and Korte 2009a,b). As munitions containing metals are frequently used on Army training lands, metals deposited by rounds can accumulate in soils. Although the deposition of metals at military ranges has only been studied on a limited basis, like explosives, metal deposition appears largely spatially heterogeneous. Anthropogenic metals are heterogeneously distributed over training ranges as particles of various sizes, shapes, and compositions. To obtain representative samples (i.e., to ensure mean contaminant concentrations in the samples will be similar to the mean concentrations in the environmental population) and to obtain reproducible estimates of the mean, the sampling design and laboratory sample preparation and analytical methods need to address compositional and distributional heterogeneity.

1.1 Background

The development of ISM began with the realization in the early-1990s that energetic residues were heterogeneously distributed on ranges and that the current sampling methodologies did not address this issue. Consequently, early studies of energetics yielded non-reproducible and non-representative results, the result being that some sites potentially underwent remediation that was not necessary; or conversely, some sites implemented no needed remedial activities. Studies conducted in the early 2000s resulted in the development of a modified sample collection and processing methodology for energetic constituents, referred to as ISM.

Anthropogenic metals are also heterogeneously distributed over active training ranges as particles of various sizes, shapes, and compositions. To address the compositional and distributional heterogeneity (e.g., to obtain a representative and reproducible estimate of the mean concentration), the sampling strategy must acquire an adequate number of particles of the constituents of interest; and these particles must be present in the sample in roughly the same proportion as the in Decision Unit (DU). The DU is an area or volume of soil of interest (i.e., an environmental population) about which one plans to make some decision based on the outcome of the sample data. Unless stated otherwise, each DU refers to a single Sampling Unit (SU), an area (or volume) of soil over which a set of grab samples or incremental samples have been randomly collected. In general, a DU can consist of multiple SUs, where each SU is characterized or represented by a set of n grab or incremental samples, where n denotes the “sample size” (i.e., the number of independent grab or incremental samples collected from the SU).

The ISM approach is not limited to laboratory sample processing; it also includes field sampling procedures and project planning (Table 1).

Table 1. Incremental sampling methodology for metallic residues.

Project Stage	Specific Activity	
Project Planning	Development of conceptual site model	
	Determination of investigation objectives	
	Identification of data needs	
	Decision Unit identification	
	Determination of sample depth interval	
	Number of increments per sample	
Field Implementation	Sample tool selection	
	Decision Unit delineation	
	Collection of soil sample	
Sample Processing	Air drying	
	Sieving	
	Particle size reduction (milling)	
	Less than 2 mm (examined)	Greater than 2 mm (examined and archived)
	Splitting (if necessary)	
	Subsampling to build digestate	
	Metals digestion	Energetics extraction
Analysis	ICP-MS or ICP-OES	HPLC

Well-defined Data Quality Objectives usually need to be established during project planning to successfully implement ISM. Key elements to address during the planning phase include

1. The conceptual site model (CSM).
2. The project's objectives.
3. The spatial boundaries of each DU.
4. The sampling depths.
5. The number of increments per sample.
6. The number of samples per DU.

To reduce the influence of compositional and distributional heterogeneity for estimating mean concentrations of energetic analytes within a DU, Method 8330B recommends collecting 30 or more evenly spaced increments to build a sample with a total sample mass greater than 1 kg (Clausen et al. 2012b, 2013a; Jenkins et al. 2004a,b, 2005, 2006; Walsh et al. 2005; Hewitt et al. 2005, 2007). The objective of this sampling technique is to obtain a representative portion of every particle size, composi-

tion, and configuration (e.g., spheres or elongated particles) and to avoid over- or under-sampling any portion of the DU. This same situation applies to small-arms ranges where residues of antimony, copper, lead, zinc, and others are present. Instead of collecting and analyzing individual grab samples and integrating the results over an area of interest (DU) or assuming a single grab represents the entire area, samples are prepared by combining a number of soil increments collected over the entire DU to obtain a sample mass of approximately 1 kg. The increments can be collected using simple random sampling or systematic random sampling. For systematic random sampling, the sampler selects a random starting point and collects evenly spaced increments as the sampler walks back and forth from one corner of the DU to the opposite corner (Fig. 1).

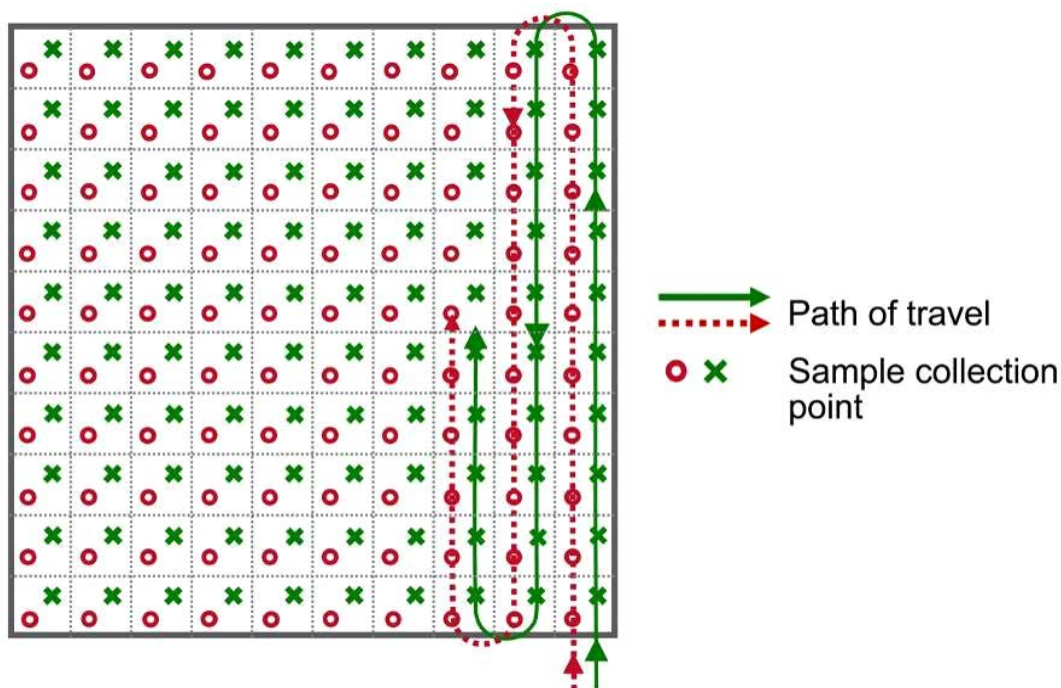


Figure 1. Example of multi-increment sampling using a systematic random sampling design for collecting two separate 100-increment samples.

1.2 Objective of the demonstration

The primary objective was to demonstrate that the ISM approach can be readily implemented for soils with metallic residues and provides a higher quality data than conventional grab sampling and analysis methods. Specifically we wanted to demonstrate that

1. Using ISM provides reproducible results.
2. Milling of soil for the analytes of interest (antimony, copper, lead, and zinc) does not introduce significant bias.
3. Application of ISM yields sample results more representative of the mean soil concentration than do conventional grab samples.
4. ISM involves lower total project soil sample costs.

Further, we wanted to demonstrate the robustness of ISM with a variety of soil types by selecting three field sites for the demonstration. The working hypothesis was that the current field sampling approach using grab or discrete samples and sample processing procedures following USEPA Method 3050B for metals in soil do not yield representative and reproducible results for military sites where the metals were heterogeneously introduced into the environment as a solid residue. This demonstration of ISM met all of the objectives as discussed in Section 3.

1.3 Regulatory drivers

In 2001, the Defense Environmental Restoration Program established the US Army's Military Munitions Response Program (MMRP) to manage the environmental and health and safety issues associated with unexploded ordnance (UXO), discarded military munitions, and munitions constituents on non-operational ranges in active installations, Defense Base Realignment and Closure (BRAC) sites, and Formerly Used Defense Sites (FUDS). Under the MMRP, the DoD is required to (1) inventory non-operational ranges that contain or are suspected to contain munitions-related material released before September 2002; (2) identify, characterize, track, and report data on MMRP sites and clean-up activities; and (3) develop a process to prioritize site cleanup and to estimate costs. The Army completed their inventory of non-operational ranges in 2003 and began Site Investigations (SI) for these MMRP sites. Based on the SI findings, some ranges may require additional assessment under the Remedial Investigation (RI) process. In addition, established directives mandate that all active DoD facilities implement procedures to assess environmental impacts from munitions on training and testing ranges (DoD 2007; DoD 2005).

Because of the success of ISM for energetics, members of the environmental community are increasingly requiring its use for other hazardous par-

ticulate constituents, such as metals (ITRC 2012; Alaska 2009; Hewitt et al. 2011, 2009; Hawaii 2008). The approach is frequently used for SIs conducted under FUDS. The USEPA has issued guidance for characterizing MMRP sites using ISM (Hewitt et al. 2011) as has the US Army Corps of Engineers (USACE) (USACE 2009). Several state and federal agencies now require ISM designs. These currently include the states of Alaska, Hawaii, and the USEPA Region 6. Other states, such as Florida, are considering rewriting their environmental regulations to require ISM. Additionally, in Massachusetts, Michigan, Missouri, and New Jersey, formal guidance is not available; but in some situations, where appropriate, these states require ISM. It is likely that additional states and USEPA regions will increasingly require ISM, thus requiring DoD MMRP and Operational Range Assessments (ORAP) to employ ISM.

2 Technology

2.1 Technology description

The technology demonstrated is the use of ISM to characterize surface soil in an area of interest (DU) containing metallic residues from training with military munitions constituents. To address the heterogeneous nature of contaminated soils, ISM entails changes to conventional field sampling methodology and to laboratory sample preparation procedures. ISM also requires well-defined project data quality objectives (DQOs) during systematic planning (Table 1). The planning phase should happen prior to any field sampling. The Interstate Technology Regulatory Council (ITRC) document (ITRC 2012) provides an overview of the ISM process, and Clausen et al. (2013b) discuss applications specific to sites with deposition of metallic residues.

In the field, the first step is to define the boundaries of the DU with markers (typically flags or stakes). The next step is to determine the approximate spacing between increments (e.g., if increments are collected using systematic random sampling) and the number of rows of increments needed to achieve the total number of increments for each ISM sample. Once the DU is identified, distributional heterogeneity is addressed by collecting a 1- to 2-kg incremental sample prepared from at least 30 to 100 “increments” collected randomly over the entire DU (Fig. 1). The objective of this sampling technique is to obtain a representative portion of every particle size, composition (antimony, copper, lead, zinc, etc.), and configuration (e.g., spheres or elongated particles) and to avoid over- or under-sampling any portion of the DU. The increments can be collected using simple random sampling or systematic random sampling. For systematic random sampling, a random starting point is selected; and evenly spaced increments are collected as the sampler walks back and forth from one corner of the DU to the opposite corner (Fig. 1).

The “increments” that are combined to prepare each incremental sample typically refer to cylindrical soil cores collected by using a coring device such as the “Cold Regions Research and Engineering Laboratory (CRREL) Multi-Increment Sampling Tool” (CMIST) (Walsh 2009).

The number of replicate ISM samples required for each DU needs to be determined during project planning. The sample size will depend on the degree of variability, the tolerances for decision errors, and the differences in concentrations considered significant. Typically, at least three replicates are independently collected from the DU to characterize the total variability. Additional replicates (e.g., a minimum of eight samples) will likely be needed for statistical evaluations (e.g., for two-sample hypothesis tests) when a normal distribution cannot be assumed.

If metal residues and energetics are both contaminants of interest and separate incremental samples are not collected in the field for metals and explosives, to control distributional and compositional heterogeneity, each sample must be split in the laboratory in a manner that is consistent with Gy's sampling theory and practice (Gy 1992). Standard operating procedures to split, mill, and subsample should be developed on a project-specific basis and should be consistent with the guidance in ASTM D6323 (ASTM 2003) and Gerlach and Nocerino (2003). As shown in Clausen et al. (2012b), successful splitting of unmilled samples with a high degree of heterogeneity usually does not occur even with a rotary splitter where many increments are collected for each split. However, using a rotary splitter following drying, sieving, and milling, Clausen et al. (2012b, 2013a) repeatedly demonstrated reproducible results with splitting for samples containing metal particles.

Clausen et al. (2012b, 2013a,b) discuss in detail protocols for the laboratory processing of incremental samples for metals. In general, when soils contain metal particulates (e.g., bullet fragments), the entire sample should be dried, sieved, and then mechanically pulverized. Table 2 summarizes the proposed changes to the sampling processing procedures for Method 3050B. Unfortunately, unlike for explosives, a "universal" grinder is not currently available for processing samples for metals although Clausen et al. (2012b, 2013a) obtained good success with the puck and roller mill. The grinder needs to be selected on the basis of the metals that are of primary interest for each project. Most commercial crushing or grinding equipment possess working surfaces composed of metal alloys containing iron, chromium, tungsten (carbide), etc. These grinding surfaces have been demonstrated to introduce metal contamination during sample processing (Clausen et al. 2012b) though Felt et al. (2008) indicate the effect on soil concentrations is minimal. However, if needed or desired,

non-metallic materials are available, such as an agate bowl and puck for the puck mill and Teflon coated cans and ceramic chips for the roller mill.

Table 2. Salient differences between Method 3050B and proposed Method 3050C.

Activity	Method 3050B/ Conventional Sampling	Method 3050C/ Incremental Sampling Method
Field Sampling	Not explicitly addressed in method. Typically, grab samples are collected with a metal scoop from biased sample locations.	An incremental sample consists of 30–100 “increments” collected randomly over the entire Sampling Unit (e.g., using a systematic sampling). For cohesive surface soils, an “increment” typically consists of a small soil cylinder (e.g., 2–5 cm in length) that was collected using a 2- to 4-cm diameter coring device (e.g., as shown in Fig. 2).
Sample Mass and Containers	Typically, about 200 g of soil in 4-oz, wide-mouth, screw-top jars.	Typically, 1–2 kg of soil in clean large (e.g., 15 × 15 in., 6-mm thick) polyethylene plastic bags sealed with Ty-wraps,
Sample Drying	Sample drying is optional and is not typically done.	Sample is air-dried at room temperature by spreading it onto a tray to form a relatively thin uniform slab.
Sieving	“...sieve, if appropriate and necessary, using a USS #10 sieve...” Soil samples are typically not sieved.	Samples are routinely passed through a #10 (2-mm) sieve. Both size fractions are weighed and a less than 2-mm fraction is additionally processed.
Milling	“Wet samples may be dried, crushed, and ground to reduce sample variability...” Milling is typically not done.	Samples are routinely milled using appropriate mechanical grinders, such as puck mill or roller mill. Milling must result in finely ground material of uniform appearance and texture.
Laboratory Subsampling	“Mix the sample thoroughly to achieve homogeneity...” Soil is often stirred with a spatula or a similar device (often in original container), and a single aliquot (e.g., scooped from the top of the container) collected as the subsampled for digestion and analyses.	After grinding, the soil is spread onto a large tray to form a thin slab of material of uniform thickness. At least 20 small aliquots are randomly collected over the entire slab with a spatula or similar device and composited to prepare a subsample for digestion and analysis.
Subsample Mass	1- 2-g wet weight or 1-g dry weight.	2–10-g dry weight.

Digestion generally follows the procedures outlined in USEPA Method 3050B with the following changes. Clausen et al. (2013b) recommends that more than 1 g of material be digested, preferably 5 g, maintaining a 1:1 ratio of acid to soil. Digestion masses greater than 5 g are potentially problematic because of foaming or effervescence that can overtop the 100-ml vials used with standard digestion blocks. Clausen et al. (2012b) found im-

proved sample reproducibility and reduced sample variability with increasing mass with the differences statistically significant although the magnitude of change was small.

The ISM discussed above is based on studies with energetics (Hewitt et al. 2009; Walsh et al. 2009) and transitioning into metals (Clausen et al. 2012b). Table 3 provides a chronological summary of the development of ISM.

Table 3. Chronological summary of multi-increment sampling.

Time Period	Activity	References
1960s–1990s	Recognition of the role of heterogeneity in distribution of metals in mining samples and development of methods to obtain representative samples	Duncan 1962, Johanson 1978 Elder et al. 1980 Gy 1992, 1999 Wallace and Kratochvil 1985 Pitard 1993 Leutwyler 1993 Studt 1995
Early 1990s–2004	Demonstration of presence of energetic residues on ranges	Racine et al. 1992 Jenkins et al. 1997a,b, 1998, 2001 Walsh and Collins 1993 Walsh et al. 1997 Thiboutot et al. 1998, 2000a,b, 2003 Ampleman et al. 2003a,b Clausen et al. 2004 Pennington et al. 2004 Taylor et al. 2004
1990s	Recognition of heterogeneity issues associated with environmental samples	Pitard 1993 Jenkins et al. 1996
Mid-1990s–Early 2000s	Recognition of heterogeneity issues for energetic constituents on military ranges	Racine et al. 1992 Jenkins et al. 1997a,b, 1999, 2001 Taylor et al. 2004 Walsh et al. 1993, 1997
2001–2009	Development of sampling and sample processing methods for soils containing energetic constituents	Jenkins et al. 2001, 2004a,b, 2005, 2006 Thiboutot et al. 2002 Walsh et al. 2002, 2003, 2005, 2006 Walsh and Lambert 2006 Hewitt and Walsh 2003 Hewitt et al. 2005, 2007, 2009

Time Period	Activity	References
2004–2007	Demonstration and comparison of ISM with traditional grab sampling approach for soils with energetic constituents	Jenkins et al. 2004a,b Walsh et al. 2004 Hewitt et al. 2005 Nieman 2007
2007–2010	Demonstration of heterogeneous distribution of metals in soils from military ranges	Clausen et al. 2007, 2010 Clausen and Korte 2009a,b
2008–present	Adoption of ISM for soils with metals	Hawaii 2008 Alaska 2009 ITRC 2012
2009–present	ESTCP ER-0918 Project	Clausen et al. 2012b, 2013a,b

Although our report is specifically focused on the application of ISM at small-arms ranges, it has potential application to any site where solid metallic residues are introduced into the environment. At military installations, this could include impact areas where artillery, mortar, or anti-tank rockets were fired as these munitions contain metals in the ordnance casing. In addition, many pyrotechnic devices contain metallic salts (Clausen et al. 2012a); so if training or maneuver areas are being sampled where these devices have been used, then ISM is appropriate. The protocol outlined in Clausen et al. (2013b) was successfully used for sampling metallic residues derived from pyrotechnic training (Clausen et al. 2012a).

2.2 Advantages and limitations of the technology

The advantages of ISM include (1) a soil sample representative of the area of interest (i.e., DU), (2) the ability to quantify the uncertainty for field sampling and laboratory sample and analysis, and (3) the reduction in the number of field samples collected for laboratory analysis (Table 4). The disadvantages of ISM include (1) an increased volume of individual samples sent to the analytical laboratory; (2) the necessity of a particles size reduction step (e.g., milling) during sample preparation; and (3) the alteration of the soil matrix during the particle size reduction step, possibly changing the availability of some metals during acid digestion (Table 4).

Table 4. Comparison of the advantages and disadvantages of the Incremental Sample Methodology.

Activity	Advantages	Disadvantages	Comment
Total Sample Error	☑		Quantification of error possible with ISM
Number of Soil Samples	☑		Fewer samples needed with ISM
Individual Sample Mass		☑	Greater sample mass to handle heterogeneity
Precision of Result	☑		Greater sample result precision with ISM
Laboratory Preparation		☑	More involved with ISM (drying, milling, subsampling)
Field Costs	☑		Fewer samples to collect and ship with ISM
Sample Preparation Costs		☑	Higher costs due to more involved processing
Soil Matrix Alteration		☑	Possible changes to metal recovery due to milling
Milling Cross-Contamination		☑	Possible metal cross-contamination from milling when using metallic components
Metal Ratios Analysis		☑	The averaging effect of ISM is not conducive for metal ratio analysis

Processing of soil samples so that they can be reproducibly subsampled often involves a particle size reduction step, such as milling (Clausen et al. 2012a, 2013a). Increasing the surface area of a soil matrix may make some metals more available for acid digestion. A recent study using three soil types and three grinding techniques compared results with those obtained for samples that were blended without pulverization. Overall, the milling step increased precision and only slightly increased metal concentrations (Felt et al. 2008). One potential drawback is that samples from areas of concern (DU) and from background locations will need to be collected and processed using the same protocols. This requirement may increase the number of samples that need to be collected, processed, and analyzed, thereby increasing costs as compared to using conventional approaches. However, collection of an inadequate number of samples with conventional sampling designs often results in large data variances, making reliable quantitative statistical comparisons difficult. Background comparisons using ISM can often be done using smaller numbers of samples as the approach tends to reduce the variability and to normalize distributions. It is

anticipated that the use of ISM will significantly increase the data quality of background samples (Clausen et al. 2013a).

The advantages of ISM include lower variability and an absence of skewed distributions (Clausen et al. 2012b, 2013a). In contrast, large variability and positive skewed distributions are normally observed for grab samples. In addition, ISM yielded highly reproducible results and high precision with percent relative standard deviations (RSDs) of less than 30% for replicate samples, suggesting that distributional heterogeneity was reasonably controlled. In contrast, measured RSDs for the grab samples typically yielded values greater than 30% and, in some cases, in the hundreds of percent. The results of the demonstration study also suggested that ISM improved the accuracy of estimates of the mean; grab samples often underestimate the population mean. In general, the ISM results exhibit a higher mean concentration than grab sample results. This situation occurred in 60% of the sample results for the three demonstration sites and the one experimental site for the metals of interest (antimony, copper, lead, zinc). In instances where the ISM mean was less than the grab sample mean, the data exhibited greater variability than desired. It seems likely that sample precision and accuracy could have been improved by taking all or some of the following steps:

1. Increasing the number of increments collected from the DU.
2. Increasing the sample mass collected from the DU.
3. Increasing the number of subsampling increments to build the digestion aliquot.
4. Increasing the digestion mass.

One of the advantages of ISM is the ability to assess the total sample error or the error associated with specific steps of the ISM process, allowing for the establishment of performance criteria. If the criteria are not met initially the ISM process can be altered to meet one's sample-quality objectives.

One of the limitations of ISM is the necessity for collecting, at minimum, 30 increments from the DU; otherwise, collecting fewer than 30 increments results in poorer data precision (Clausen et al. 2012b). An RSD of less than 30% was generally achieved when the number of increments exceeded 30. However, fewer than 30 increments collected resulted in RSDs

greater than 30%. Thus, for the situation studied, more than 50 increments were not necessary; but clearly fewer than 30 were inadequate to obtain reproducible results. Additionally, owing to the large number of increments collected within a DU, ISM tends to result in better spatial coverage and, therefore, in more representative samples than the conventional grab sampling approach. However, ISM is insufficient in and of itself to overcome the distributional and compositional heterogeneity in the soil samples. Modifications to laboratory sample preparation procedures are also necessary to reduce variability owing to sample heterogeneity (Gy 1992, 1993; Walsh et al. 1993, 2002, 2005; Gerlach and Nocerino 2003; Hewitt et al. 2007, 2009, 2011; Clausen et al. 2012b, 2013a,b; ITRC 2012).

Another potential limitation is that the typical puck mill used by commercial environmental laboratories contains metal components. Studies by Clausen et al. (2012b) indicated a potential for a significant increase in chromium, manganese, nickel, and vanadium concentrations as a result of cross-contamination from a metallic puck and bowl. However, contamination from the puck mill seems minimal for the small-arms range metals antimony, copper, lead, and zinc (Clausen et al. 2013a; Felt et al. 2008). Further, the cross-contamination issue becomes less important as the metal concentration of the sample increases relative to the regulatory action level. For antimony, copper, lead, and zinc, the potential concentration increase from cross-contamination resulting from milling is less than 5 mg/kg. This may be problematic in situations where the expected DU soil concentration is within several mg/kg of an action level. However, cross-contamination issues can be avoided by using an agate puck and bowl, although these are more expensive than the metallic versions and process less material owing to their smaller size. Another alternative is the use of a Teflon lined roller mill with Teflon chips, which yielded acceptable results (Clausen et al. 2012b). It should be kept in mind that many of the small-arms range bermed soils we have sampled often have lead levels, typically the principal metal of interest, in the 1000s to 100,000s mg/kg range; but the decision limit for lead is often only 400 mg/kg.

3 Performance Objectives

There were three quantitative performance objectives and one qualitative performance objective for the demonstration and validation of the technology (Table 5). The quantitative performance objectives were sample reproducibility, lack of sample bias, and cost reduction. The qualitative performance objective was ease of technology use. The effectiveness of the technology for soil sampling is predominately a function of the precision of replicate laboratory subsample results from the same field ISM sample and the precision of replicate field ISM sample results from the same DU. ISM's effectiveness was evaluated by collecting replicate ISM soil samples from each DU and comparing them against multiple grab samples collected from the same DU (Clausen et al. 2013a). Fifteen replicate ISM samples were collected at the small-arms firing range berm DU at three sites: Fort Wainwright, Fort Eustis, and Kimama Training Site (TS). From the same DUs, 50 grab samples were collected from Fort Wainwright, 30 from Kimama TS, and 33 from Fort Eustis.

An evaluation of sample variability was performed using statistical comparisons at the 95% level of confidence. The null hypothesis is no difference between the variances of the population of grab and ISM samples. The results indicated a significant difference between variances for the two populations with lower variances evident for ISM as compared to grab samples; observations were consistent with earlier findings (Clausen et al. 2013a). A secondary goal was to achieve a percent RSD of less than 30 for field replicates from the same DU and less than 15 for laboratory subsample replicates with ISM. This objective was met with ISM but not with the grab sampling approach.

Positive bias (e.g., owing to milling during sample preparation) was also evaluated by processing method blanks consisting of glass material (Clausen et al. 2012b). Glass samples were milled before and after a batch of soil samples were milled. A total of 7 soil samples were processed in this manner. Bias was evaluated using the criteria summarized in Table 5. For the metals of interest (antimony, copper, lead, and zinc) there was no evidence of an increase in the glass blank samples between pre- and post-milled samples (Clausen et al. 2012b, 2013a). Laboratory Control Samples

(LCS) were also processed with each sample batch to evaluate bias. Again there was no evidence of sample bias for the analytes of interest.

Table 5. Performance objectives for ER-0918.

Performance Objective	Data Requirements	Success Criteria	Performance Objective Met
Quantitative Performance Objectives			
Obtain reproducible results for surface soil samples containing metal particles	Field and laboratory replicates analyzed for metals.	Demonstrate statistically significant decreases for variability (with 95% confidence) for replicate field samples and replicate laboratory subsamples compared with replicates processed using conventional methodology. RSD $\leq$ 30% for field replicates within same DU RSD $\leq$ 15% for lab replicates (for concentrations > 100 mg/kg)	Yes Yes Yes
Evaluate bias	Method blanks (MBs) and laboratory control samples (LCSs) processed with ISM.	Concentrations < $1/10$ the ISM sample concentrations. LCS recoveries should be 70%–130% of the expected values or of the manufacturer's specifications.	Yes
Compare performance of ISM sampling and grab sampling for metals in soils	Samples collected using multi-increment sampling approach and grab sampling designs. Hours or cost of field sampling effort and of sample preparation and analysis recorded.	ISM sampling design results in equivalent or superior estimates of the mean with less analyses and results in a cost savings of at least 20%.	Yes
Qualitative Performance Objectives			
Implementability	Feedback collected from field and laboratory personnel.	ISM sampling approach can be readily implemented given appropriate equipment and planning.	Yes

The third quantitative performance objective was that ISM yields a total reduction in cost of 20%. Total reduction accounts for both physical collection of soil samples in the field and sample preparation back in the laboratory. This objective was evaluated by monitoring the manpower and length of time needed for

1. Field mobilization preparation.
2. DU/sample location determination and flagging.
3. Surveying of sample locations/DU.

4. Physical collection of samples.
5. Shipping costs of samples.
6. Sample preparation costs.
7. Sample analysis during the demonstration study (Clausen et al. 2013a).

Sample preparation activities assessed the unmilled approach following USEPA Method 3050B and the milled approach following our modified Method 3050B approach, referred to as Method 3050C. These activities were done for both ISM and grab samples. The outcome of this cost comparison was that total ISM costs were at least 20% less than the conventional grab sampling approach (Clausen et al. 2013a).

Because the number of grab samples is often determined subjectively, environmental consultants experienced with SI and RI were polled for a hypothetical small-arms range berm. The consensus seems that for a 3-m high by 100-m long berm, 7 to 10 grab samples would be appropriate; however, those with a statistical background and training preferred a greater number of grab samples. ISM becomes less cost effective relative to grab sampling as the number of grab samples decreases. However, precision and, presumably, accuracy (for estimating the DU mean) for grab sampling significantly decreases as the sample size decreases and is usually poor relative to that provided by an equal number of incremental samples. As ISM and conventional judgmental sampling designs using grab samples do not result in comparable data quality, comparisons based solely on the per-unit costs of sampling and analysis do not accurately characterize the cost of ISM relative to conventional sampling.

Implementability is a qualitative performance objective that assesses feedback from field and laboratory personnel about the ease of use of ISM. Ease of use also includes the availability of tools to implement the ISM and sample processing procedures. The discussions about field sampling indicate little difference between ISM and the conventional grab sampling approach as the same field equipment is used for both (Clausen et al. 2013a). The only major difference during sample preparation in the laboratory is with milling the sample. Implementation of this step is limited in the sense that the majority of commercial environmental analytical laboratories do not have milling equipment. However, for those laboratories that have milling equipment, the ISM approach is readily implementable.

4 Site Description

Instead of conducting the demonstration and validation at a single site, the demonstration and validation was conducted at three sites to ensure robustness of the technology. The three sites selected and discussed below are Kimama TS in Idaho, Fort Eustis in Virginia, and Fort Wainwright in Alaska. Clausen et al. (2013a) provides detailed site descriptions.

4.1 Site location and history

4.1.1 Kimama Training Site

The Kimama TS is located in south-central Idaho in Lincoln County, approximately 17 miles northwest of Minidoka off of Highway 24, and was used by the Idaho Army National Guard for armored and small-arms training (Fig. 2). The area where the demonstration was conducted is referred to as Training Area 3, which encompasses approximately 14,322 acres (Fig. 2). Training Area 3 was used for armored maneuver training from 1974 through 1993. Three small-arms ranges are located in the northwest portion of Training Area 3 and were used from 1969 through 1993. The three small-arms ranges encompass 2355 acres of Training Area 3.

Munitions used at Training Area 3 included small arms, star clusters, riot control grenades, trip flares, practice mortar fuzes, 40-mm practice rifle grenades, and M69 practice hand grenades. The ordnance used on the small-arms ranges included 7.62 mm, .45 cal, .22 cal, and .50 cal. The munitions constituents (lead and antimony) were detected at levels on the small-arms range during a SI, warranting further investigation (FPM 2009). An RI/FS (feasibility study) project plan was prepared in October 2010 with the fieldwork anticipated to be complete by July 2011 (FPM 2010). Clausen et al. (2013a) presents additional details about the firing range.

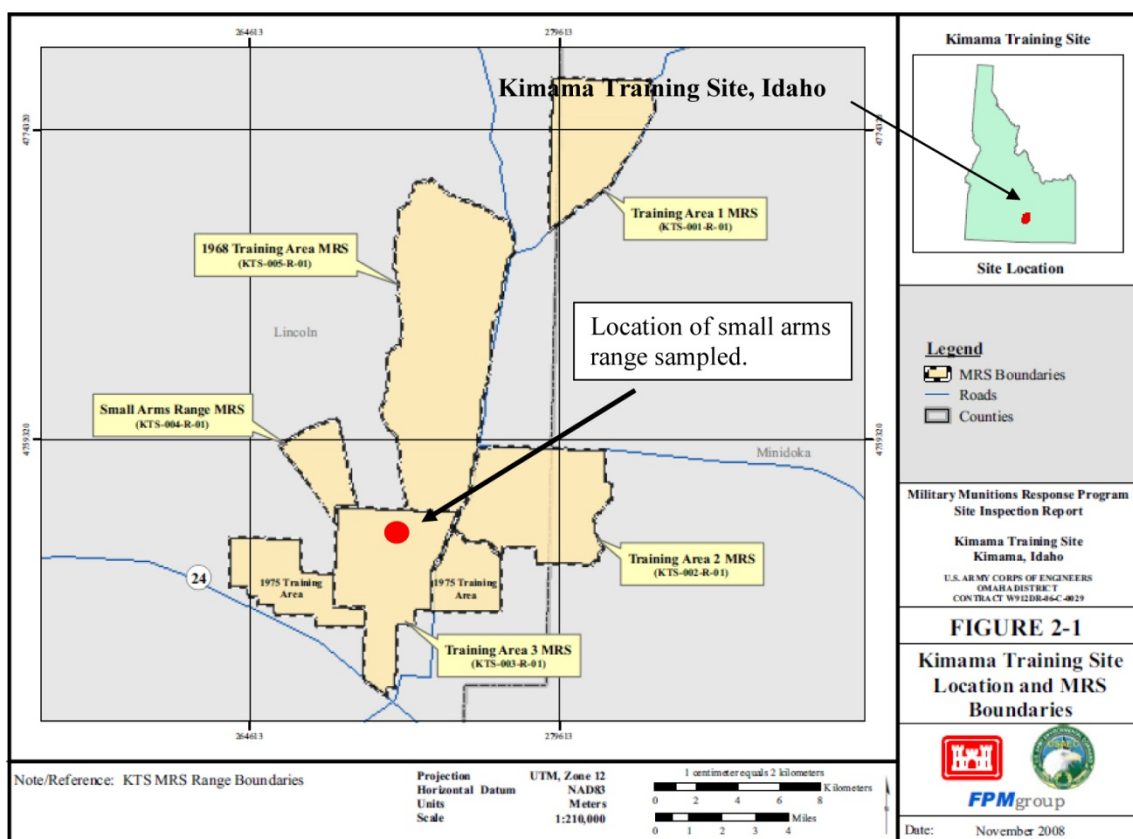


Figure 2. Location of the Kimama Training Site, Idaho, and location of Training Area 3 (FPM 2009).

4.1.2 Fort Eustis

Fort Eustis is located within the geographic boundaries of Newport News, Virginia, in the southeastern portion of the state. The demonstration conducted within the cantonment area of Fort Eustis occurred on the northern berm of the 1000-in. Rifle Range (Fig. 3). The 1000-in. Rifle Range is a former small-arms training range used between 1920 and 1941 for target practice using .22-, .30-, and .45-cal munitions. The 1000-in. Rifle Range is estimated to cover 18.5 acres. Clausen et al. (2013a) presents additional details about the firing range.

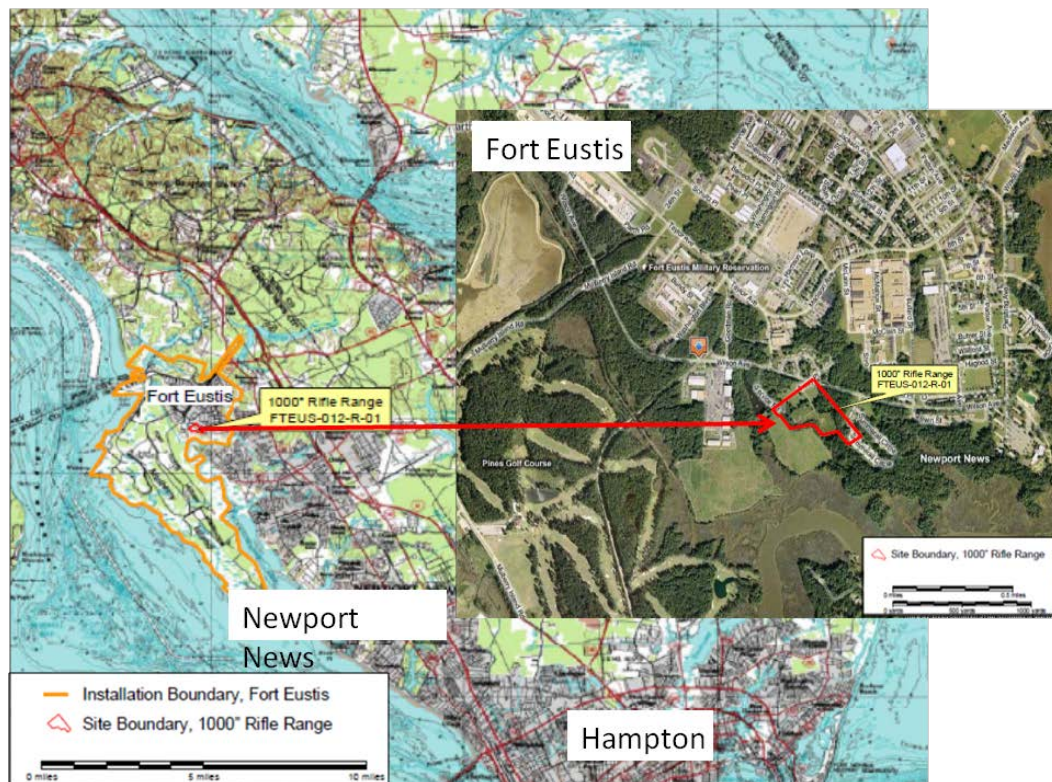


Figure 3. Map showing location of Fort Eustis, VA, and the 1000-in. Rifle Range.

4.1.3 Fort Wainwright

Fort Wainwright is located in central Alaska near Fairbanks and covers approximately 910,498 acres (Fig. 4). The demonstration was conducted on the active Range 16 Records Range of the Fort Wainwright Small Arms Range Complex located off of (south) Richardson Highway. All manner of small-arms ammunition is used on the Range 16 Record Range.

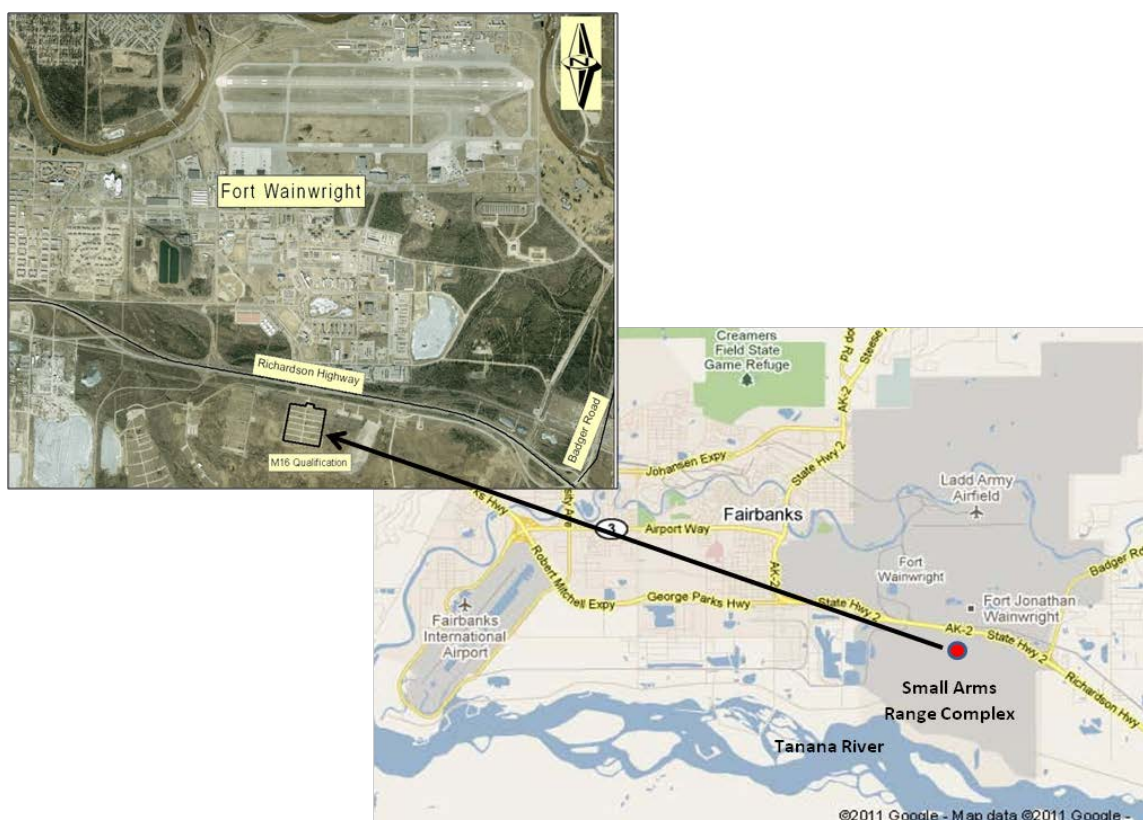


Figure 4. Map of Fort Wainwright, AK, and the Range 16 Record Range.

4.2 Site geology and hydrogeology

4.2.1 Kimama Training Site

The Kimama TS is located in the Snake River Plain, and the surface lithology consists largely of approximately 3- to 15-m-thick sections of interbedded lacustrine and fluvial sediments of volcanic origin (Idaho Geological Survey 2011). Soils are also volcanic in character and consist mainly of silt loam with some small areas of sandy loam. Analysis of two samples (KTS45 and KTS48) yielded a determination of poorly graded sand with silt (Clausen et al. 2013a). Some aeolian soil erosion has occurred in the vicinity of previous tank activity, particularly in the sandier soils (USACE 1972).

The Kimama TS overlies the Snake River Aquifer, which occurs in basalt and sediments of the Snake River Group. The aquifer is located at an estimated depth of 100–150 m below ground surface. Groundwater flows most rapidly in the upper 60 m, which is the most productive portion of

the aquifer and is associated with permeable zones consisting of the tops and bottoms of the basalt lava flows (USACE 1972).

4.2.2 Fort Eustis

Fort Eustis lies within the Atlantic Coastal Plain and consists of unconsolidated and interbedded sands and clays with minor amounts of gravel and shell fragments. Locally, the site geology consists of impermeable clays, silts, and clayey sand with sand and silty sand lenses. Previous grain size analysis of soils from the 1000-in. Range was characterized as silty sand (URS 2010), which is consistent with particle size samples (BCK1C) collected during this study (Clausen et al. 2013a). Groundwater is present at a depth of 4–6 m below ground surface with flow towards the Warwick River (URS 2010). The 1000-in. Rifle Range drains through marshes into the Warwick River.

4.2.3 Fort Wainwright

The Fort Wainwright Small Arms Complex is located on surficial material consisting of well-stratified layers and lenses of unconsolidated sand and rounded river gravel overlain by as much as 5 m of silt. Gravel consists mostly of quartz and metamorphic rock with clasts ranging from 0.6 to 7 cm in diameter. The gravel is 3 to more than 125 m thick. It is locally perennially frozen down to 90 m with low ice content. In addition, there are discontinuous swales and slough deposits consisting of poorly stratified lenses and layers of stream-laid silt and silty sand, which are well sorted and can contain up to 30% clay. These swale and slough deposits are locally perennially frozen with moderate to high ice content. Analysis of a surface soil from the study area (MI 15) yielded a particle distribution consistent with silty sand containing gravel (Clausen et al. 2013a).

4.3 Contaminant distribution

4.3.1 Kimama Training Site

The munitions constituents (lead and antimony) were detected at levels during an SI, warranting further investigation (FPM 2009). Grab samples were collected in 2008. A subsequent RI/FS of the small-arms range at Training Area 3 yielded no metal results exceeding USEPA screening criteria (FPM 2010). However, only five composite samples were collected,

consisting of seven increments each from a radius of 0.3 m (FPM 2010). The composite sampling locations were selected at random; none of the samples were collected from the berm face, where the highest metal concentrations are expected. As discussed in Clausen et al. (2013a) Section 5.5.1, the ISM lead sample results collected from the berm face as part of our demonstration were slightly less than 300 mg/kg; the grab samples had a mean slightly less than 500 mg/kg.

4.3.2 Fort Eustis

The site is managed under the MMRP and had recently undergone an SI completed in 2007 (URS 2007). The SI involved the collection of 24 shallow soil samples taken at depths of 0–12 in. in the berms and firing lanes of the 1000-in. Rifle Range. All but four of these samples indicated lead concentrations were higher than the base-wide background level (23 mg/kg). Three of the soil samples had lead concentrations higher than the recommended soil screening level for residential use (400 mg/kg)—two in the Northern Berm and one in the Central Berm remnant.

4.3.3 Fort Wainwright

The small-arms ranges at Fort Wainwright have not been previously sampled and analyzed for metals, principally because they are active ranges. Given the use of projectiles containing antimony, copper, lead, and zinc, the presence of these metals above background concentrations seemed likely.

5 Test Design

5.1 Conceptual experimental design

The experimental design for the demonstration was generally the same for all three sites and is described in Clausen et al. (2013a). The entire impact berm face for each site was considered the DU (Fig. 5). Both the Kimama TS and Fort Eustis sites had a single intact berm (Fig. 6 and 7). In contrast, the range at Fort Wainwright consisted of multiple (16) individual berms located at varying distances downrange from an individual firing point (Fig. 8). At Fort Wainwright samples were collected from all 16 berms at the 100-m downrange distance from the firing point to form the DU (Fig. 9). In the case of Fort Wainwright, the firing point was also designated a DU and was sampled using ISM to demonstrate that the methodology developed for metals was equally applicable to energetics (i.e., a single sample could be collected for both analyses). As the firing points at Kimama TS and Fort Eustis were no longer identifiable, samples were not collected from these areas during the demonstration (Clausen et al. 2013a).

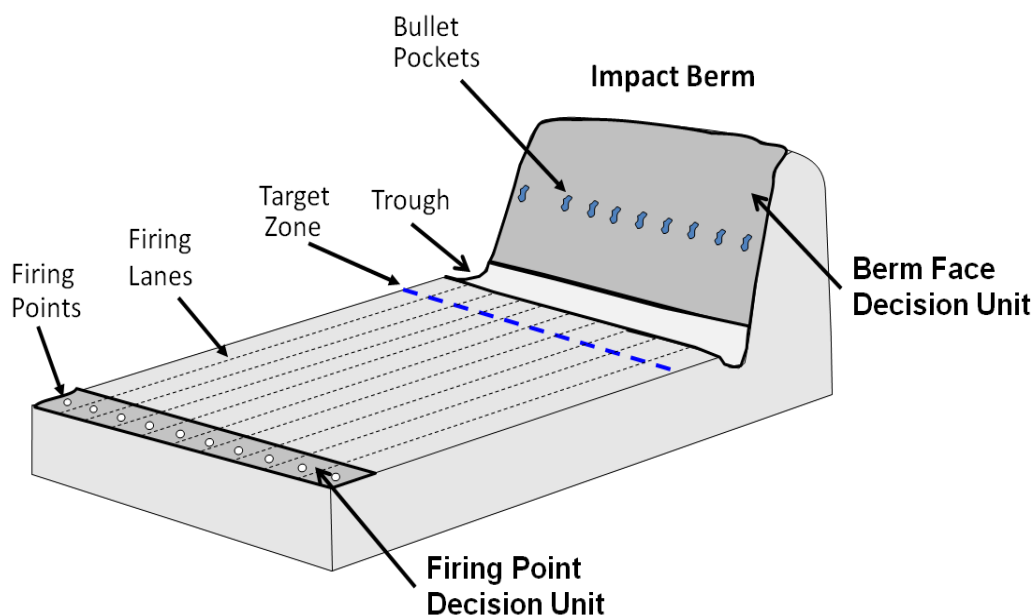


Figure 5. Generic example of a typical small-arms firing range.



Figure 6. The northernmost small-arms range berm face located in Training Area 3 of Kimama TS.



Figure 7. The 1000-in. small-arms range berm face at Fort Eustis.



Figure 8. The small-arms firing Range 16 Record berms at Fort Wainwright.

Using the random systematic approach, the team collected 15 replicate ISM samples from both the berm face DU and the firing point DU at Fort Wainwright to yield 30 individual samples (Fig. 1). Each replicate ISM sample consisted of approximately 100 increments. In the case of Fort Wainwright, because a single contiguous berm was not present, 6 increments were collected from each of the 16 individual bermlets to yield a single sample representative of the berms on the entire range at the 50-m distance (Figs. 8 and 9). In addition, within each berm face DU, a grid was created and individual grab samples were collected. For each of the 16 individual berms, three discrete samples were also collected using the pattern shown in Figure 10. Further, the berms were subdivided into 3 SUs with samples pooled from the upper left, upper right, and lower bottom to look at the distribution of metal within the berm. Finally, Berm 11 was subdivided into two SUs (left and right) to determine whether the results for the right and left sides of the berm were similar.

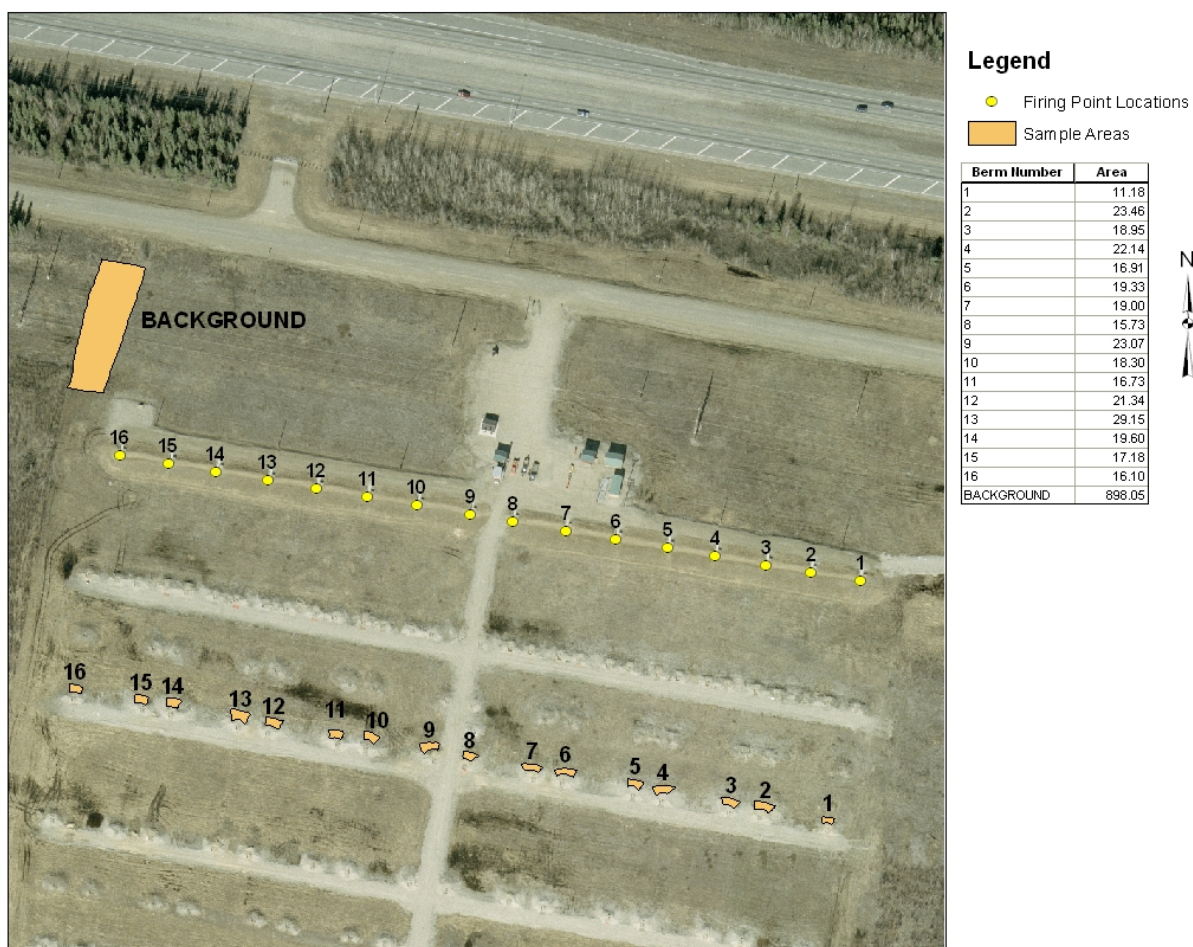


Figure 9. Location of berms sampled using ISM and grab techniques at the Range 16 Record Range at Fort Wainwright.

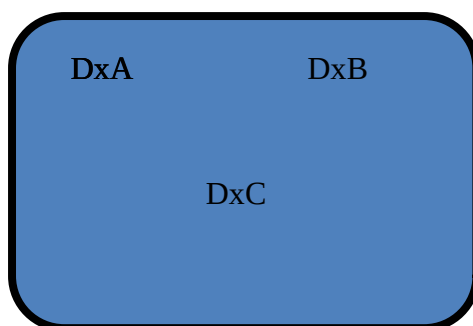


Figure 10. Grab sample grid layout for Fort Wainwright.

The grid sampling pattern used at the Fort Eustis and Kimama TS is shown in Figures 11 and 12, respectively. Two increments and two grab samples were collected per grid at Fort Eustis (Fig. 13). The berm at Fort

Eustis is a single DU that was divided into three SUs. Previous sampling (URS 2007) indicated an area of elevated metal content on the northern end of the berm (Fig. 13). Because the area was highly disturbed, no background samples were collected at Fort Eustis. At the Kimama TS, a small-arms range berm and background DU was established (Fig. 14). Grab samples were also collected within the small-arms range berm DU for comparison with the ISM samples.

1	2	3	4	5	6	7	8	9	10
11	12	13	14	15	16	17	18	19	20
21	22	23	24	25	26	27	28	29	30

Figure 11. Grab sample grid layout for Kimama TS berm face.

11	1	9	8	7	6	5	4	3	2	1
22	21	20	19	18	17	16	15	14	13	12
33	32	31	30	29	28	27	26	25	24	23

Figure 12. Grab sample grid layout for Fort Eustis berm face.

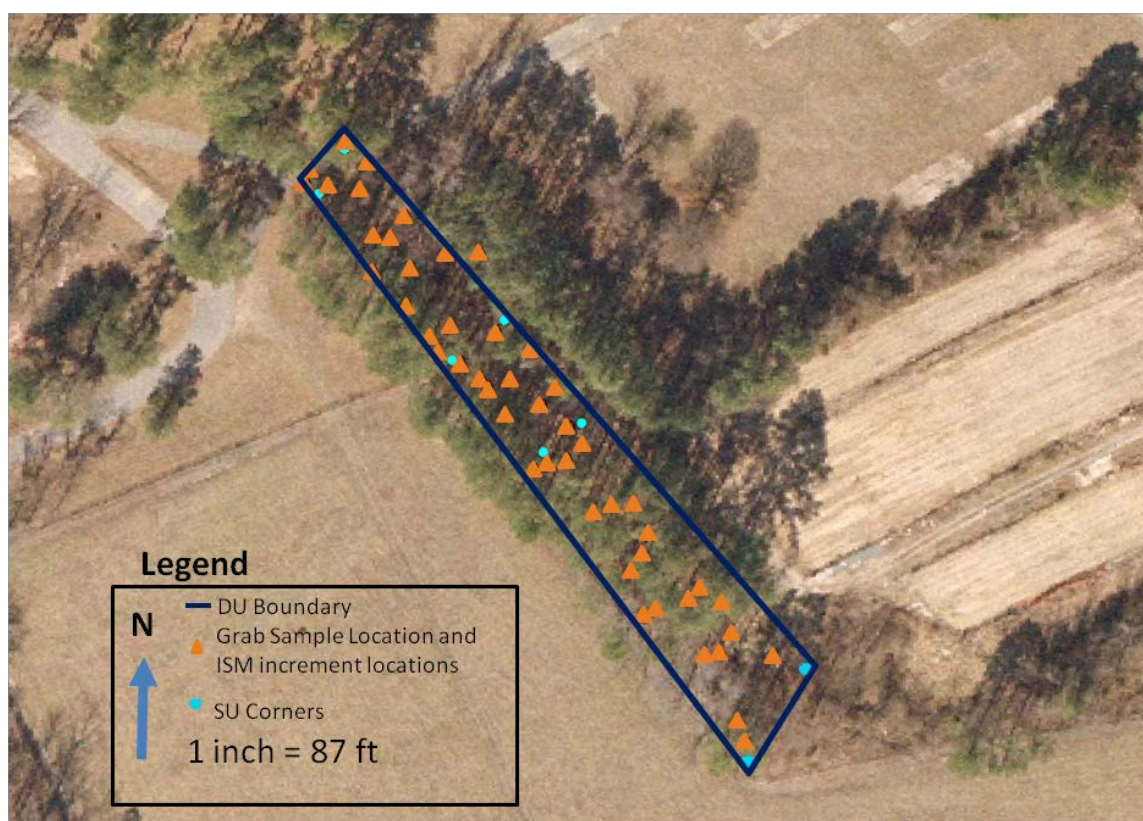


Figure 13. Aerial view of grab sample grid locations (orange triangles) and DU boundaries (blue circles) for the 1000-in. Range berm face at Fort Eustis.

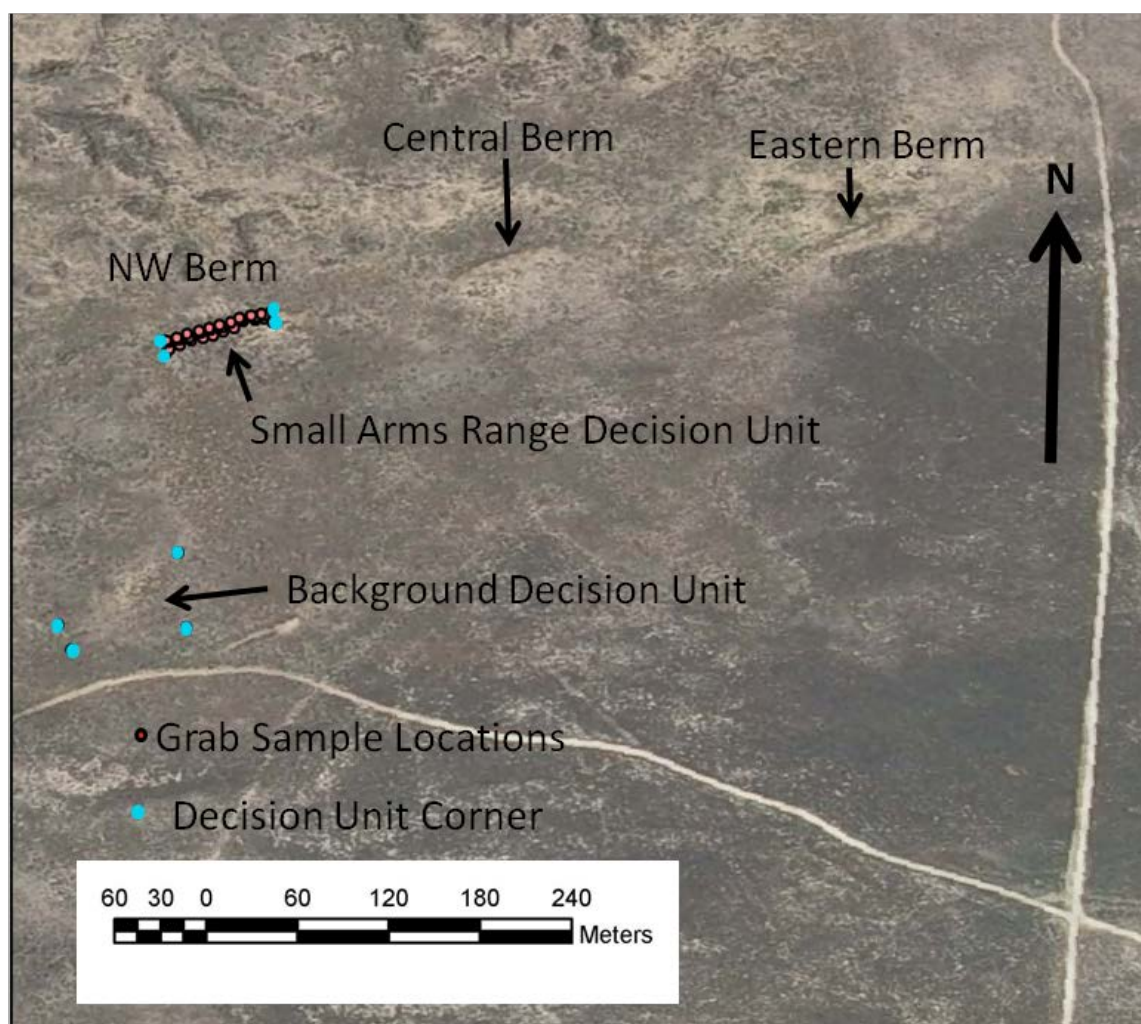


Figure 14. Aerial view of grab sample grid locations (orange circles) and DU boundaries (blue circles) for the NW berm face at Kimama TS.

5.2 Baseline characterization

Background surface soil samples were collected at Kimama TS and Fort Wainwright and analyzed for metals for comparison with samples obtained from the berm face and firing point DUs. A background sample was not collected from Fort Eustis because there did not appear to be any undisturbed soil locations in the vicinity of the range.

At the Kimama TS, the background samples did not have detectable concentrations of antimony (and thorium) (Clausen et al. 2013a). The concentration of copper ranged from 5.7 to 12.5 mg/kg with a mean of 8.0 mg/kg. Lead concentrations ranged from 5.2 to 1.2 mg/kg with a mean of 7.6 mg/kg.

At Fort Wainwright, the background samples did not have detectable concentrations of antimony (as well as silver, beryllium, cadmium, and thorium) (Clausen et al. 2013a). The concentration of copper ranged from 16.7 to 28.8 mg/kg with a mean of 26.8 mg/kg. Lead concentrations ranged from 8.1 to 136 mg/kg with a mean of 63.7 mg/kg.

In addition to the metal content of the background samples, several additional physical and chemical characteristics for the native soils were determined, including grain size, Total Organic Carbon (TOC), Cation Exchange Capacity (CEC), and soil pH (Clausen et al. 2013a).

5.3 Treatability or laboratory study results

The results from earlier laboratory studies under Task 1 of this project were presented in Clausen et al. (2012b) Section 2.2 *Technology Development*. No new treatability or laboratory studies were conducted for this demonstration.

5.4 Field testing

There are basically four phases to ISM: (1) Project Planning, (2) Mobilization, (3) Surveying/Sampling, and (4) Demobilization (Table 6). A draft demonstration report was submitted to ESTCP in June 2013 (Clausen et al. 2013a) followed by a protocol for ISM implementation at small-arms ranges (Clausen et al. 2013b). The Project Planning phase involves developing the conceptual site model, determining the study objectives, identifying the data needs, establishing the DU, and defining the depth and number of increments per ISM sample. Mobilization involves gathering the field equipment together and traveling to the site. The third phase involves demarcating the DU in the field, surveying the DU boundary or a corner of the DU, and sampling. The first three steps are identical to current conventional sampling methods. Sampling involves collecting conventional grab samples from within the DU and collecting ISM samples. Demobilization involves packing up the sampling equipment, shipping samples back to the laboratory, and traveling. Again, this is no different than the current conventional method. No investigative derived waste is created nor is equipment left at the sites.

Table 6. Gantt chart for field demonstration activities.

	September 2011			October 2011				November 2011				December 2011				
Activity	12-16	19-22	26-30	3-7	11-14	17-21	24-28	31-4	6-10	14-18	21-23	28-2	5-9	12-16	19-23	27-30
Fort Wainwright																
Project Planning																
Mobilization																
Surveying/Sampling																
Demobilization																
Kimama Training Site																
Project Planning																
Mobilization																
Surveying/Sampling																
Demobilization																
Fort Eustis																
Project Planning																
Mobilization																
Surveying/Sampling																
Demobilization																

Each ISM sample consisted of collecting approximately 100 increments within a DU using systematic random sampling (Fig. 1). A DU was established at the firing point (Fort Wainwright) and berm face of each small-arms range. Firing points could not be established for the Fort Eustis or Kimama TS (Clausen et al. 2013a). The firing point DU at Fort Wainwright extended 5 m behind and 5 m in front of the firing point and encompassed all firing lanes at the small-arms range. The berm face DUs encompassed an area including all firing lanes and from the base of the berm face to the top. In addition, at Fort Eustis, three SUs within the DU were established for comparison with previous contractor-conducted SI studies. At Fort Wainwright, a continuous berm is not present; so the 16 individual berms located at 100 m down range from the firing point were considered a contiguous berm. The area sampled represented approximately 0.5 acres.

Within each DU, approximately 100 evenly spaced increments were collected to form an individual ISM surface soil sample. The CMIST (Walsh 2009) was used to extract cylindrical soil cores referred to as “increments.” The coring bit used had a diameter of 2 cm. The sampling depth used at all 3 sites was 5 cm, which yielded an ISM sample of approximately 1–2 kg. The required mass is a function of the soil volume of each core (the sampling depth and core diameter), the number of cores and increments collected, and the mean soil density. Although, the typical approach is to collect three replicate ISM samples for each DU to assess uncertainty, this demonstration collected 15 replicate samples to facilitate statistical comparisons between the ISM and grab sample results.

Using a grid-node approach, 48 grab samples were collected from Fort Wainwright, 33 from Fort Eustis, and 30 from Kimama TS (Clausen et al. 2013a). From within each node, a single increment was collected using the CMIST sampler and was placed in individual amber 4 oz containers, yielding a sample mass of approximately 0.2 kg. Typically, grab samples would be collected with a metal scoop. However, for direct comparison between the ISM and grab samples, the same sample device, CMIST, was used. For a typical small-arms range, usually a half-dozen to a dozen grab samples would be obtained. However, to facilitate statistical analysis of the data and comparisons with the ISM data, more samples were collected (greater than 30) than is typical. The grid-node layout encompassed the same area as the DUs where the ISM samples were collected.

5.5 Sampling methods

The demonstration included collecting 63 ISM surface soil samples and 48 conventional grab samples at Fort Wainwright, 18 ISM and 30 grab samples from Kimama TS, and 27 ISM and 33 grab samples at Fort Eustis (Clausen et al. 2013a). Conventional grab and ISM samples were collected from three small-arms range berm DUs at each of the military installations. One firing point DU sample was collected at Fort Wainwright to assess the suitability of collecting and processing a single sample for both metal and energetic analysis. A boundary for each DU was determined in the field largely based on the observable extent of the impact berm or firing point.

5.5.1 Grab samples

Grab surface soil samples were collected following the conventional grid node approach from within each DU. All grab sample locations were individually surveyed. Instead of using a metal scoop, the CMIST was used to collect the grab sample from the center of the grid. The 2-cm-diameter corer was used with a 5-cm-deep sample collected. The grab samples were processed following the USEPA Method 5030B, which basically involved scooping a 1-g sample out from the top of the sample jar and then performing the standard digestion. Metals analysis followed USEPA Method 6010.

5.5.2 Incremental sampling methodology samples

Collection of ISM surface soil samples followed the methodology outlined in Clausen et al. (2013a) Section 2.1 *Technology Descriptions*; and the steps, including sample preparation, are summarized in the present report in Table 1 with the differences between ISM and conventional grab approaches shown in Table 2. The ISM samples were collected in plastic bags by combining 100 increments for a total mass of 1–2 kg. All samples for this demonstration were processed, extracted, and digested at CRREL. Samples were analyzed following EPA Method 8330B for Nitroglycerin (NG); 2,4-dinitrotoluene; and 2,6-dinitrotoluene or a modified Method 3050B/6010B for metals. Energetic analysis was performed at CRREL and all metals analysis and additional analytes were analyzed at the Environmental Laboratory (EL).

Except for at Fort Eustis, we selected a background location close to the small-arms ranges at each installation but upwind of the prevailing wind direction. Triplicate ($n = 3$) 100-increment samples were collected from these background DUs, which covered an area of 0.5 acres at each site.

At Fort Wainwright, 15 replicate ISM samples were collected from the firing point and were analyzed for energetics and metals. The intent was to demonstrate the metals ISM process would work with soils containing energetics. For each berm DU, $n = 15$ ISM field replicates were collected to evaluate total precision.

Table 7. Comparison of Grab versus ISM for this demonstration.

Activity	Conventional Grab Sampling	Incremental Sampling Method
Surveying	Each individual sample location was flagged and surveyed.	The DU corners were determined and demarcated with flagging as were lane boundaries for sample collection. The four corners of the DU were surveyed.
Soil Sampling	Not explicitly addressed in Method 3050B. Grab samples collected with CMIST from biased locations. Typically, about 200 g of soil was collected in 4-oz, wide-mouth, amber, screw-top jars.	A 100-increment sample was collected randomly over the entire DU (e.g., using a systematic sampling) using CMIST. Typically, 1–2 kg of soil was collected in clean, large (e.g., 15 × 15 in., 6 mm thick) polyethylene plastic bags sealed with Ty-wraps.
Sample Drying	Not performed.	Samples were air-dried at room temperature by spreading them onto trays to form a relatively thin, uniform slab.
Sieving	Not performed.	Samples were passed through a #10 (2-mm) sieve. Both size fractions were weighed and the < 2 mm fraction was additionally processed.
Milling	Not performed, although disaggregation with a mortar and pestle is sometimes performed.	Samples were milled using a puck mill for 5 x 60 s.
Laboratory Subsampling	A single aliquot was scooped from the top of the container for digestion and analyses.	After milling, the soil was spread onto a large tray to form a thin slab of material of uniform thickness. With a flat spatula, at least 20 small aliquots “increments” were randomly collected over the entire slab to prepare a subsample for digestion and analysis.
Subsample Mass	1-g wet weight.	2-g dry weight.
Analysis	EPA Method 6010.	EPA Method 6010 (EPA Method 8330B for Firing Point sample from Fort Wainwright).

5.6 Sampling results

5.6.1 Kimama Training Site

At the Kimama TS, copper and lead were detected in ISM surface soil samples from the small-arms Northern Berm in Training Area 3 at levels higher than the background sample. The mean copper and lead levels in the background ISM sample were 7.96 and 7.61 mg/kg and 35.3 and 292 mg/kg in the berm sample, respectively (Clausen et al. 2013a). In contrast, the mean copper and lead results for the grab samples were 23.0 and 493 mg/kg, respectively (Fig. 15) indicating the grab sample mean for lead (493 mg/kg) was nearly double that of the ISM mean (292 mg/kg). Note that the ISM mean and median for Pb were nearly the same (292 versus 287 mg/kg) whereas the mean and median lead grab sample values (493 versus 73.5 mg/kg) differed by nearly a factor of seven. A total of 15 replicate ISM and 30 grab samples were collected, and the calculated RSD for lead was significantly higher for the grab samples (334%) as compared to the ISM samples (18%). Figure 15 depicts the individual grab results for the berm face. The western side of the Northern Berm had higher copper, lead, and zinc concentrations than the eastern side of the berm. Clausen et al. (2013a) provides individual sample results for all ISM and grab samples, including all metal analytes, along with a more detailed analysis of the data.

Lead Results for Discrete Samples (mg/kg)										Legend	
75	1240	9060	1050	523	278	103	60	85	45	<50	
39	287	97	325	556	278	72	31	24	14	51-299	
11	19	30	40	142	198	28	49	25	13	>300	
ISM Mean = 292, ISM Median = 287, Discrete Mean = 493, Discrete Median = 73.5											
Copper Results for Discrete Samples (mg/kg)											
20	57	74	32	41	42	26	15	18	18	<20	
14	24	21	19	39	42	14	16	13	13	21-39	
10	10	12	14	14	22	13	18	11	12	<40	
ISM Mean = 35., ISM Median = 29.2, Discrete Mean = 23.0, Discrete Median = 18.1											
Antimony Results for Discrete Samples (mg/kg)											
<2.00	<2.00	<2.00	<2.00	<2.00	<2.00	<2.00	<2.00	2.2	<2.00	<0.02	
<2.00	<2.00	<2.00	<2.00	<2.00	<2.00	<2.00	<2.00	<2.00	<2.00	>0.021	
<2.00	<2.00	<2.00	<2.00	<2.00	<2.00	<2.00	<2.00	<2.00	<2.00		
ISM Mean = <2.00, ISM Median = < 2.00, Discrete Mean = 19.5, Discrete Median = 3.01											
Zinc Results for Discrete Samples (mg/kg)											
53	53	56	46	55	51	48	41	47	41	<25	
54	52	45	50	42	51	22	43	40	39	26-49	
45	44	42	46	44	51	44	42	37	40	>50	
ISM Mean = 30.0, ISM Median = 28.9, Discrete Mean = 45.4, Discrete Median = 44.9											

Figure 15. Grab surface soil results for lead, copper, antimony, and zinc (mg/kg) from the Kimama Training Site small-arms range berm face with Incremental Sampling Methodology comparisons.

5.6.2 Fort Eustis

At the 1000-in. Firing Range on Fort Eustis, copper and lead levels were elevated in grab and ISM surface soil samples for the entire berm face (Clausen et al. 2013a). The mean copper and lead grab sample results were 43.3 and 434 mg/kg, respectively, whereas the mean ISM results were 51.2 and 496 mg/kg, respectively. Unlike the Kimama TS results, the mean values for the grab and ISM samples were similar for both copper and lead. Although the ISM mean and median for lead values were nearly the same (496 versus 509 mg/kg), the mean and median lead grab sample values (434 versus 94.3 mg/kg) differed by nearly a factor of five, indicating the grab sample results are highly skewed. A total of three replicate ISM and 33 grab samples were collected, and the calculated RSD for copper and lead was significantly higher for the grab samples (298% and 350%, respectively) as compared to the ISM samples (104% and 72%), indicating greater precision with the ISM versus the grab samples. Although, the performance criteria of less than 30% RSD was not met with the ISM samples, suggesting an insufficient sample mass resulting from an insufficient number of increments per sample, an insufficient mass per increment collected, or an insufficient mass for the digestion aliquot.

Because previous SI data indicated most of the copper and lead was concentrated on the right side of the berm (URS 2007), a focused sampling effort was conducted in this area as well. A total of 12 grab and 15 ISM samples were collected from the first four grids (northern side of Fig. 13). The mean grab sample copper and lead results were 93.5 and 1002 mg/kg, respectively (Clausen et al. 2013a). The mean ISM copper and lead results were 114 and 932 mg/kg, respectively. Similar to the observation for the full berm, the ISM mean and median lead values were nearly the same at 932 and 934 mg/kg, respectively. In contrast, the mean and median lead values for the grab samples were significantly different at 1002 and 212 mg/kg, respectively, indicating a highly skewed dataset. An assessment of the precision of results indicated that the grab samples ($n = 12$) had RSDs of 224%, 241%, and 166% for copper, lead, and zinc, respectively. In contrast, the RSD for the ISM data ($n = 14$) yielded values of 44%, 30%, and 4% for copper, lead, and zinc, respectively. Clausen et al. (2013a) provides individual sample results for all ISM and grab samples, including all metal analytes. Copper did not meet the performance objective of less than 30% RSD, suggesting a longer milling interval or larger digestion aliquot mass may be necessary for improved sample precision.

These results indicate there is a large long-range spatial heterogeneity of metallic residues. The results from the three SUs indicate most of the metal contamination is present in the northern end of the berm, an observation that would likely not have been apparent (e.g., without historical knowledge) if a small number of grab samples (e.g., $n = 10$) were collected in a judgmental manner to characterize the berm. The long-range spatial heterogeneity and the large variability (e.g., RSDs) of grab sample results suggest that the mean concentration of the entire berm will likely not be accurately estimated without collecting a large number of grab samples over the entire berm (e.g., using systematic random sampling).

5.6.3 Fort Wainwright

At the Range 16 Record Range at Fort Wainwright, copper and lead levels in the background surface soil samples appeared to be at similar levels as the ISM entire berm sample, suggesting that the background location selected behind the firing point has been anthropogenically impacted. The mean copper and lead levels in the background ISM sample were 38.2 and 416 mg/kg, respectively (Clausen et al. 2013a). In contrast, the mean copper and lead levels in the ISM firing point sample were 135 and 50.5 mg/kg, respectively, and at the berm, 92 and 453 mg/kg, respectively. In addition to the metals, NG was detected at the firing points with a mean concentration of 335 mg/kg. The RSDs for metals and NG for replicate firing point incremental samples met the target value of 30% for total precision. The measurement performance objective of the incremental samples that were collected from the berm was also met for lead and zinc. However, the RSD for copper was 64%. It has been our observation that copper has a tendency to plate out in the puck mill. It is possible that a longer milling interval, greater than 300 s, would result in better precision.

The mean copper, lead, and antimony grab sample results for the entire berm were 81.0, 432, and 14.0 mg/kg, respectively (Clausen et al. 2013a). The lead grab samples' mean (432 mg/kg) for the entire berm was similar to the ISM mean (453 mg/kg). The entire berm ISM mean and median for lead was nearly the same (453 versus 468 mg/kg) whereas the mean and median lead grab sample values (432 versus 85.7 mg/kg) differed by nearly a factor of five, indicating the grab sample results were highly skewed.

A total of 15 replicate ISM and 48 grab samples were collected. The calculated grab sample RSDs for copper, lead, and zinc were 218%, 228%, and 42%, respectively (Clausen et al. 2013a). In contrast, the calculated copper,

lead, and zinc RSDs for the ISM samples were 64%, 24%, and 12%, respectively. Clausen et al. (2013a) provides individual sample results for all ISM and grab samples, including all metal analytes.

6 Performance Assessment

6.1 Sample reproducibility with Incremental Sample Methodology

A quantitative performance objective for ISM was to obtain reproducible sample results through collection and analysis of replicate field samples. Two different statistical success criteria were developed for this evaluation. The first approach involved collecting replicate incremental samples and grab samples in the same DU and subsequently using statistical hypothesis tests to determine whether the ISM approach resulted in significantly smaller variances. The smaller ISM variances were compared with the grab variances to determine whether they were significantly different with at least 95% confidence (Fig. 16). The evaluation indicated that the ISM approach generally resulted in better measurement precision for all three of the demonstration sites; the ISM approach generally resulted in smaller variances though the target level of confidence (95%) was not met for all of the metals and DUs. The largest differences in the variance were observed for lead, which is usually the primary contaminant of concern for small-arms ranges. The large reductions for the lead variances is illustrated in Figure 16 for the natural-logarithm-transformed ISM ($n = 15$) replicates and the ($n = 48$) grab replicates from the Fort Wainwright DU. The Levene's test indicates the variance of the set of incremental samples (denoted as "ISM") is significantly smaller than variance of the set of grab samples (denoted by "Discrete") with well over 95% confidence (as the "P-value" is much smaller than 0.05).

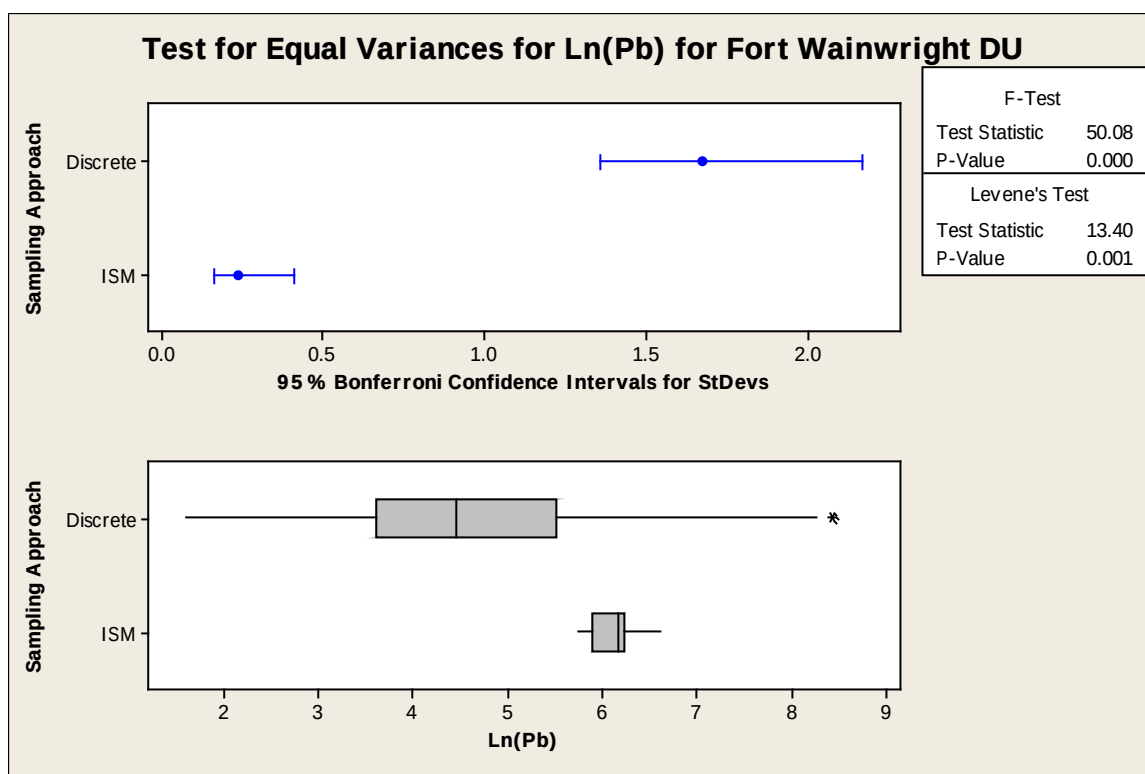


Figure 16. Analysis of variance for lead at Fort Wainwright.

The Levene's test was not able to detect significant differences between the grab and ISM variances for copper and zinc with at least 95% confidence, but the square ranks test for the variances detected significant differences with at least 95% confidence for lead and copper (the level of confidence was greater than 99% for lead, 99% for copper, but only 85% for zinc). Although the hypothesis tests did not detect significant differences between the variances at the target level of confidence, the ISM approach seems to result in better precision than conventional grab sampling. The standard deviation for the set of grabs (22 mg/kg) is over three times larger than the ISM standard deviation (7 mg/kg). As shown in the zinc box plot (Fig. 17), the grab replicates also exhibit greater skewness than ISM replicates. Values that exceeded the upper or lower quartiles by $1.5 \times \text{IQR}$ (Interquartile Range) or more were identified as "outliers," consistent with conventional terminology and plotting procedures for box plots.

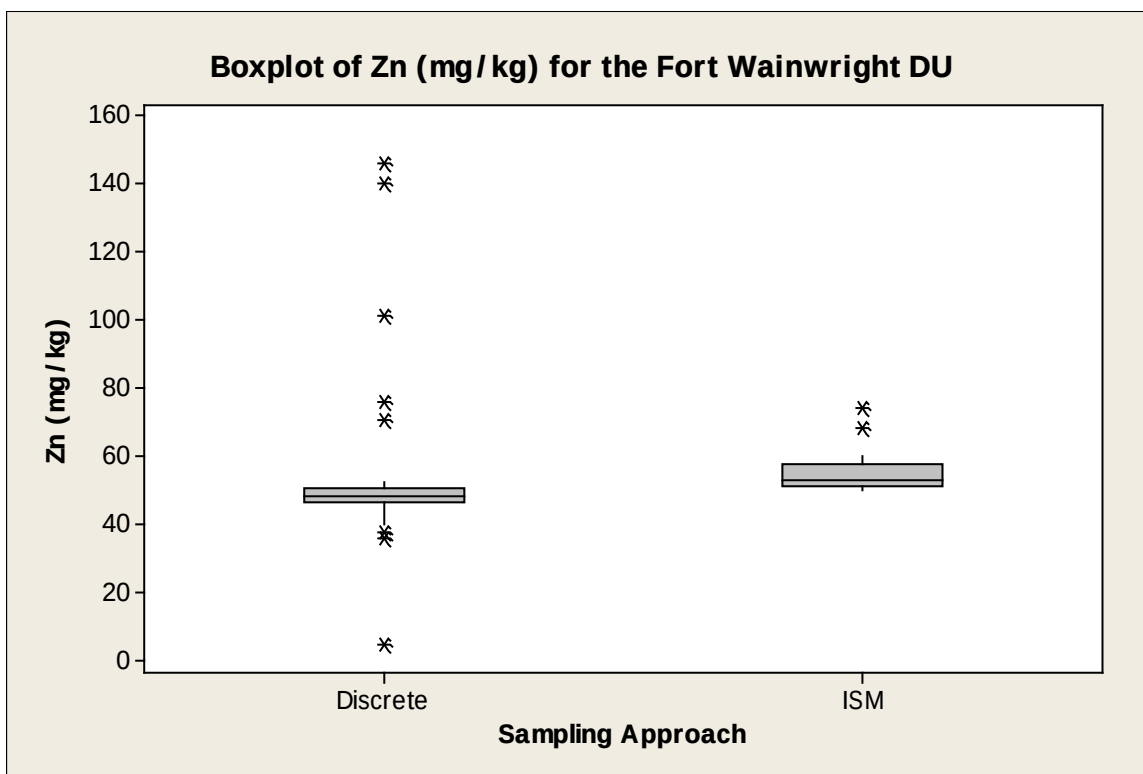


Figure 17. Analysis of variance for zinc at Fort Wainwright.

A second qualitative evaluation involved calculating the RSD for each population (Table 8). Again, for all three sites, the RSDs for the ISM samples were significantly lower than for the conventional grab samples with a few situations where the measurement performance objective of less than 30% RSD was not met for some metals with ISM. As shown at Fort Eustis and at Fort Wainwright, an ISM sample with a larger number of increments and a greater mass yielded lower RSDs (Table 8). It is believed in those situations where the performance metric of RSD less than 30% was not met that the target would have been achievable by resampling and collecting an ISM sample with a greater number of increments, by using a larger sampling tool to increase the recovered mass, by increasing the digestion mass, or by increasing the number of subsampling increments to build the digestion aliquot. This leads to the observation that when the degree of expected analyte heterogeneity is unknown, the sampler should err on the conservative side by collecting a sample from a DU with a larger number of increments or a greater mass. The other approach is to select a DU with a smaller area that will thus increase the increments per DU area or the mass of the sample per DU area. Overall, sample reproducibility precision was improved with the collection of ISM samples compared with conventional grab samples. Laboratory subsampling precision was previously

demonstrated during the technology development of this project (Clausen et al. 2012b, 2013a).

6.2 Bias evaluation

Clausen et al. (2013a) evaluated the method blanks and LCS for bias with none indicated. Glass blank samples analyzed as a control indicated some increase in aluminum, chromium, iron, and manganese as a result of milling with the puck mill although none of the analytes of interest (copper, lead, antimony, or zinc) increased significantly (Clausen et al. 2012b, 2013a), which is consistent with the findings of Felt et al. (2008) for reference soils studied. Therefore, there is no significant cross-contamination from the puck mill into the samples for the metals of interest (copper, lead, antimony, or zinc).

Table 8. Comparison of relative standard deviations for the analytes of interest (copper, lead, antimony, zinc) for ISM and conventional grab samples at three demonstration sites.

	Kimama TS	Fort Eustis ¹	Fort Wainwright ²
	Incremental Samples		
Mean Mass (g)	881	1112/1485	2403/1208/1237
Number of Samples	15	15/15	15/15/15
Copper RSD (%)	53	104/44	64/52/125
Lead RSD (%)	18	72/30	24/56/58
Antimony RSD (%)	ND	500/60	ND/ND/ND
Zinc RSD (%)	23	5/4	8/17/20
	Conventional Grab Samples		
Mean Mass (g)	100	78.8/69.6	159
Number of Samples	30	33/12	48
Copper RSD (%)	66	298/224	218
Lead RSD (%)	334	350/247	226
Antimony RSD (%)	172	219/295	97
Zinc RSD (%)	15	23/27	69

Green = < 30% RSD, Yellow = ~30% RSD, Red = > 30% RSD

RSD—relative standard deviation; ND—not detected

¹ Entire berm/right side of berm

² Entire berm/left side Berm 11/right side Berm 11

6.3 Performance comparison

As demonstrated in Clausen et al. (2012b), measurement of the improved accuracy as compared to the “true” mean is difficult to quantify because the true mean is typically unknown. Either knowing the total amount of metal introduced into the DU or digestion of all the soil in the DU are the only means to know the true metal concentration. Neither of these are practical approaches as the number of projectiles fired into a berm DU is typically unknown, and it is not practical to digest several hundred or thousand of kilograms of soil. The approach taken in Clausen et al. (2012b, 2013a) was to pool all of the ISM samples from a DU to calculate the overall mean, the assumption being this pooled mean is more representative of the true site condition. This assumption is not entirely unreasonable as the DUs studied typically had several dozen ISM samples collected with each ISM sample containing 50 to 100 increments. Thus, the total number of increments collected from a given DU was several hundred up to 1500 in one instance. Comparing the pooled ISM mean to the grab sample results indicated accuracy errors approaching 100% for individual grab samples. The only means to reduce the accuracy error with grab samples was to increase the number of grab samples collected.

As shown, in this demonstration (Table 8), total sample error or precision as measured qualitatively using the RSD is often several hundred percent for conventional grab samples whereas RSDs for ISM samples are typically less than 30% for soils containing metallic residues (Clausen et al. 2012b, 2013a).

Improved accuracy or precision may not be necessary when the soil metal concentration is well above or well below a regulatory action level or some other criteria for comparison. However, when the measured soil concentration is close to an actual level, the precision and accuracy of the measurements becomes increasingly important. As observed from this demonstration, in the case of lead, different decisions are likely to be made whether conventional grab or ISM samples are collected. Clausen et al. (2013a) provide an example for lead at the Fort Wainwright Record Range berm where different decisions could be made when relying on the grab sample data alone whereas a consistent decision would be made each time if the ISM data were used. Assuming a regulatory action level of 400 mg/kg for lead, 13% of the time the grab sample value would be less than 400 mg/kg; and 87% of the time it would be greater than 400 mg/kg. However, the ISM data clearly indicates the mean lead level for the DU is

above 400 mg/kg. Consequently, reliance on grab samples would result in a relatively high false negative rate (i.e., erroneously concluding contamination is less than the 400-mg decision limit for remedial action).

In terms of assessing performance by cost, each stage of the sampling activity and laboratory processing step was evaluated for the time to perform the task for both the ISM and conventional grab samples at each of the three sites. For the field sampling activities, seven variables were considered for surface soil samples:

1. Mobilization preparation time.
2. Shipping of field equipment to the site.
3. Surveying and flagging in the field.
4. Sampling.
5. Labeling of samples.
6. Demobilization.
7. Shipping of samples to the laboratory.

Mobilization, shipping equipment to the field, and demobilization costs are essentially the same. The remaining field activities take twice as long with ISM versus conventional grab sampling on a per sample basis. Laboratory preparation procedures assessed for the soil samples were (1) air drying, (2) sieving, (3) milling, and (4) subsampling to prepare the digestion aliquot. With the conventional grab sampling approach, the collection of a 0.5- to 2-g aliquot from the top of the jar takes minimal time, thus there are no sample preparation costs. Based on our experience at CRREL, the laboratory preparation time per sample is roughly 30 minutes of labor. There is essentially no difference between analyses for a conventional grab sample versus ISM. Table 9 shows the breakdown of costs by field sampling, sample preparation, and analysis activity based on a field technician and laboratory technician at a rate of \$50/hr. On a per sample basis, the cost of ISM is approximately 55% to 65% higher than conventional grab sampling. However, a cost savings is apparent when one considers the number of samples. The more grab samples needed to characterize the DU (7 versus 15 grab samples), the more favorable the cost comparison with ISM (Table 9). For a typical small-arms range DU, three replicate ISM samples would be commonly collected versus 7 to 15 conventional grab samples for the same DU. Thus, the total ISM project cost is 1 to 3 times (5% to 65%) lower as compared to the conventional grab sampling total project cost. Therefore, ISM met our performance criteria of at least a 20%

reduction in sample cost even when the quality of the grabs results (e.g., the poorer measurement precision) relative to set of incremental samples is not taken into account.

Table 9. Comparison of labor costs between ISM and conventional grab sampling on a per sample and total cost basis based on demonstrations at Kimama Training Site, Fort Eustis, and Fort Wainwright.

Activity	Per Sample Costs (\$)		Total Project Costs (\$)		Total Project Costs (\$)	
	ISM	Grab	ISM ¹	Grab ²	ISM ¹	Grab ³
Field Sampling	35–50	10–15	105–150	70–105	105–150	150–225
Laboratory Preparation	40–60	0–10	120–180	0–70	120–180	0–150
Analysis	225–275	125–135	675–825	875–945	675–825	1875–2025
Total	300–385	135–160	900–1155	945–1120	900–1155	2025–2400

¹ Based on collection of 3 replicates.

² Based on the collection of 7 grab samples.

³ Based on the collection of 15 grab samples.

7 Cost Assessment

The costs associated with field-sampling activities include travel, related lodging and meals, labor, and the shipment of samples off site. Unique to the costs associated with sampling activities on military training ranges is the need to acquire the services of explosive ordnance disposal (EOD) personnel or UXO technicians. However, the expenses associated with gaining site access, engaging EOD support, travel, and labor are anticipated to be equivalent for grab and ISM. Additionally, in the case of small-arms ranges, an EOD escort is not needed because UXO presence is unlikely. The major cost differences between ISM and conventional grab sampling arise predominately from the ISM requirement of handling and processing of larger environmental samples. However, this cost increase is greatly offset by the need for fewer ISM samples than grab samples to adequately characterize a DU. The cost differential between conventional grab samples and ISM is quantifiable. However, the cost of making a wrong decision is not easily quantifiable. The potential cost of implementing a remedial remedy when it is not necessary could be quite large, ranging from tens of thousands to tens of millions of dollars. The potential remedial action of a false positive finding is not likely to be known unless both conventional grab and ISM samples are collected. However, implementation of ISM will result in a lower false positive rate, which is associated with fewer unnecessary remedial actions. Conversely, the ISM also has a more reliable detection rate, thus avoiding false negatives.

7.1 Cost model

To aid in our cost analysis, the labor hours in all phases or actual costs of the field demonstration (site preparation, locating the sample with GPS, sample labeling, sample collection, shipment of samples, etc.) were collected and tracked in an Excel spreadsheet (Clausen et al. 2013a) (Table 10). In addition, the cost or labor hours to process the samples and to analyze them in the laboratory were tracked. Labor categories and labor rates were ascertained so that labor hours could be converted to actual costs. Because the actual number of samples collected for this demonstration was greater than for a typical project, the following approach was taken. Three ISM replicate samples from a single DU were compared against a typical number of conventional grab samples for the same DU. Because the number of grab samples collected varies by objective, analyte of con-

cern, desires of the interested stakeholders, etc., two scenarios were considered: 7 and 15 grab samples. This range was based on discussions with individuals involved with using conventional grab sampling techniques at MMRP sites.

Table 10. Comparison of labor hours¹ or costs by cost element between ISM and conventional grab sampling on a per sample and total cost basis based on demonstrations at Kimama Training Site, Fort Eustis, and Fort Wainwright.

Stage	Activity	Fort Wainwright				Fort Eustis				Kimama Training Site			
		ISM	Grab	ISM	Grab	ISM	Grab	ISM	Grab	ISM	Grab	ISM	Grab
		Per Sample		Total Project		Per Sample		Total Project		Per Sample		Total Project	
Mobilization	Preparation	=	=	=	=	=	=	=	=	=	=	=	=
	Expendables	\$8.62	\$1.29	\$129.35	\$61.92	\$8.62	\$0.89	\$129.35	\$42.57	\$8.62	\$1.29	\$129.35	\$38.70
	Shipping Field Equipment	=	=	=	=	=	=	=	=	=	=	=	=
Field	Surveying/ Flagging	1	1	16	53	2	1	30	45	1	2	8	54
	Sampling	12	2	90	37	20	4	150	60	20	3	150	45
	Decontamination	0	2	0	96	0	5	0	165	0	5	0	150
	Labeling	2	2	35	113	1	1	15	33	1	1	18	36
	Demobilization	=	=	=	=	=	=	=	=	=	=	=	=
	Shipping Samples	\$25.27	\$2.46	\$379.00	\$118.00	\$7.54	\$2.59	\$113.14	\$85.47	\$22.27	\$3.67	\$334.00	\$110.00
	Shipping Field Equipment	=	=	=	=	=	=	=	=	=	=	=	=
Laboratory	Air Drying Prep	2	0	30	0	2	0	30	0	2	0	30	0
	Sieving	2	0	30	0	2	0	30	0	2	0	30	0
	Milling	5	0	75	0	5	0	75	0	5	0	75	0
	Cleaning Milling Equipment	10	0	150	0	10	0	150	0	10	0	150	0
	Sub-Sampling	10	0	150	0	10	0	150	0	10	0	150	0
	QA/QC	\$26.67	\$33.33	\$400.00	\$1,600.00	\$26.67	\$26.67	\$400.00	\$800.00	\$26.67	\$26.67	\$400.00	\$800.00
	Laboratory Supplies	<	>	<	>	<	>	<	>	<	>	<	>
	Analysis	\$225.00	\$100.00	\$3,375.00	\$4,800.00	\$225.00	\$100.00	\$3,375.00	\$3,300.00	\$225.00	\$100.00	\$3,375.00	\$3,000.00

¹ Units are hours unless denoted by \$.

= indicates equivalent cost; > or < denotes a minor lower or greater cost not tracked.

We also calculated the costs based on our labor at CRREL and EL for sample preparation and analysis and obtained actual costs from several commercial environmental laboratories (Table 11). A labor rate of \$50/hr was used for converting labor into dollars. A typical soil digestion and target analyte list (TAL) analysis of 13–18 metals by a commercial environmental laboratory costs \$100/sample. If ISM sample preparation (air drying, sieving, subsampling) including milling is required, this adds approximately \$125/sample, based on discussions with several commercial environmental laboratories. Use of ISM sample preparation without milling adds approximately \$75/sample.

Table 11. Comparison of costs for ISM and conventional grab sampling on a per sample and total cost basis based on demonstrations at Kimama Training Site, Fort Eustis, and Fort Wainwright.

Stage	Activity	Fort Wainwright				Fort Eustis				Kimama Training Site			
		ISM	Grab	ISM	Grab	ISM	Grab	ISM	Grab	ISM	Grab	ISM	Grab
		Per Sample		Total Project		Per Sample		Total Project		Per Sample		Total Project	
Mobilization	Preparation	=	=	=	=	=	=	=	=	=	=	=	=
	Expendables	\$9	\$1	\$129	\$62	\$9	\$1	\$129	\$43	\$9	\$1	\$129	\$39
	Shipping Field Equipment	=	=	=	=	=	=	=	=	=	=	=	=
Field	Surveying/ Flagging	\$1	\$0.92	\$13	\$44	\$2	\$1	\$30	\$45	\$0	\$2	\$7	\$45
	Sampling	\$10	\$1	\$75	\$31	\$17	\$4	\$150	\$60	\$17	\$3	\$125	\$38
	Decontamination	\$0	\$2	\$0	\$80	\$0	\$5	\$0	\$165	\$0	\$4	\$0	\$125
	Labeling	\$2	\$2	\$29	\$94	\$1	\$1	\$15	\$33	\$1	\$1	\$15	\$30
	Demobilization	=	=	=	=	=	=	=	=	=	=	=	=
	Shipping Samples	\$25	\$2	\$379	\$118	\$8	\$3	\$113	\$85	\$22	\$4	\$334	\$110
	Shipping Field Equipment	=	=	=	=	=	=	=	=	=	=	=	=
Laboratory	Air Drying Prep	\$2	\$0	\$25	\$0	\$2	\$0	\$25	\$0	\$2	\$0	\$25	\$0
	Sieving	\$2	\$0	\$25	\$0	\$2	\$0	\$25	\$0	\$2	\$0	\$25	\$0
	Milling	\$4	\$0	\$63	\$0	\$4	\$0	\$62	\$0	\$4	\$0	\$63	\$0
	Cleaning Milling Equipment	\$8	\$0	\$125	\$0	\$8	\$0	\$125	\$0	\$8	\$0	\$125	\$0
	Sub-Sampling	\$8	\$0	\$125	\$0	\$8	\$0	\$125	\$0	\$8	\$0	\$125	\$0
	QA/QC	\$27	\$33	\$400	\$1,600	\$27	\$27	\$400	\$800	\$27	\$27	\$400	\$800
	Laboratory Supplies	<	>	<	>	<	>	<	>	<	>	<	>
	Analysis	\$225	\$100	\$3,375	\$4,800	\$225	\$100	\$3,375	\$3,300	\$225	\$100	\$3,375	\$3,000
	Total	\$323	\$143	\$4,763	\$6,829	\$311	\$141	\$4,574	\$4,531	\$325	\$141	\$4,748	\$4,186

From a statistical basis (e.g., the Central Limit Theorem), n incremental samples, each prepared from k increments, will produce data that is roughly of similar quality for estimating the population mean of the DU as $n \times k$ grab samples. On the basis of this simplistic theoretical model (which likely over estimates the quality of the ISM results to a degree in practice), the cost of sampling n ISM samples of k incremental in the field will be no greater than the $n \times k$ grab samples. The cost of the former would be expected to be less than that of the latter because ISM would entail the use of fewer sample containers and less labeling and documentation. Even if comparable field sampling costs are estimated for the grab and ISM samples, the ISM approach will result in a cost savings owing to the smaller number of laboratory analyses required.

Similarly, if each grab sample is assumed to weigh on average 150 g and each ISM weighs 1500 g but $n \times k$ grab samples produce the same quality of data as n ISM, it follows that the total weight of the grab and ISM samples that need to be shipped is $150 \text{ g} \times n \times k$ and $1500 \text{ g} \times n$, respectively. Therefore, the cost of shipping n ISM samples should be about one tenth the cost of shipping $n \times k$ grab samples. The cost of sample disposal would similarly be reduced.

The ISM approach will result in a cost savings for the determinative (instrumental) portion of the analytical method of a factor of k . However, the additional sample preparation steps needed for the ISM approach increases the total per sample laboratory analytical cost for each metal analysis

(which includes the cost of sample preparation and instrumental analysis). Additional sample preparation is required for both the environmental samples and the batch QC samples, such as method blanks and LCSs. Owing to the sample mass that needs to be processed (e.g., milled), the preparation of LCS for the ISM approach is more costly than for the grab sampling approach. It is conservatively estimated that the ISM approach will increase the total per sample laboratory analysis cost by a factor of no more than two. The sample preparation procedures (e.g., which entail drying, sieving, and milling) are similar to those used for Method 8330B, which increased the cost of these analyses by about \$150 (Hewitt et al. 2009), approximately doubling the per sample cost of an explosive analysis. Similarly, an LCS from Environmental Research Associates (ERA) for the analysis of metals using the ISM approach is similar to that for an LCS for the analysis of the ISM method for explosives using Method 8330B. As the per sample cost for the analysis of a soil sample by a commercial environmental testing laboratory for TAL by Method 3050B/6010B is about \$100, it is reasonable to conclude that the per sample cost for the ISM approach will increase by no more than a factor of two. As $k = 100$ for this effort, the total laboratory portion for the ISM laboratory analyses should be smaller by a factor of $k/2 = 50$ than the total laboratory cost for discrete sample analyses.

7.2 Cost drivers

The main cost drivers that should be considered in selecting the ISM technology for future implementation include the number of DUs or SUs sampled, the number of replicates collected, and the mass of the sample. The key site-specific characteristic that will significantly impact cost is the degree of contaminant heterogeneity expected. If an aqueous metal release occurred, then contaminant heterogeneity is likely to be low; and milling of the sample may not be necessary because adequate precision can likely be achieved without this sample preparation step. However, if metal residues were released into the environment, then sample heterogeneity is likely to be high; and milling would be required. Milling adds approximately \$100 to the per sample cost (Table 10) although the total project costs are likely to be lower depending upon the number of grab samples that would have been collected.

Sample theory indicates that cylindrical cores should be collected. The use of scoop-type samplers, therefore, is discouraged. A device such as CMIST, which collects a cylindrical core, is desired; and the cost investment is

modest and can be recouped through repetitive use of the sampler. A low cost alternative includes the use of 50 cc syringes, with the tip cut off, that can then be pushed into the ground to collect a core; and the plunger is used to eject the soil. A single syringe can be used for multiple increments from the same DU. This device works well in soft unconsolidated soils.

There should be no impact on ISM implementation based on these cost drivers. The biggest cost driver is the purchase of milling equipment by the environmental laboratory and the setup of a dedicated sample processing room. At present, a limited number of commercial laboratories have made the investment in milling equipment. However, as the demand for milling increases as a result of regulatory requirements, it is anticipated that more laboratories will add this to their service capability.

7.3 Cost analysis

For our cost analysis we considered the need to sample a single DU using triplicate ISM samples as opposed to collecting 7 or 15 conventional grab samples (Table 10). The DU consisted of a small-arms range berm 100 m long by 9 m high. The sample depth was 5 cm and the standard metal TAL was assumed. We also assumed a onetime sampling event. We assumed a field sampling crew of 2 individuals although for the demonstration we used either 3 or 4 individuals to speed up the sampling process.

Sample preparation costs were considered equivalent for ISM and grab samples although it has been our experience that the same degree of planning used for ISM is not afforded to conventional grab sampling. However, our assumption was that the same degree of planning and organization of field equipment would occur for both sampling approaches. The expendables used in the demonstration were slightly different and account for a slight cost difference. In the case of grab sampling, the samples were collected in 4-oz, wide-mouth, amber, screw-top jars, which is the norm in environmental sampling. The ISM samples were collected in 15 × 15 in. 3-mm plastic bags and secured with a sample label and a twist tie. This type of container was used to accommodate the larger volume of soil collected. It should be noted that similar sample collection containers could be used for both sampling approaches. Since the same field equipment is used, the mobilization and shipping costs are largely equivalent.

In the field, each individual grab sample is typically surveyed since it has a unique sampling location. In contrast, with ISM, the DU is the unique lo-

cation; so only a single corner of the DU needs to be surveyed if the DU is easily demarcated on a map or an aerial photograph. If the boundaries are less clear, then each corner of the DU can be surveyed. Individual increment sample locations do not need to be surveyed. The sampling activity is essentially the same although with ISM multiple increments are collected and combined to form a single sample. Cylindrical cores need to be collected with ISM to satisfy sample theory. In the demonstration, both ISM and grab samples were collected with the CMIST. However, individuals working in the environmental consulting industry typically use a metal scoop to collect grab samples.

Decontamination of the sampling tool used for ISM is not needed between increments as long as the samples are being collected from within the same DU. While this is not true of replicate samples collected from the same DU, cleaning of the sample tool between replicates is a good management practice and should help to ensure the statistical independence of the replicates.

In contrast, each grab sample is unique and thus requires either disposable sample tools or decontamination between samples. Our decontamination procedure consisted of an acetone rinse followed by a triplicate deionized water rinse. Our cost analysis does not include disposal of rinse of water. However, recovery of this waste water would add slightly to the per sample and total project costs for grab sampling. Because there are typically more samples to label using the grab approach, more time is required for this activity. Costs for field demobilization and shipping of equipment are essentially equivalent as discussed earlier in regards to mobilization activities. There is a difference in sample shipping costs between the two approaches. Although fewer soil samples are collected with ISM, the mass of material collected is 5 to 10 times larger than a grab sample. This more than offsets the greater number of grab samples collected.

The conventional grab sampling approach typically does not involve sample preparation. In some cases, a portion of the soil may be air-dried and given a couple turns in a mortar and pestle. However, discussions with commercial environmental laboratories indicate this is not typical unless the client specifies this activity. The typical approach is for the laboratory to open the 4-oz jar and scoop 0.5 to 2 g of material off the top to use for digestion. As discussed previously, ISM involves air-drying, sieving, milling, and subsampling.

Because more grab samples are collected than with ISM, the associated quality assurance/quality control (QA/QC) costs are higher. The QA/QC analysis includes matrix spike, matrix spike duplicate, laboratory duplicates, and process blanks. This also holds true for the analysis cost, which is the same for both types of samples; but more samples are collected and analyzed with the grab approach.

Our cost analysis indicates field sampling using ISM is \$20–\$40 higher per sample than conventional grab sampling (Table 10). This is largely a function of the greater amount of time needed to collect the ISM sample (i.e., the collection of multiple increments). Similarly, laboratory preparation costs run \$40–\$60 higher with ISM; and analysis, which includes QA/QC, is double the grab-sample cost. This is largely a function of ISM requiring processing of the sample whereas conventional grab sampling typically does not involve sample preparation activities. Therefore, on a per sample basis, the cost of ISM is approximately 55% to 65% higher than conventional grab sampling. The per sample cost for ISM sampling of soil with metal residues ranges from \$300–\$385.

However, the total project cost with ISM is lower than the conventional grab method. This is due to more samples typically collected with grab sampling. For a typical small-arms range DU, three replicate ISM samples would be collected versus 7 to 15 conventional grab samples for the same DU. Therefore, total project costs are to 5% to 50% lower with ISM. The cost savings become greater as the number of DUs/SUs sampled increases. The reduction of costs with ISM is primarily a function of the fewer number of samples needed to adequately characterize an area.

8 Implementation Issues

Promulgation of USEPA Method 8330B (USEPA 2006) has resulted in increased application of ISM and recognition that this approach may be applicable to other analytes in addition to energetics. ITRC (2012) recently published a guidance document that discussed in great technical detail the theory and application of ISM. The DoD and the Army, other government agencies, federal and state regulators, environmental consultants, and commercial laboratories have increasingly discussed the application of ISM to metals. Clausen et al. (2013b) recently published a possible protocol for applying ISM to sites with surface soil containing metallic residues. The ISM sampling strategy is also now mandated by the states of Alaska (Alaska 2009) and Hawaii (Hawaii 2008) for analytes and all surface soil sampling situations. Presently, the authors of this document are working with the USEPA to modify and update SW-846 Method 3050B with the new Method referred to as Method 3050C to accommodate ISM. Proposed Method 3050C includes changes to the laboratory sample preparation procedures, including milling of the samples, and the addition of an Appendix discussing the application of ISM in the field. The proposed changes to Method 3050B are similar to those recommendations presented in Clausen et al. (2013b). Much of the Appendix language is similar to the additions made and promulgated in USEPA Method 8330B. The US Army Corp of Engineer Environmental and Munitions Center of Expertise (EM CX) and DoD are considering changes to existing guidance to incorporate ISM.

The field demonstrations conducted at the three test sites indicate that ISM is readily implementable (Clausen et al. 2013a). No special field equipment is required beyond what is typically used for collecting conventional grab surface soil samples. It is recommended that, to adhere to Gy's theory (1999, 1992), a sample corer device be used so that a cylindrical soil sample is collected. Environmental sampling performed with metal scoops should be discouraged because they do not provide a representative sample (ITRC 2012).

The additional laboratory processing steps outlined with ISM are more involved than what has been used for conventional grab sample processing (Clausen et al. 2012b). The larger sample volume and the need for sample drying, sieving, and milling necessitates a dedicated room at the laborato-

ry for sample processing. This may be problematic for some of the smaller commercial environmental laboratories, but discussions with the larger firms indicate that this is not an impediment to implementability. A number of the larger commercial environmental laboratories have such dedicated sample processing facilities.

Equipment for milling of the soil samples is a potential limiter to the implementability of ISM. Few commercial environmental laboratories have a puck, puck and ring, or roller/ball mill. Although a puck or puck and ring mill is expensive, a roller/ball mill is more affordable. If one is interested in the potential cross-contamination from metallic components of the puck or puck and ring mill, agate bowls and pucks are available. However, currently available agate bowls and pucks are small (i.e., they generally hold less than 600 g of material), thus requiring multiple milling operations to process the entire lot of a single sample. Furthermore, agate milling equipment is expensive compared to steel. As demonstrated by Clausen et al. (2012b), a roller/ball mill with Teflon lined cans and ceramic chips is a lower cost alternative to the puck and puck and ring mills.

The other laboratory changes involve subsampling to prepare the digestion aliquot; digesting an aliquot minimum mass of 5 g; using a consistent acid-to-soil ratio; and the addition of alternative acid solutions for some metals, such as antimony and tungsten, that have poor recoveries with the standard acid digestion procedure of Method 3050B. Discussions with a number of commercial environmental laboratories indicate all of these proposed changes are readily implementable but result in larger unit costs for the metal analyses.

References

- Alaska. 2009. *Draft Guidance on MULTI INCREMENT Soil Sampling*. Anchorage, Alaska: Alaska Department of Environmental Conservation, Division of Spill Preventions and Response, Contaminated Sites Program.
- Ampleman, G., S. Thiboutot, J. Lewis, A. Marois, S. Jean, A. Gagnon, M. Bouchard, R. Martel, R. Lefebvre, C. Gauthier, J. M. Ballard, T. A. Ranney, and T. F. Jenkins. 2003a. *Evaluation of the Impacts of Live Fire Training at CFB Shilo*. DRDC-Valcartier-TR-2003-066. Quebec, Canada: Defence Research Development Canada—Valcartier.
- Ampleman, G., S. Thiboutot, J. Lewis, A. Marois, S. Jean, A. Gagnon, M. Bouchard, T. Jenkins, A. Hewitt, J. C. Pennington, and T. A. Ranney. 2003b. *Evaluation of the Contamination by Explosives in Soils, Biomass and Surface Water at Cold Lake Air Weapons Range (CLAWR), Alberta, Phase 1 Report*. DRDC-Valcartier-TR-2003-208-Annex. Quebec, Canada: Defence Research Development Canada—Valcartier.
- ASTM. 2003. *Standard Guide for Laboratory Subsampling of Media Related to Waste Management Activities*. ASTM D6323-98. West Conshohocken, PA: American Society for Testing and Materials.
- Clausen, J., and N. Korte. 2009a. The distribution of metals in soils and pore water at three U.S. military training facilities. *Soil and Sediment Contamination Journal: An International Journal* 18(5):546–563.
- Clausen, J. L. and N. Korte. 2009b. Environmental fate of tungsten from military use. *The Science of the Total Environment* 407(8):2887–2893.
- Clausen, J. L., J. Robb, D. Curry, and N. Korte. 2004. A case study of contaminants on military ranges: Camp Edwards, Massachusetts, USA. *Environmental Pollution* 129:13–21.
- Clausen, J., S. Taylor, S. Larson, A. Bednar, M. Ketterer, C. Griggs, D. Lambert, A. Hewitt, C. Ramsey, S. Bigl, R. Bailey, and N. Perron. 2007. *Fate and Transport of Tungsten at Camp Edwards Small Arms Ranges*. ERDC/CRREL TR-07-05. Hanover, NH: US Army Engineer Research and Development Center.
- Clausen, J. L., A. Bednar, D. Lambert, R. Bailey, M. Kuhlbrush, S. Taylor, and S. Bigl. 2010. *Phase II Tungsten Fate-and-Transport Study for Camp Edwards*. ERDC/CRREL TR-10-3. Hanover, NH: US Army Engineer Research and Development Center.
- Clausen, J. L., J. Richardson, N. Korte, G. Gooch, T. Hall, N. Perron, E. Butterfield, M. Walsh, and S. Taylor. 2012a. *Metal Residue Deposition from Military Pyrotechnic Devices and Field Sampling Guidance*. ADA562327. Prepared for US Army Environmental Command, Fort Sam Houston, TX. Hanover, NH: US Army Engineer Research and Development Center.
<http://handle.dtic.mil/100.2/ADA562327>.

- Clausen, J. L., T. Georgian, J. Richardson, A. Bednar, N. Perron, L. Penfold, D. Anderson, G. Gooch, T. Hall, and E. Butterfield. 2012b. *Evaluation of Sampling and Sample Preparation Modifications for Soil Containing Metal Residues*. ERDC TR-12-1. Hanover, NH: US Army Engineer Research and Development Center. [http://acwc.sdp.sirsi.net/client/search/asset:asset?t:ac=\\$N/1006020](http://acwc.sdp.sirsi.net/client/search/asset:asset?t:ac=$N/1006020).
- Clausen, J. L., T. Georgian, A. Bednar, N. Perron, A. Bray, P. Tuminello, G. Gooch, N. Mulherin, A. Gelvin, M. Beede, S. Saari, W. Jones, and S. Tazik. 2013a. *Demonstration of Incremental Sampling Methodology for Soil Containing Metallic Residues*. ERDC TR-13-9. Hanover, NH: US Army Engineer Research and Development Center.
- Clausen, J. L., T. Georgian, and A. Bednar. 2013b. *Incremental Sampling Methodology (ISM) for Metallic Residues*. ERDC TR-13-5. Hanover, NH: US Army Engineer Research and Development Center.
- DoD. 2007. *Environmental and Explosives Safety Management on Operational Ranges Within the United States*. DoD Directive 4715.11. Washington, DC: Department of Defense. <http://www.dtic.mil/whs/directives/corres/pdf/471511p.pdf>.
- DoD. 2005. *Operational Range Assessments*. DoD Instruction 4715.14. Washington, DC: Department of Defense. <http://www.dtic.mil/whs/directives/corres/pdf/471514p.pdf>.
- Duncan, A. J. 1962. Bulk sampling: Problems and lines of attack. *Technometrics* 4:319–344.
- Elder, R. S., W. O. Thompson, and R. H. Myers. 1980. Properties of composite sampling procedures. *Technometrics* 22:179–186.
- Felt, D. R., A. J. Bednar, T. Georgian. 2008. The effects of grinding methods on metals concentrations in soil. *Talanta* 77(1):380–387.
- FPM. 2009. *Final Site Inspection Report—Kimama Training Site, Rupert, Idaho*. Rome, NY: FPM Group, Ltd.
- FPM. 2010. *Project Management Plan Military Munitions Response Program Remedial Investigation/Feasibility Study Training Area 3 MRS (KTS-003-R-01)*. Rome, NY: FPM Group, Ltd.
- Gerlach, R. W., and J. M. Nocerino. 2003. *Guidance for Obtaining Representative Laboratory Analytical Subsamples from Particulate Laboratory Samples*. EPA/600/R-03/027. Washington, DC: US Environmental Protection Agency.
- Gy, P. 1992. *Sampling of Particulate Materials Theory and Practice*. New York, NY: Elsevier Scientific Publishing Company.
- Gy, P. 1999. *Sampling for Analytical Purposes*. New York, NY: John Wiley and Sons.
- Hawaii. 2008. Section 4, Soil Sample Collection Approaches. In *Interim Final Technical Guidance Manual for the Implementation of the Hawaii State Contingency Plan*. Hawaii Department of Health, Office of Hazard Evaluation and Emergency Response. <http://www.hawaiidoh.org/>.

- Hewitt, A. D., and M. E. Walsh. 2003. *On-site homogenization and sub sampling of surface samples for analysis of explosives*. ERDC/CRREL TR 03-14. Hanover, NH: US Army Engineer Research and Development Center.
- Hewitt, A. D., T. F. Jenkins, C. A. Ramsey, K. L. Bjella, T. A. Ranney, and N. M. Perron. 2005. *Estimating energetic residue loading on military artillery ranges: Large decision units*. ERDC/CRREL TR-05-7. Hanover, NH: US Army Engineer Research and Development Center.
http://www.crrel.usace.army.mil/techpub/CRREL_Reports/reports/TR05-7.pdf.
- Hewitt A. D., T. F. Jenkins, M. E. Walsh, M. R. Walsh, S. R. Bigl, and C. A. Ramsey. 2007. *Protocols for collection of surface soil samples at military training and testing ranges for the characterization of energetic munition constituents*. ERDC/CRREL TR-07-10. Hanover, NH: US Army Engineer Research and Development Center.
- Hewitt, A. D., T. F. Jenkins, M. E. Walsh, S. R. Bigl, and S. Brochu. 2009. *Validation of Sampling Protocol and the Promulgation of Method Modifications for the Characterization of Energetic Residues on Military Testing and Training Ranges*. ERDC/CRREL TR-09-6. Hanover, NH: US Army Engineer Research and Development Center.
<http://libweb.erdcl.usace.army.mil/uhtbin/cgisirsi/20110317135155/SIRSI/0/518/0/CRREL-TR-09-6.pdf>.
- Hewitt, A. D., T. F. Jenkins, S. R. Bigl, J. L. Clausen, H. Craig, M. E. Walsh, R. Martel, and K. Nieman. 2011. *EPA Federal Facilities Forum Issue Paper: Site Characterization for Munitions Constituents*. EPA-505-S-11-01. Washington, DC: US Environmental Protection Agency, Solid Waste and Emergency Response, Federal Facilities Forum Issue.
- Idaho Geological Survey. 2011. *Geologic Mapping*. Moscow, ID: Idaho Geological Survey.
<http://www.idahogeology.org/Services/GeologicMapping/>.
- ITRC. 2012. *Technical and Regulatory Guidance: Incremental Sampling Methodology*. ISM-1. Washington, DC: Interstate Technology and Regulatory Council, Incremental Sampling Methodology Team. <http://itrcweb.org/ism-1/>.
- Jenkins, T. F., C. L. Grant, G. S. Brar, P. G. Thorne, T. A. Ranney, and P. W. Schumacher. 1996. *Assessment of Sampling Error Associated with Collection and Analysis of Soil Samples at Explosives-Contaminated Sites*. Special Report 96-15. Hanover, NH: US Army Cold Regions Research and Engineering Laboratory.
- Jenkins, T. F., M. E. Walsh, P. G. Thorne, S. Thiboutot, G. Ampleman, T. A. Ranney, and C. L. Grant. 1997a. *Assessment of Sampling Error Associated with the Collection and Analysis of Soil Samples at a Firing Range Contaminated with HMX*. Special Report 97-22. Hanover, NH: US Army Cold Regions Research and Engineering Laboratory.
- Jenkins, T. F., C. L. Grant, G. S. Brar, P. G. Thorne, P. W. Schumacher, and T. A. Ranney. 1997b. *Sampling Error Associated with Collection and Analysis of Soil Samples at TNT Contaminated Sites*. *Field Analytical Chemistry Technology* 1:151–163.

- Jenkins, T. F., M. E. Walsh, P. G. Thorne, P. H. Miyares, T. A. Ranney, C. L. Grant, and J. R. Esparza. 1998 *Site Characterization for Explosives Contamination at a Military Firing Range Impact Area*. Special Report 98-9. Hanover, NH: US Army Cold Regions Research and Engineering Laboratory.
- Jenkins, T. F., C. L. Grant, M. E. Walsh, P. G. Thorne, S. Thiboutot, G. Ampleman, and T. A. Ranney. 1999. Coping with spatial heterogeneity effects on sampling and analysis at an HMX—Contaminated Antitank Firing Range. *Field Analytical Chemistry Technology* 3:19–28.
- Jenkins, T. F., J. C. Pennington, T. A. Ranney, T. E. Berry, P. H. Miyares, M. E. Walsh, A. D. Hewitt, N. M. Perron, L. V. Parker, C. A. Hayes, and E. G. Wahlgren. 2001. *Characterization of Explosives Contamination at Military Firing Range*. ERDC TR-01-5. Hanover, NH: US Army Engineer Research and Development Center.
- Jenkins, T. F., T. A. Ranney, A. D. Hewitt, M. E. Walsh, and K. L. Bjella. 2004a. *Representative sampling for energetic compounds at an antitank firing range*. ERDC/CRREL TR-04-7. Hanover, NH: US Army Engineer Research and Development Center. <http://www.crrel.usace.army.mil/library/technicalreports/TR04-7.pdf>.
- Jenkins, T. F., A. D. Hewitt, T. A. Ranney, C. A. Ramsey, D. J. Lambert, K. L. Bjella, and N. M. Perron. 2004b. *Sampling strategies near a low-order detonation and a target at an artillery impact area*. ERDC/CRREL TR-04-14. Hanover, NH: US Army Engineer Research and Development Center. <http://www.crrel.usace.army.mil/library/technicalreports/TR04-14.pdf>.
- Jenkins, T. F., A. D. Hewitt, M. E. Walsh, T. A. Ranney, C. A. Ramsey, C. L. Grant, and K. L. Bjella. 2005. Representative Sampling for Energetic Compounds at Military Training Ranges. *Environmental Forensics* 6:45–55.
- Jenkins T. F., A. D. Hewitt, C. L. Grant, S. Thiboutot, G. Ampleman, M. E. Walsh, T. A. Ranney, C. A. Ramsey, A. J. Palazzo, and J. C. Pennington. 2006. Identity and distribution of residues of energetic compounds at Army live-fire training ranges. *Chemosphere* 63:1280–1290.
- Johanson, J. R. 1978. Particle segregation and what to do about it. *Chemical Engineering* May:183–188.
- Leutwyler, K. 1993. Shaking conventional wisdom. *Scientific American* September:24.
- Nieman, K. C. 2007. Personal communication with Jay Clausen. Hill Air Force Base, UT: Select Engineering Services, 75 CEG/CEVC.
- Pennington, J. C., et al. 2004. *Distribution and fate of energetics on DoD test and training ranges: Interim Report 4*. ERDC TR-04-4, Vicksburg, MS: US Army Engineer Research and Development Center. <http://el.erdcl.usace.army.mil/elpubs/pdf/tr04-4.pdf>.
- Pitard, F. F. 1993. *Pierre Gy's Sampling Theory and Sampling Practice: Heterogeneity, Sampling Correctness, and Statistical Process Control*. Boca Raton, FL: CRC Press.

- Racine, C. H., M. E. Walsh, C. M. Collins, D. J. Calkins, B. D. Roebuck, and L. Reitsma. 1992. *Waterfowl Mortality in Eagle River Flats, Alaska: The Role of Munition Residues*. CRREL Report 92-5. Hanover, NH: Cold Regions Research and Engineering Laboratory.
- Studt, T. 1995. For material researchers, It's back to the sandbox. *R&D Magazine* July:41–42.
- Taylor S., A. Hewitt, J. Lever, C. Hayes, L. Perovich, P. Thorne, and P. Daghalin. 2004. TNT particle size distribution for detonated 155-mm howitzer rounds. *Chemosphere* 55:357–367.
- Thiboutot, S., G. Ampleman, A. Gagnon, A. Marois, T. F. Jenkins, M. E. Walsh, P. G. Thorne, and T. A. Ranney. 1998. *Characterization of Antitank Firing Ranges at CDB Valcartier, WATC Wainwright and CFAD Dundurn*. Quebec, Canada: Defence Research Establishment—Valcartier.
- Thiboutot, S., G. Ampleman, A. Gagnon, and A. Marois. 2000a. *Characterization of an Unexploded Ordinance Contaminated Range (Tracadie Range) for Potential Contamination by Energetic Materials*. DREV-TR-2000-102. Quebec, Canada: Defence Research Establishment—Valcartier .
- Thiboutot, S., G. Ampleman, P. Dube, C. Dubois, R. Martel, R. Lefebvre, M. Mailloux, G. Sunahara, P. Y. Roubidoux, and J. Hawari. 2000b. *Characterization of DND Training Ranges Including Anti-Tank Firing Ranges and Ecotoxicological Assessment*. Quebec, Canada: Defence Research Establishment—Valcartier.
- Thiboutot, S., G. Ampleman, and A.D. Hewitt. 2002. *Guide for Characterization of Sites Contaminated with Energetic Materials*. ERDC/CRREL TR-02-1. Hanover, NH: US Army Engineer Research and Development Center.
- Thiboutot, S., G. Ampleman, J. Lewis, D. Faucher, A. Marois, R. Martel, J. M. Ballard, S. Downe, T. F. Jenkins, A. Hewitt. 2003. *Environmental Conditions of Surface Soils and Biomass Prevailing in the Training Area at CFB Gagetown, New Brunswick*. DRDC-TR-2003-152, Quebec, Canada: Defence Research Development Canada—Valcartier.
- URS. 2007. *Final Site Inspection Report, Fort Eustis, Fort Eustis, Virginia, Military Munitions Response Program*. Gaithersburg, MD: URS Group Inc.
- URS. 2010. *Remedial Investigation/Feasibility Study 1000" Rifle Range Fort Eustis, Virginia Military Munitions Response Program, Volume 1: Remedial Investigation*. Gaithersburg, MD: URS Group Inc.
- USACE. 1972. *Environmental Analysis Record: Amended Special Land-use Permit Application, I-2407*. Seattle, WA: US Corps of Engineers, Seattle District.
- USACE. 2009. *Implementation of Incremental Sampling (IS) of Soil for the Military Munitions Response Program, Environmental and Munitions Center of Expertise Interim Guidance Document (IGD) 09-02*. Huntsville, AL: Department of the Army, Huntsville Center, Corps of Engineers.

- USEPA. 2006. Method 8330B: Nitroaromatics, nitramines, nitrate esters by high performance liquid chromatography (HPLC). In *Test Methods for Evaluating Solid Waste, Physical/Chemical Methods, Office of Solid Waste and Emergency Response*. SW-846. Washington, DC: US Environmental Protection Agency. <http://www.epa.gov/epaoswer/hazwaste/test/pdfs/8330b.pdf>.
- Wallace, D., and B. Kratochvil. 1985. Use of a mill and spinning riffle for subsampling laboratory samples of oil sand. *Austria Journal of Research* 2:233–239.
- Walsh, M. E., and C. M. Collins. 1993. *Distribution of White Phosphorus in Residues from the Detonation of 81 mm Mortar WP Smoke Rounds*. Special Report 93-18. Hanover, NH: US Army Cold Regions Research and Engineering Laboratory.
- Walsh, M. E., and D.J. Lambert. 2006. *Extraction Kinetics of Energetic Compounds from Training Range and Army Ammunition Plant Soils: Platform Shaker versus Sonic Bath Methods*. ERDC/CRREL TR-06-6. Hanover, NH: US Army Engineers Research and Development Center.
- Walsh, M. E., T. F. Jenkins, P. H. Miyares, P. S. Schnitker, J. W. Elwell, and M. H. Stutz. 1993. *Evaluation of SW846 Method 8330 for Characterization of Sites Contaminated with Residues of High Explosives*. CRREL Report 93-5. Hanover, NH: US Army Cold Regions Research and Engineering Laboratory.
- Walsh, M. E., C. M. Collins, R. N. Bailey, and C. L. Grant. 1997. *Composite Sampling of Sediments Contaminated with White Phosphorus*. Special Report 97-30. Hanover, NH: US Army Cold Regions Research and Engineering Laboratory.
- Walsh, M. E., C. A. Ramsey, and T. F. Jenkins. 2002. The Effect of Particle Size Reduction on Sub sampling Variance for Explosives Residues in Soil. *Chemosphere* 49:1265–1271.
- Walsh, M. E., C. M. Collins, T. F. Jenkins, A. D. Hewitt, J. Stark, and K. Myers. 2003. Sampling for Explosives-Residues at Ft. Greely. *Soil and Sediment Contamination* 12:631–645.
- Walsh, M. E., C. M. Collins, A. D. Hewitt, M. R. Walsh, T. F. Jenkins, J. Stark, A. Gelvin, T. S. Douglas, N. Perron, D. Lambert, R. Bailey and K. Meyers. 2004. *Range characterization studies at Donnelly Training Area, Alaska: 2001 and 2003*. ERDC/CRREL TR-04-3. Hanover, NH: US Army Engineer Research and Development Center. <http://www.crrel.usace.army.mil/library/technicalreports/TR04-3.pdf>.
- Walsh, M. E., C. A. Ramsey, C. M. Collins, A. D. Hewitt, M. R. Walsh, K. Bjella, D. Lambert, and N. Perron. 2005. *Collection methods and laboratory processing of samples from Donnelly Training Area Firing Points Alaska 2003*. ERDC/CRREL TR-05-6. Hanover, NH: US Army Engineer Research and Development Center. <http://www.crrel.usace.army.mil/library/technicalreports/TR05-6.pdf>.
- Walsh, M. E., C. A. Ramsey, S. Taylor, A. D. Hewitt, K. Bjella, C. M. Collins. 2006. Sub sampling Variance for 2,4-DNT in Firing Point Soils. *Soil and Sediment Contamination: an International Journal* 16(5):459–472.
- Walsh, M. R. 2009. *User's Manual for the CRREL Multi-Increment Sampling Tool*. ERDC/CRREL SR-09-1. Hanover, NH: US Army Engineer Research and Development Center. <http://acwc.sdp.sirsi.net/client/default>.

Appendix A: Points of Contact

POINT OF CONTACT Name	ORGANIZATION Name Address	Phone Fax E-mail	Role in Project
Jay Clausen	ERDC/CRREL 72 Lyme Road Hanover, NH 03755	603-646-4597 603-646-4785 jay.l.clausen@us.army.mil	PI
Anthony Bednar	ERDC/EL 3909 Halls Ferry Road EP-C Building 3299 RM 106 Vicksburg, MS 39180-6199	601-634-3652 601-634-2742 anthony.j.bednar.usa.army.mil	Co-PI
Thomas Georgian	USACE EM CX 1616 Capitol Ave Omaha, NE 68102	402-697-2567 thomas.georgian@us.army.mil	Co-PI
Larry Penfold	Test America 4955 Yarrow St Arvada, CO 80002	303-736-0119 303-502-6559 larry.Penfold@testamericainc.com	Industry Partner
Diane Anderson	APPL, Inc. 908 N. Temperance Ave. Clovis, CA 93611	559-275-2175 559-275-4422 danderson@applinc.com	Industry Partner
Amber Michel	AGEISS, Inc. 733d MSG/CED/AMF/EE 1407 Washington Blvd. JBLE (Fort Eustis), Virginia 23604-5306	757-878-4123 ext 296 757-878-4589 Amber.Michel@us.army.mil	Fort Eustis Site Access

REPORT DOCUMENTATION PAGE				Form Approved OMB No. 0704-0188	
Public reporting burden for this collection of information is estimated to average 1 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing this collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to Department of Defense, Washington Headquarters Services, Directorate for Information Operations and Reports (0704-0188), 1215 Jefferson Davis Highway, Suite 1204, Arlington, VA 22202-4302. Respondents should be aware that notwithstanding any other provision of law, no person shall be subject to any penalty for failing to comply with a collection of information if it does not display a currently valid OMB control number. PLEASE DO NOT RETURN YOUR FORM TO THE ABOVE ADDRESS.					
1. REPORT DATE (DD-MM-YYYY) September 2013		2. REPORT TYPE Technical Report/Final		3. DATES COVERED (From - To)	
4. TITLE AND SUBTITLE Cost and Performance Report of Incremental Sampling Methodology for Soil Containing Metallic Residues				5a. CONTRACT NUMBER	
				5b. GRANT NUMBER	
				5c. PROGRAM ELEMENT NUMBER	
6. AUTHOR(S) Jay L. Clausen, Thomas Georgian, and Anthony Bednar				5d. PROJECT NUMBER ER-0918	
				5e. TASK NUMBER	
				5f. WORK UNIT NUMBER	
7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES) Cold Regions Research and Engineering Laboratory (CRREL) US Army Engineer Research and Development Center 72 Lyme Road Hanover, NH 03755-1290				8. PERFORMING ORGANIZATION REPORT NUMBER ERDC TR-13-10	
9. SPONSORING / MONITORING AGENCY NAME(S) AND ADDRESS(ES) Environmental Security Technology Certification Program (ESTCP) Arlington, VA 22203				10. SPONSOR/MONITOR'S ACRONYM(S) ESTCP	
				11. SPONSOR/MONITOR'S REPORT NUMBER(S)	
12. DISTRIBUTION / AVAILABILITY STATEMENT Approved for public release; distribution is unlimited.					
13. SUPPLEMENTARY NOTES					
14. ABSTRACT Objectives of this project were to demonstrate improved data quality for metal constituents in surface soils on military training ranges and to develop a methodology that would result in the same or lower cost. The demonstration was conducted at two inactive small-arms ranges at Fort Eustis, VA, and Kimama Training Site (TS), ID, and at one active small-arms range at Fort Wainwright, AK. The samples included 63 Incremental Sampling Methodology (ISM) and 50 conventional grab from Fort Wainwright, 18 ISM and 30 grab from Kimama TS, and 27 ISM and 33 grab from Fort Eustis. The variability in metal concentrations as measured with replicate samples and evaluated using percent relative standard deviation (RSD) were less than 10% for all metals using ISM. In contrasts, RSDs were often greater than 50% for conventional replicate grab samples. Calculated mean ISM metal concentrations were statistically greater than the mean for conventional grab samples.					
15. SUBJECT TERMS Antimony Copper		Decision unit (DU) Incremental sampling methodology (ISM) Lead		Metals Ranges Surface soil	
16. SECURITY CLASSIFICATION OF:			17. LIMITATION OF ABSTRACT	18. NUMBER OF PAGES	19a. NAME OF RESPONSIBLE PERSON
a. REPORT	b. ABSTRACT	c. THIS PAGE			19b. TELEPHONE NUMBER (include area code)
U	U	U	U	80	