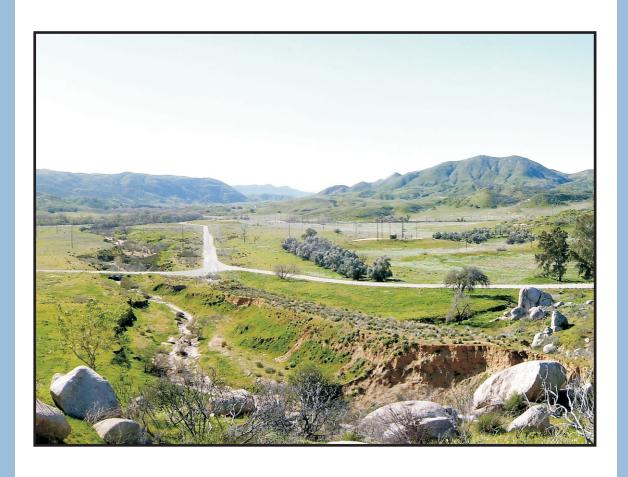
Semiannual Groundwater Monitoring Report Third Quarter and Fourth Quarter 2012 Potrero Canyon (Lockheed Martin Beaumont Site 1) Beaumont, California







Lockheed Martin Enterprise Business Services Energy, Environment, Safety and Health 2950 North Hollywood Way, Suite 125 Burbank, CA 91505 Telephone 818•847•0197 Facsimile 818•847•0256



April 23, 2013

Mr. Daniel Zogaib Southern California Cleanup Operations Department of Toxic Substances Control 5796 Corporate Avenue Cypress, CA 90630

Subject: Submittal of the Semiannual Groundwater Monitoring Report, Third Quarter and Fourth Quarter 2012, Lockheed Martin Corporation, Potrero Canyon (Lockheed Martin Beaumont Site 1), Beaumont, California

Dear Mr. Zogaib:

Please find enclosed one hard copy of the body of the report and two compact disks with the report body and appendices of the Semiannual Groundwater Monitoring Report, Third Quarter and Fourth Quarter 2012, Lockheed Martin Corporation, Potrero Canyon (Lockheed Martin Beaumont Site 1), Beaumont, California for your review and approval or comment.

If you have any questions regarding this submittal, please contact me at 818-847-9901 or brian.thorne@lmco.com.

Sincerely,

Bin 7. Um

Brian T. Thorne Project Lead

Enclosure: Semiannual Groundwater Monitoring Report, Third and Fourth Quarter 2012, Lockheed Martin Corporation, Potrero Canyon (Lockheed Martin Beaumont Site 1), Beaumont, California

Copy: Gene Matsushita, LMC (electronic and hard copy)
Barbara Melcher, CDM Smith (electronic copy)
Tom Villeneuve, Tetra Tech (electronic copy)
Alan Bick, Gibson Dunn (electronic copy)

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SEMIANNUAL GROUNDWATER MONITORING REPORT THIRD QUARTER AND FOURTH QUARTER 2012 POTRERO CANYON (LOCKHEED MARTIN BEAUMONT SITE 1) BEAUMONT, CALIFORNIA

Prepared for:

Lockheed Martin Corporation

Prepared by:

Tetra Tech, Inc.

April 2013

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ACRONYMS AND ABBREVIATIONS

BPA Burn Pit Area

BGS below ground surface

BTOC below top of casing

cfs cubic feet per second

1,1-DCA 1,1-dichloroethane

1,1 -DCE 1,1-dichloroethene

cis-1,2-DCE cis-1,2-dichloroethene

DG downgradient

DO dissolved oxygen

DOC dissolved organic carbon

DWNL California Department of Public Health drinking water notification level

EC electrical conductivity

GPS global positioning system

HCP Habitat Conservation Plan

LCS laboratory control sample

MCL California Department of Public Health maximum contaminant level

MCEA Massacre Canyon Entrance Area

MDL method detection limit

MEF Mount Eden formation

mg/L milligrams per liter

μg/L micrograms per liter

μg/L/yr micrograms per liter per year

MS/MSD matrix spike/matrix spike duplicate

msl mean sea level

NA not analyzed / not applicable / not available

ND non-detect

nM nanoMoles

NPCA Northern Potrero Creek Area

NTU nephelometric turbidity unit

NWS National Weather Service

ORP oxidation-reduction potential

PQL practical quantitation limit

psi pounds per square inch

QAL Quaternary alluvium

QAL/MEF Quaternary alluvium/Mount Eden formation

QA/QC quality assurance/quality control

Radian Corporation, Inc.

RMPA Rocket Motor Production Area

RPD relative percent difference

site Potrero Canyon (Lockheed Martin Beaumont Site 1)

1,1,1-TCA 1,1,1-trichloroethane

Tetra Tech, Inc.

TCE trichloroethene

UG upgradient

USEPA United States Environmental Protection Agency

USFWS United States Fish and Wildlife Service

VFA volatile fatty acid

VOC volatile organic compound

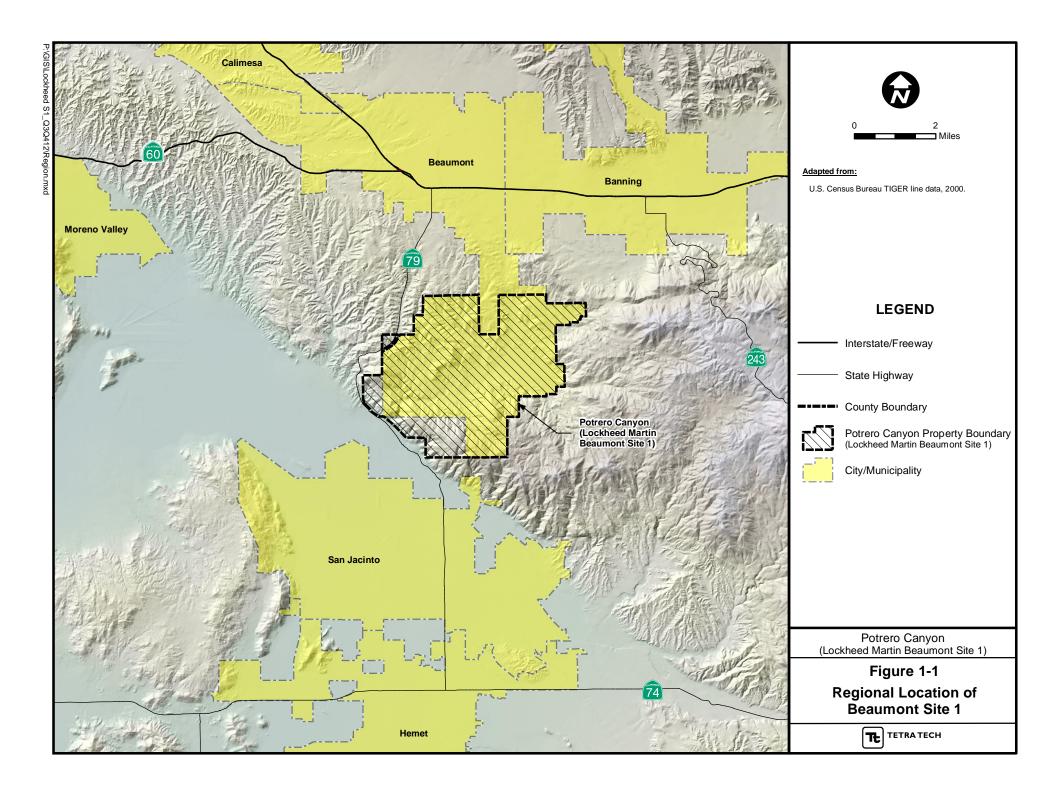
SECTION 1 INTRODUCTION

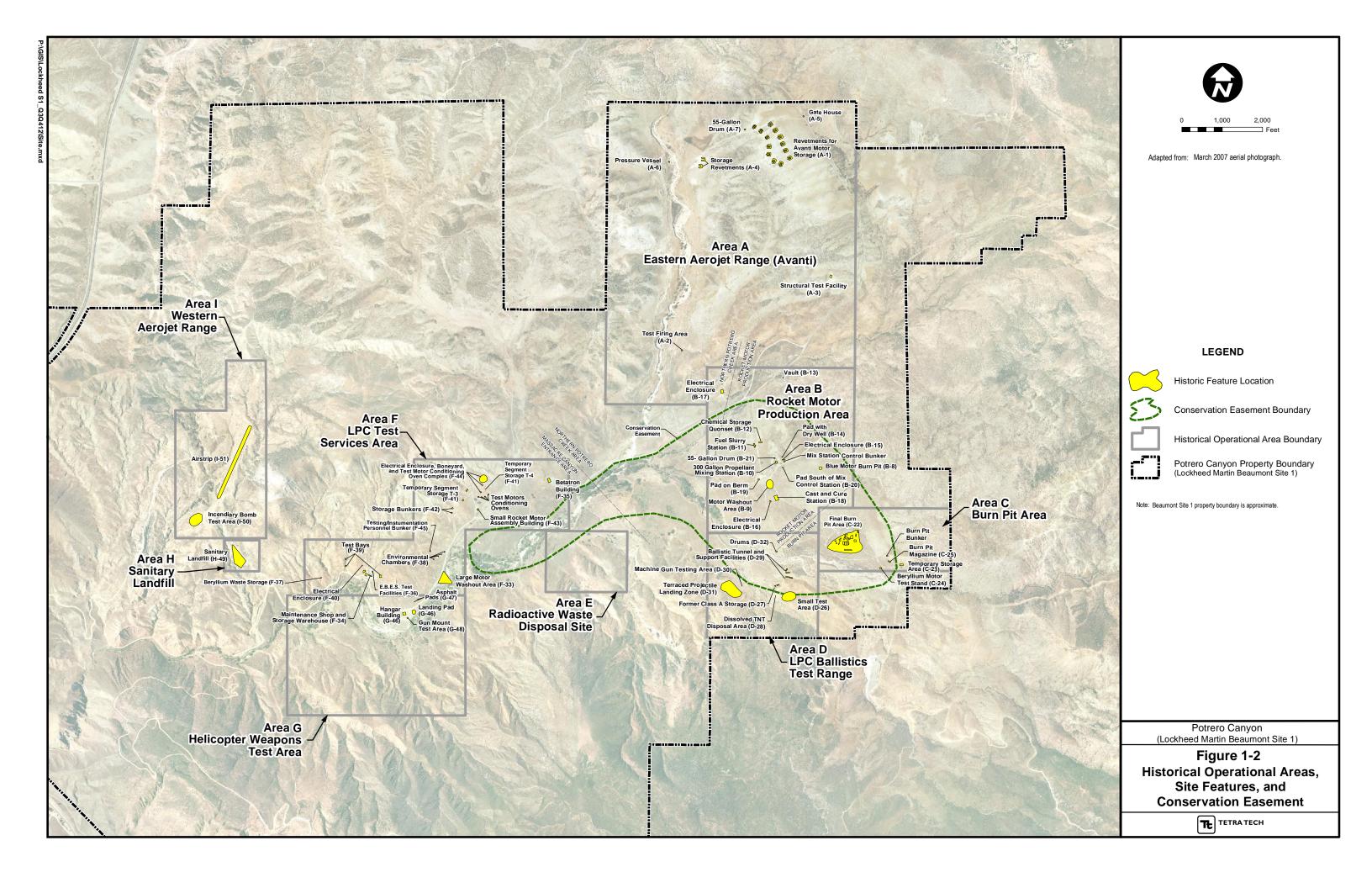
On behalf of Lockheed Martin Corporation, Tetra Tech, Inc. has prepared this Semiannual Groundwater Monitoring Report, which presents the results of the Third Quarter 2012 (1 July 2012 through 30 September 2012) and Fourth Quarter 2012 (1 October 2012 through 31 December 2012) water quality monitoring for the Potrero Canyon (Lockheed Martin Beaumont Site 1) Groundwater Monitoring Program. The site is located in an undeveloped area in the southern portion of the city of Beaumont in Riverside County, California (Figure 1-1). Currently, the site is inactive except for environmental investigations performed under Consent Order 88/89-034 and Operation and Maintenance Agreement 93/94-025 with the California Department of Toxic Substances Control. The State of California owns approximately 94% (8,552 acres) of the site. The remaining 565 acres, referred to as the conservation easement, were retained by Lockheed Martin Corporation (Figure 1-2).

The Groundwater Monitoring Program includes quarterly, semiannual, annual, and biennial monitoring tasks with both groundwater and surface water collected and sampled as shown in Appendix A, Table 1-1. The annual and biennial events are larger major monitoring events, and the quarterly and semiannual events are smaller minor events. All new wells are sampled quarterly for one year. Semiannual wells are sampled the second and fourth quarter of each year, annual wells are sampled the second quarter of each year, and biennial wells are sampled during the second quarter of even-numbered years.

The objectives of this report are to accomplish the following:

- Briefly summarize the site history
- Document water level and water quality monitoring activities and results
- Analyze and evaluate the groundwater elevation and water quality monitoring data generated





This report is organized into the following sections: (1) Introduction, (2) Summary of Monitoring Activities, (3) Groundwater Monitoring Results, (4) Summary and Conclusions, and (5) References. Appendix A provides a brief description of the previous site environmental investigations and the current conceptual site model.

1.1 SITE BACKGROUND

The site consists of a 9,117-acre parcel located in the southern portion of Beaumont, California. The site was primarily used for ranching before 1960. From 1960 to 1974, Lockheed Propulsion Company used the site for solid rocket motor and ballistics testing (Tetra Tech, 2003a). Activities at the site also included burning of process chemicals and waste rocket propellants in an area commonly referred to as the Burn Pit Area.

Tetra Tech identified nine primary historical operational areas at the site. A map of site historical operational areas and features is presented as Figure 1-2. Historical operational areas were used for various activities associated with rocket motor assembly, testing, and propellant incineration. A brief description of each historical operational area follows.

Historical Operational Area A – Eastern Aerojet Range

Between 1970 and 1972, Aerojet leased an area (referred to as the Eastern Aerojet Range) along the eastern portion of the site. The Eastern Aerojet Range was used periodically for ballistics research and development experimentation on several types of 30-millimeter projectiles. Avanti, a highly classified project, used the land directly east of the Eastern Aerojet Range, including several U-shaped revetments for the storage of explosive materials and rocket motors. The purpose of the Avanti project and its operational procedures are unknown, due to its classified status (Radian, 1986).

Historical Operational Area B – Rocket Motor Production Area

The Rocket Motor Production Area, also known as the Propellant Mixing Area, was used for the processing and mixing of rocket motor solid propellants. The rocket motor production process consisted of: 1) a fuel slurry station, 2) a mixing station, and 3) a cast and curing station.

If a defect was found in the solid propellant mix, the rocket motor was scrapped. The solid propellant was removed from the casings by water jetting at the motor washout south of the mixing station (Radian, 1986).

In 1973, an area east of the mixing station, known as the Blue Motor Burn Pit, was used for the destruction of four motors, which included a motor with "Malloy blue" solid propellant, also referred to as milori blue or Prussian blue (Radian, 1986).

Historical Operational Area C – Burn Pit Area

The Burn Pit Area had three primary features: (1) the chemical storage area, (2) burn pits, and (3) the beryllium test stand. Hazardous wastes generated at the site were stored in 55-gallon drums on a concrete pad east of the burn pits at the chemical storage area until enough material was accumulated for a burning event. The hazardous materials burned in the pits included ammonium perchlorate, wet propellant from motor washout, dry propellant, batches of out-of-specification propellant, various kinds of adhesives, resin curatives such as polybutadiene acrylonitrile/acrylic acid copolymer, burn rate modifiers such as ferrocene, pyrotechnic and ignition components, packaging materials (e.g., metal drums, plastic bags, and paper drums), and solvents (Radian, 1986).

On the south side of the bedrock outcrop where the burn pit instrumentation bunker was located, there was a one-time firing of small beryllium research motors.

Historical Operational Area D – Lockheed Propulsion Company Ballistics Test Range

The Lockheed Propulsion Company Ballistics Test Range facilities included gun mounts, a ballistic tunnel, and storage buildings and trailers. Guns were tested by firing through the tunnel toward a terraced hill. Live rounds were not used, although projectiles were often specially shaped and weighted to simulate actual live rounds (Radian, 1986). Another major project conducted in this area was experimentation on a rocket-assisted projectile to test penetration capability. Additional experiments included impact testing of various motors and pieces of equipment (Radian, 1986).

Class A explosives were reportedly stored in two or three 10-foot by 10-foot buildings behind a berm. A small canyon behind the hill to the south of the former storage buildings was reportedly used as a small test area for incendiary bombs. An incendiary bomb was detonated in the center of drums containing various types of fuel (e.g., jet fuel, gasoline, and diesel) set in circles of different radii to observe shrapnel and penetration patterns. (Alternatively, this test may have been

conducted in Historical Operational Area I.) At a small area near the bend in the road, acetone was used to dissolve 2,4,6-trinitrotoluene out of projectiles before they were fired (Radian, 1986).

Historical Operational Area E – Radioactive Waste Disposal Site

During 1971, low-level radioactive waste was buried in one of four canyons southeast of the Lockheed Propulsion Company test services area as reported by former site employees. In 1990, the radioactive waste was located and removed. Soil samples were collected after removal of the waste. The analytical results indicated that detected radiation levels were within the range of naturally occurring levels (Radian, 1990). Maps from the removal action report suggest the waste was removed from Canyon 2.

Historical Operational Area F – Lockheed Propulsion Company Test Services Area

The Lockheed Propulsion Company Test Services Area included the following features: (1) three bays for structural load tests, (2) a 13-foot-diameter spherical pressure vessel, (3) six temperature-conditioning chambers, (4) four environmental chambers, (5) a 25-million electron volt Betatron for X-raying large structures, (6) personnel and instrumentation protection bunkers, and (7) supporting workshops and storage areas (Radian, 1986).

If defects were identified during the integrity and environmental testing activities, the rocket motors were taken to the large motor washout area south of the conditioning chambers adjacent to Potrero Creek (Radian, 1986).

Rocket motor structural load testing under static and captive firing conditions occurred at the Lockheed Propulsion Company test bays. During several of the initial tests conducted at Bay 309, the readied motor exploded instead of firing (Radian, 1986).

Historical Operational Area G – Helicopter Weapons Test Area

The helicopter weapons test area was used to develop equipment for handling helicopter weapons systems. The facilities in this area included a hangar (Building 302), helicopter landing pad, stationary ground-mounted gun platforms, and a mobile target suspended between towers. The primary project at this area was testing of both stationary guns and guns mounted on helicopters. Experimentation was also performed on the solid propellant portion of an armor-piercing round. Most rounds were fired into the side of the creek wash, about 100 yards to the south of the hangar. A longer impact area labeled with distance markers was located in the canyon to the south of the

wash. Projectiles were steel only; warheads were not used during tests at this facility (Tetra Tech, 2003a).

Historical Operational Area H – Sanitary Landfill

A permitted sanitary landfill was located along the western side of the site. The permit for the landfill authorized Lockheed Propulsion Company to dispose of trash such as paper, scrap metal, concrete, and wood generated during routine daily operations. Lockheed policy strictly dictated that no hazardous materials were to have been disposed at this landfill. The trenches were later covered and leveled, with only an occasional tire, metal scrap, or piece of wood remaining on the surface (Tetra Tech, 2003a).

Historical Operational Area I – Western Aerojet Range

Between 1970 and 1972, Aerojet leased an area (referred to as the Western Aerojet Range) along the western portion of the site. Lockheed Propulsion Company conducted an incendiary test with a 500-pound bomb at the southwest end of the Western Aerojet Range. This test was reportedly similar to testing performed at the Lockheed Propulsion Company Ballistics Test Area. According to Radian's historical report, the Western Aerojet Range was originally leveled to be used as an airstrip (Radian, 1986). Based on employee interviews, the airstrip may have been used only on one occasion (Tetra Tech, 2003a). During investigations performed in 2006 for munitions and explosives of concern (Tetra Tech, 2007), it was discovered that inert 27.5-millimeter projectiles were tested in this area.

Post Lockheed Propulsion Company and Aerojet Facility Usage

Lockheed Martin Corporation leased portions of the site to several outside parties for use in various activities (Radian, 1986; Tetra Tech, 2003a). The International Union of Operating Engineers used the site from 1971 through 1991 for surveying and heavy equipment training. The Union's main office was located in Bunker 304 of Historical Operational Area F (Lockheed Propulsion Company Test Services Area). The Union's earth-moving actions involved maintaining roads and reshaping various parts of the site, primarily in Historical Operational Areas F and G.

On several occasions, General Dynamics used Historical Operational Area B (Rocket Motor Production Area) for testing activities (Radian, 1986). In 1983 and 1984, General Dynamics conducted weapons testing of a Viper Bazooka and Phalanx Gatling gun.

Structural Composites used the steep terrain of the site for vehicle rollover tests on a number of occasions. Structural Composites also conducted heat and puncture tests on pressurized fiberglass and plastic reinforced cylinders. The tests involved shooting a single 30-caliber round at the cylinders and recording the results (Radian, 1986).

SECTION 2 SUMMARY OF MONITORING ACTIVITIES

Section 2 summarizes the Third Quarter 2012 and Fourth Quarter 2012 groundwater monitoring events conducted at the site. The results from these monitoring events are discussed in Section 3.

2.1 GROUNDWATER LEVEL MEASUREMENTS

Groundwater level measurements are collected at the site on a quarterly basis from all available wells. Water level measurements were proposed for 182 wells for the Third Quarter 2012 and Fourth Quarter 2012 monitoring events. The Third Quarter 2012 groundwater level measurements were collected between 13 August and 15 August 2012. The Fourth Quarter 2012 groundwater level measurements were collected between 13 November and 14 November 2012. Appendix B provides copies of field data sheets from the water quality monitoring events, and Appendix C presents a summary of well construction details.

Precipitation data are collected from the local weather station in Beaumont to correlate observed changes in groundwater levels with local precipitation. During Third Quarter 2012, the Beaumont National Weather Service (NWS) station reported approximately 0.30 inches of precipitation. During Fourth Quarter 2012, the Beaumont NWS reported approximately 3.62 inches of precipitation.

2.2 SURFACE WATER FLOW

The site is primarily drained by Potrero Creek, an ephemeral stream which follows the valley from north to south before turning southwest to pass through Massacre Canyon toward its convergence with the San Jacinto River. Potrero Creek is fed by local tributary drainage and storm water runoff from the city of Beaumont as well as from other ephemeral streams in the southern and eastern portions of the site. The largest of the tributary drainages is Bedsprings Creek, which is southwest of the former Rocket Motor Production Area (RMPA) and former Burn Pit Area (BPA). In general, creeks are dry except during and immediately after periods of rainfall. However, springs and seeps occur in and adjacent to Potrero Creek in the western portion of the site. Surface water flow is not continuous through most of Potrero Valley. In Massacre Canyon, although perennial surface water flow is present during most years, during dryer periods surface water flow becomes

limited to two reaches, 50 to 100 feet in length, along the western portion of the Northern Potrero Creek Area (NPCA). In general, creeks are dry except during and immediately after periods of heavy rainfall. The areas in Potrero and Bedsprings creek where surface water was present were mapped during the Third Quarter 2012 and Fourth Quarter 2012 groundwater monitoring events. The four previously identified fixed locations were checked for flowing water and, if present, the flow rate and volume were determined through field observation and measurements.

2.3 GROUNDWATER AND SURFACE WATER SAMPLING

The frequency of groundwater monitoring depends on the well's classification in the network and intended monitoring purpose. Groundwater is sampled as frequently as quarterly and surface water samples are collected semiannually. Sampling, analytical, and quality assurance/quality control (QA/QC) procedures for the monitoring events are described in the *Programmatic Sampling and Analysis Plan, Lockheed Martin Corporation, Beaumont Sites 1 and 2, Beaumont, California* (Tetra Tech, 2010).

The Third Quarter 2012 monitoring event consisted of water level monitoring, the quarterly sampling of newly installed wells, and storm water sampling. The Fourth Quarter 2012 monitoring event consisted of water level monitoring; surface water sampling; the quarterly sampling of newly installed wells and the semiannual sampling of increasing contaminant trend wells, guard wells, and contaminant attenuation wells. Tables 2-1 and 2-2 list the locations sampled during the Third Quarter 2012 and Fourth Quarter 2012 monitoring events, respectively. Contaminant attenuation parameters include alkalinity, chloride, dissolved organic carbon (DOC), dissolved manganese, ferrous iron, total iron, sulfide, sulfate, nitrate, methane, ethane, ethene, hydrogen, and volatile fatty acids (VFAs). The tables summarize analytical methods, sampling dates, QA/QC samples collected, and field notes.

Surface water samples are collected from up to 17 fixed locations. One designated alternate surface water location (SW-17) is sampled if flowing water is not encountered at the southern end of Massacre Canyon at Gilman Springs Road (SW-16) (Figure 2-1). Certain locations are generally sampled only during or shortly after periods of precipitation, because of the ephemeral nature of the streams on the site.

Table 2-1 Sampling Schedule - Third Quarter 2012

Sample Location	Sample Date	VOCs (1)	1,4-Dioxane (2)	Perchlorate (3)	Comments and QA/QC Samples
MW-110	08/22/12	X	X	X	Sampled with Dedicated Pump, MS/MSD sample
MW-111A	08/21/12	X	X	X	Sampled with FLUTe™ System
MW-111B	08/21/12	X	X	X	Sampled with FLUTe™ System
MW-111C	08/31/12	X	X	X	Sampled with FLUTe™ System
MW-111D	08/21/12	X	X	X	Sampled with FLUTe™ System
MW-111E	08/21/12	X	X	X	Sampled with FLUTe™ System
MW-112A	08/20/12	X	X	X	Sampled with FLUTe™ System, Duplicate MW-112A-Dup
MW-112B	08/20/12	X	X	X	Sampled with FLUTe™ System
MW-112C	08/20/12	X	X	X	Sampled with FLUTe TM System

Total Sample Locations: 9
Total Samples Collected: 9

Notes:

- (1) Volatile organic compounds (VOCs) analyzed by EPA Method SW8260B
- (2) 1,4-Dioxane analyzed by EPA Method SW8270C SIM
- (3) Perchlorate analyzed by EPA Method E332.0

MS/MSD - Matrix Spike/Matrix Spike Duplicate

Table 2-2 Sampling Schedule - Fourth Quarter 2012

Sample Location	Sample Date	VOCs (1)	1,4-Dioxane (2)	Perchlorate (3)	Contaminant Attenuation Parameters (4)	Comments and QA/QC Samples		
SW-01	NA	-	-	-	-	Surface Water - Dry no sample collected		
SW-02	11/15/12	X	X	X	-	Surface Water		
SW-03	11/15/12	X	X	X	-	Surface Water, Duplicate Sample SW-03-Dup		
SW-04	11/15/12	X	X	X	-	Surface Water		
SW-06	NA	-	-	-	-	Surface Water - Dry no sample collected		
SW-07	NA	-	-	-	-	Surface Water - Dry no sample collected		
SW-08	NA	-	-	-	-	Surface Water - Dry no sample collected		
SW-09	11/15/12	X	X	X	-	Surface Water, MS/MSD Sample		
SW-10	NA	-	-	-	-	Surface Water - Dry no sample collected		
SW-11	NA	-	-	-	-	Surface Water - Dry no sample collected		
SW-12	NA	-	-	-	-	Surface Water - Dry no sample collected		
SW-13	NA	-	-	-	-	Surface Water - Dry no sample collected		
SW-14	NA	-	-	-	-	Surface Water - Dry no sample collected		
SW-15	NA	-	-	-	-	Surface Water - Dry no sample collected		
SW-16	NA	-	-	-	-	Surface Water - Dry no sample collected		
SW-17	NA				-	Surface Water - Dry no sample collected		
SW-18	11/15/12	X	X	X	-	Surface Water		
SW-19	NA	-	=	-	-	Surface Water - Dry no sample collected		
F34-TW1	11/26/12	X	X	X	-	Sampled with Peristaltic Pump		
IW-04	11/19/12	X	X	X	-	Sample with Dedicated Pump		
MW-05	11/28/12	X	X	X	X	Sample with Dedicated Pump		
MW-08	11/26/12	X	X	X	X	Sample with Dedicated Pump		
MW-13	11/26/12	X	X	X	X	Sample with Dedicated Pump, MS/MSD Sample		
MW-15	11/16/12	X	X	X	-	Sample with Dedicated Pump		
MW-18	11/16/12	X	X	X	-	Sample with Dedicated Pump, Duplicate Sample MW-18-Dup		
MW-43	11/26/12	X	X	X	X	Sample with Dedicated Pump, Duplicate Sample MW-43-Dup		
MW-48	11/28/12	X	X	X	X	Sample with Dedicated Pump		
MW-60A	11/29/12	X	X	X	-	Sample with Dedicated Pump		
MW-60B	11/29/12	X	X	X	-	Sample with Dedicated Pump		

Notes:

Well not sampled or surface water sample not collected.

- (1) Volatile organic compounds (VOCs) analyzed by EPA Method SW8260B
- (2) 1,4-Dioxane analyzed by EPA Method SW8270C SIM
- (3) Perchlorate analyzed by EPA Method E332.0
- (4) Contaminant attenuation parameters by various methods
- NA Not analyzed
- MS/MSD Matrix Spike/Matrix Spike Duplicate

Table 2-2 Sampling Schedule - Fourth Quarter 2012 (continued)

Sample Location	Sample Date	VOCs (1)	1,4-Dioxane (2)	Perchlorate (3)	Contaminant Attenuation Parameters (4)	Comments and QA/QC Samples
MW-67	11/16/12	X	X	X	-	Sample with Dedicated Pump, MS/MSD Sample
MW-68	11/28/12	X	X	X	-	Sample with Dedicated Pump
MW-70	11/16/12	X	X	X	-	Sample with Dedicated Pump
MW-76B	11/28/12	X	X	X	X	Sample with Dedicated Pump
MW-88	11/19/12	X	X	X	-	Sample with Dedicated Pump
MW-93	11/19/12	X	X	X	-	Sample with Dedicated Pump, Duplicate Sample MW-93-Dup
MW-98B	11/28/12	X	X	X	-	Sample with Dedicated Pump
OW-02	11/28/12	X	X	X	X	Sampled with Peristaltic Pump
P-02	11/26/12	X	X	X	X	Sample with Dedicated Pump
MW-100	11/16/12	X	X	X	-	Sample with Dedicated Pump
MW-101	11/28/12	X	X	X	X	Sample with Dedicated Pump
MW-102	11/28/12	X	X	X	X	Sample with Dedicated Pump
MW-103	11/27/12	X	X	X	X	Sampled with Peristaltic Pump, Duplicate Sample MW-103-Dup
MW-104	11/27/12	X	X	X	X	Sampled with Peristaltic Pump
MW-105	11/27/12	X	X	X	X	Sampled with Peristaltic Pump
MW-106	11/27/12	X	X	X	X	Sampled with Peristaltic Pump
MW-107	11/27/12	X	X	X	X	Sampled with Peristaltic Pump
MW-109	11/27/12	X	X	X	X	Sampled with Peristaltic Pump
MW-110	11/19/12	X	X	X	-	Sample with Dedicated Pump
MW-111A	11/20/12	X	X	X	-	Sampled with FLUTe TM System
MW-111B	11/20/12	X	X	X	-	Sampled with FLUTe™ System
MW-111C	NA	-	-	-	-	FLUTe™ System clogged, unable to sample
MW-111D	11/20/12	X	X	X	-	Sampled with FLUTe TM System
MW-111E	11/20/12	X	X	X	-	Sampled with FLUTe TM System
MW-112A	11/20/12	X	X	X	-	Sampled with FLUTe TM System
MW-112B	11/20/12	X	X	X	-	Sampled with FLUTe TM System
MW-112C	11/20/12	X	X	X	-	Sampled with FLUTe TM System

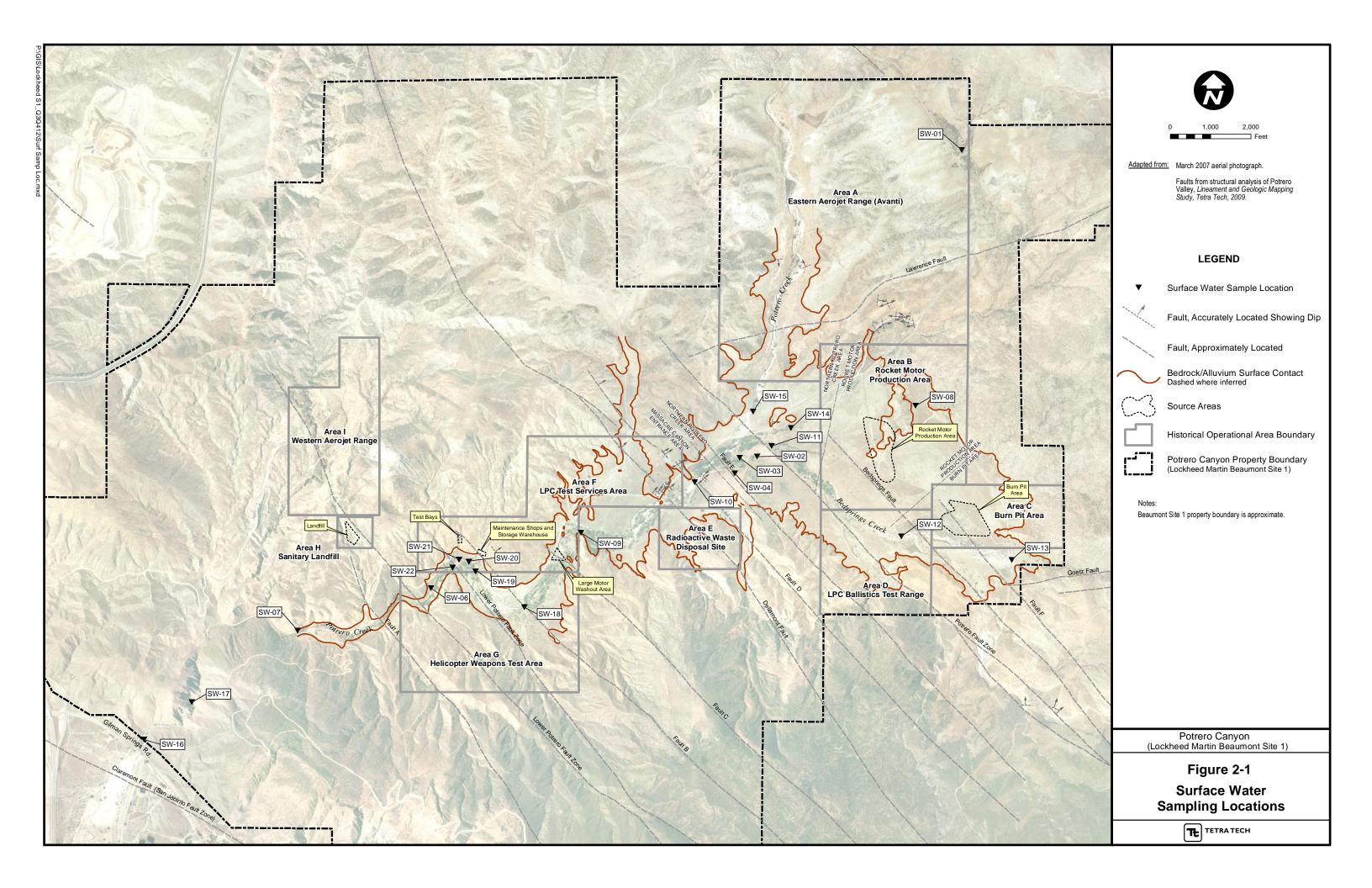
Total Sample Locations: 56 Total Samples Collected: 42

Notes:

Well not sampled or surface water sample not collected.

- (1) Volatile organic compounds (VOCs) analyzed by EPA Method SW8260B
- (2) 1,4-Dioxane analyzed by EPA Method SW8270C SIM
- (3) Perchlorate analyzed by EPA Method E332.0
- (4) Contaminant attenuation parameters by various methods
- NA Not analyzed

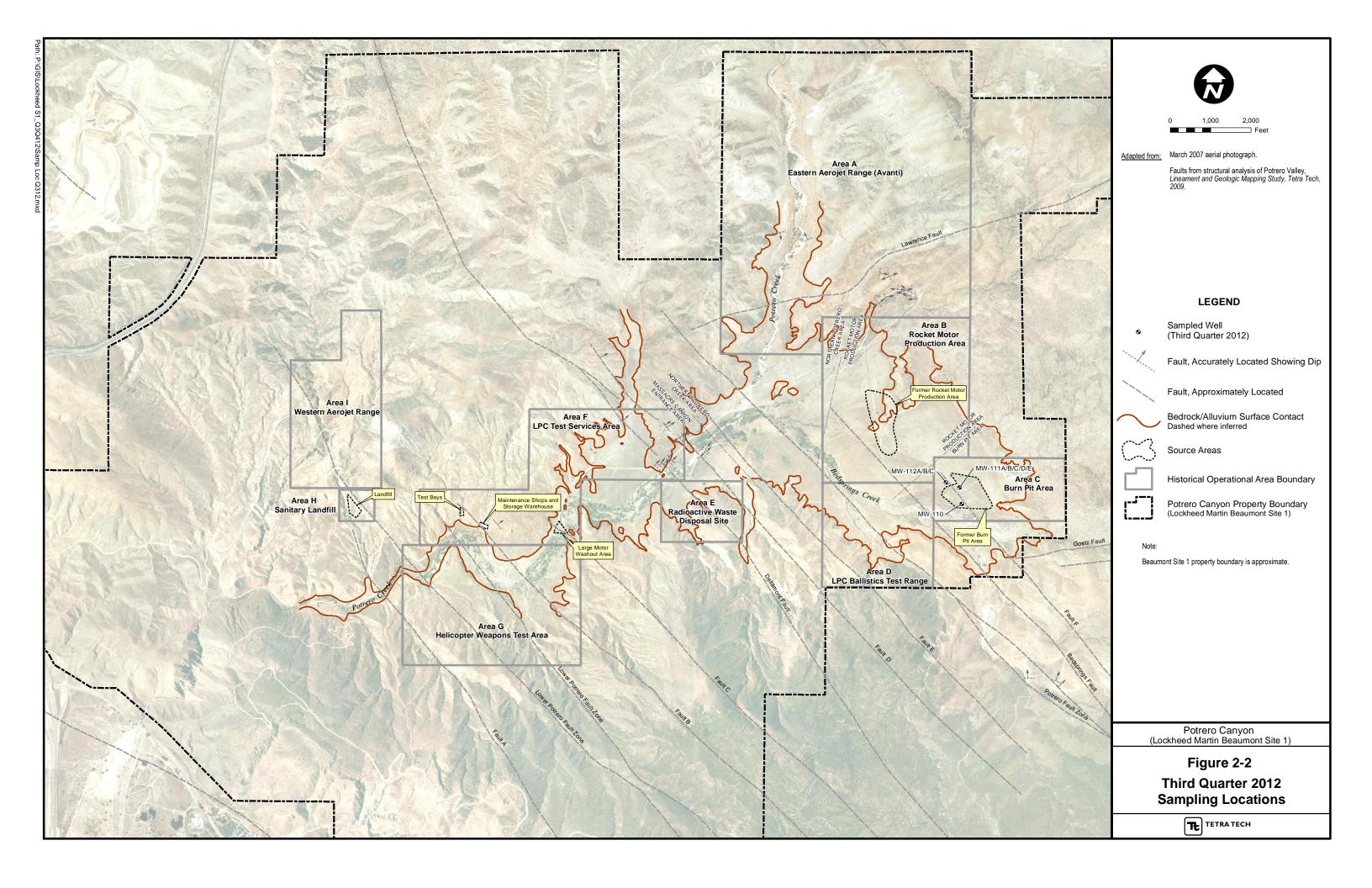
MS/MSD - Matrix Spike/Matrix Spike Duplicate

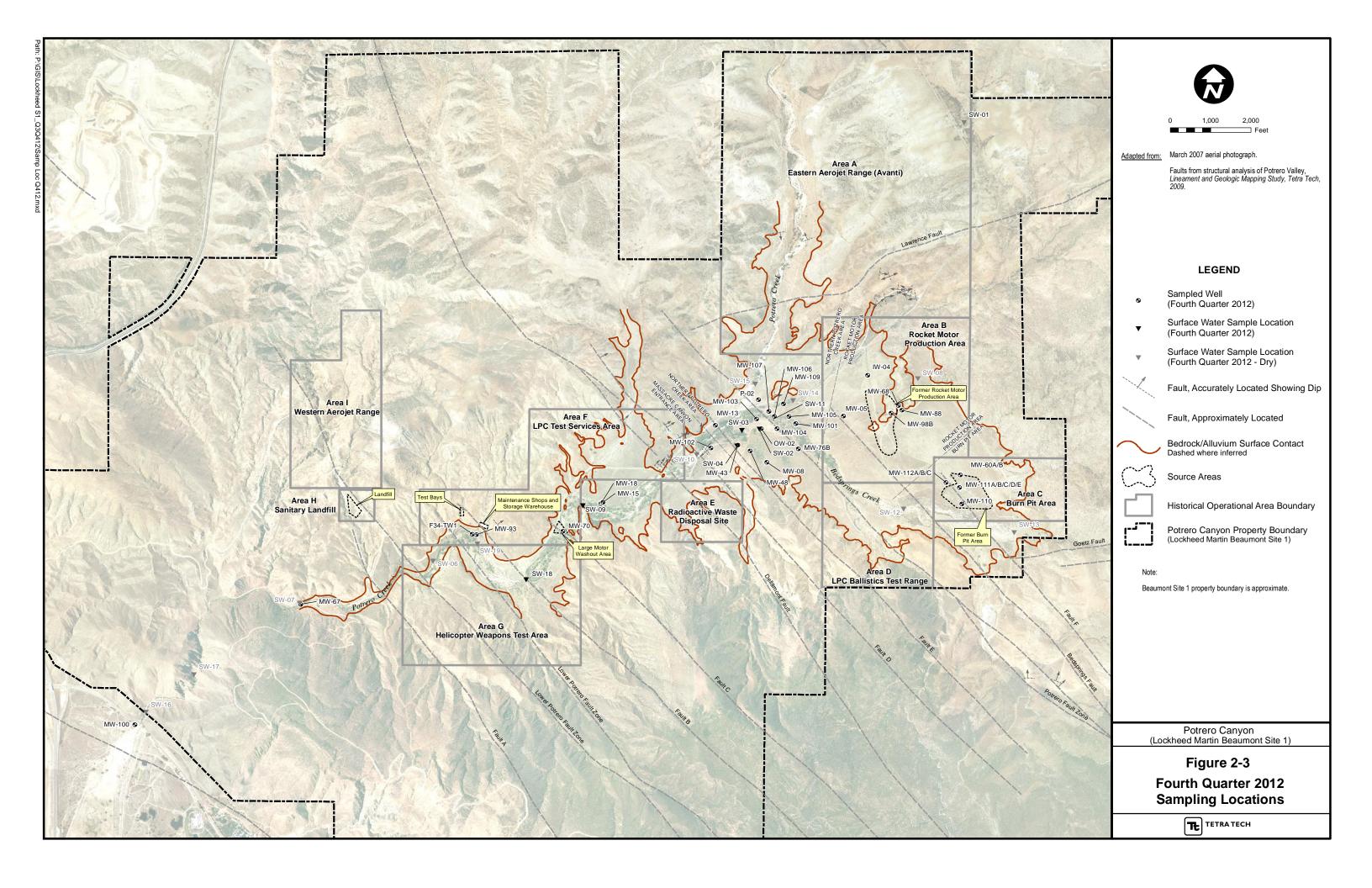


2.3.1 Proposed and Actual Surface Water and Well Locations Sampled

For the Third Quarter 2012 monitoring event, 9 monitoring wells were proposed and sampled for water quality monitoring. Figure 2-2 illustrates groundwater locations sampled for the Third Quarter 2012 monitoring event.

For the Fourth Quarter 2012 monitoring event, 56 sampling locations (17 surface water, one alternate surface water location, and 38 monitoring wells) were proposed for water quality monitoring. Twelve surface water sample locations were not sampled because the locations were dry. The alternate surface water location (SW-17) was also dry and was not sampled. Additionally one monitoring well, MW-111C, was unable to be sampled because its port in the Water FLUTeTM multilevel monitoring system was clogged with sediment. Therefore, water quality data were collected from five surface water, and 37 monitoring well locations during this event. Figure 2-3 illustrates groundwater and surface water locations sampled for the Fourth Quarter 2012 monitoring event.





2.3.2 Field Sampling Procedures

The following water quality field parameters were measured and recorded on field data sheets (Appendix B) during well purging: water level, temperature, pH, electrical conductivity (EC), turbidity, oxidation-reduction potential (ORP), and dissolved oxygen (DO). Groundwater samples were collected from monitoring wells by low-flow purging and sampling through dedicated double-valve pumps, a portable bladder pump, or a peristaltic pump.

Collection of water quality parameters started when at least one discharge hose/pump volume had been removed, and purging was considered complete when the above parameters had stabilized, or the well was purged dry (evacuated). Stabilization of water quality parameters was used as an indication that representative formation water had entered the well and was being purged. The criteria for stabilization of these parameters are as follows: water level \pm 0.1 foot, pH \pm 0.1, EC \pm three percent, turbidity < 10 nephelometric turbidity units (NTUs) (if > 10 NTUs \pm 10%), DO \pm 0.3 milligrams per liter (mg/L), and ORP \pm 10 millivolts. Sampling instruments and equipment were maintained, calibrated, and operated in accordance with the manufacturers' specifications, guidelines, and recommendations. If a well was purged dry, the well was sampled with a disposable bailer after sufficient recharge had taken place to allow sample collection.

Groundwater samples were collected in order of decreasing volatilization potential and placed in appropriate containers. A sample identification label was affixed to each container, and sample custody was maintained by chain-of-custody record. Samples collected were chilled and transported via courier to E.S. Babcock & Sons, a state-accredited analytical laboratory, thus maintaining proper temperatures and sample integrity. Trip blanks were collected for the monitoring events to assess cross-contamination potential of water samples while in transit. Equipment blanks were collected when sampling with non-dedicated equipment to assess cross-contamination potential of water samples via sampling equipment.

Surface water sampling locations were previously located using a global positioning system (GPS) and had been marked in the field. Surface water samples were collected at these GPS-mapped locations either by using a disposable bailer with the sample transferred to the laboratory-supplied water sample containers, or by collecting the water sample directly in the laboratory-supplied water sample container. Temperature, pH, EC, turbidity, ORP, and DO were measured and recorded on field data sheets at surface water sampling locations.

2.4 ANALYTICAL DATA QA/QC

The samples were tested using approved United States Environmental Protection Agency (USEPA) methods. Since the analytical data were obtained by following USEPA-approved method criteria, the data were evaluated by using the USEPA-approved validation methods described in the USEPA Contract Laboratory Program National Functional Guidelines for Organic and Inorganic Superfund Data Review (USEPA, 2008 and 2010). The National Functional Guidelines contain instructions on method-required quality control parameters and on how to interpret these parameters to confer validation to environmental data results.

Quality control parameters used in validating data results include holding times, field blanks, laboratory control samples, method blanks, duplicate environmental samples, spiked samples, and surrogate and spike recovery data.

2.5 HABITAT CONSERVATION

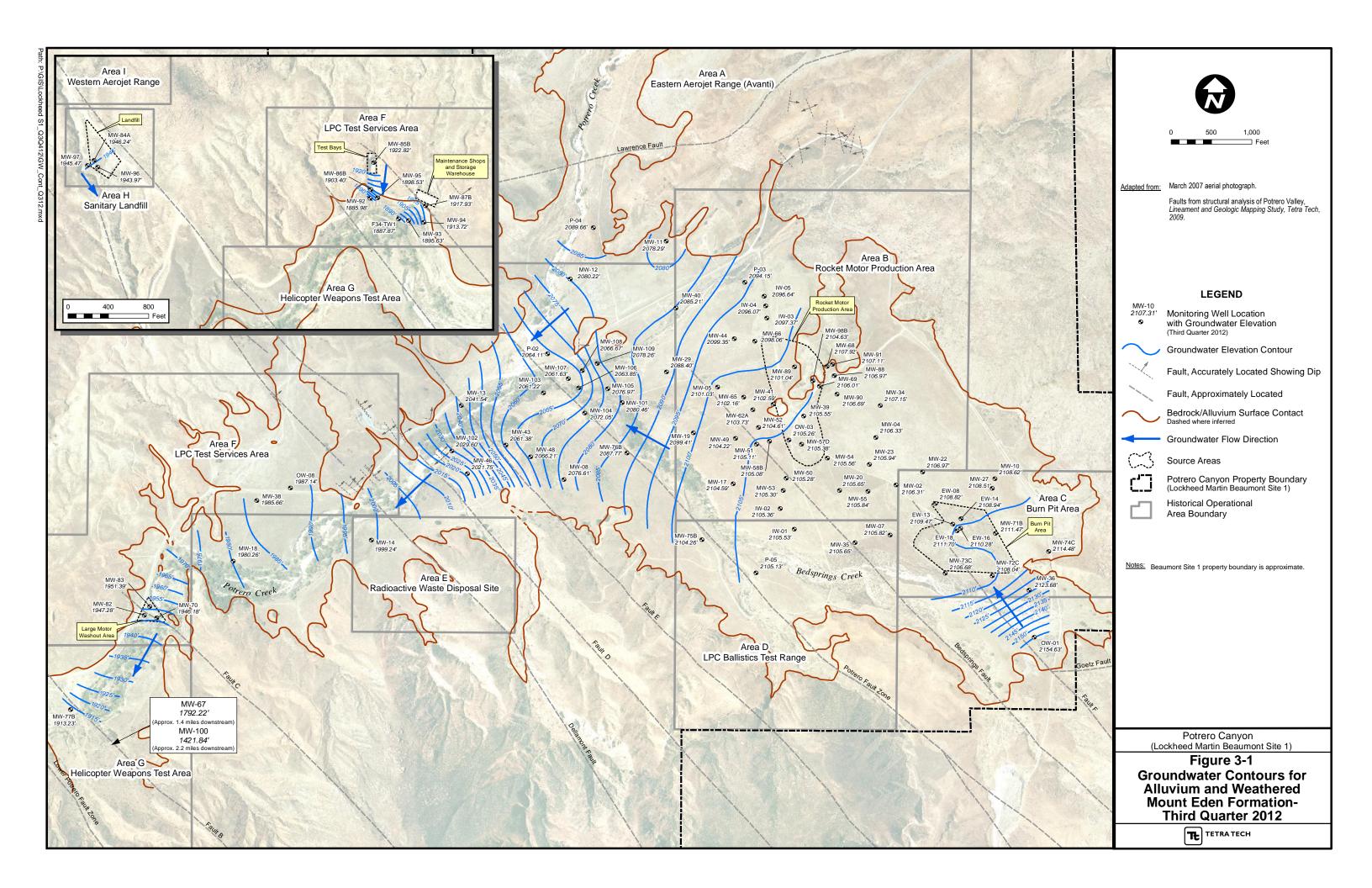
All monitoring activities were performed in accordance with the United States Fish and Wildlife Service (USFWS)-approved Habitat Conservation Plan (HCP) (USFWS, 2005) and subsequent clarifications (Lockheed Martin Corporation, 2006a, 2006b, and 2006c) of the HCP. Groundwater sampling activities were conducted with light-duty vehicles, and were supervised by a USFWS-approved biologist as specified in the Low Effect HCP.

SECTION 3 GROUNDWATER MONITORING RESULTS

Section 3 presents the results and interpretations of the Third Quarter 2012 and Fourth Quarter 2012 groundwater monitoring events. The following subsections include tabulated summaries of groundwater elevation and water quality data, contour maps, and primary chemicals of potential concern results. Plots of groundwater elevation versus time (hydrographs) and concentration versus time (time series graphs) for primary and secondary chemicals of potential concern are presented in Appendices D and E, respectively.

3.1 GROUNDWATER ELEVATION

Groundwater elevations during the Third Quarter 2012 monitoring event ranged from approximately 2,155 feet mean sea level (msl) upgradient of the former Burn Pit Area to approximately 1,792 feet msl in the Massacre Canyon Entrance Area. Groundwater elevations during the Fourth Quarter 2012 monitoring event ranged from approximately 2,153 feet msl upgradient of the former Burn Pit Area to approximately 1,789 feet msl in the Massacre Canyon Entrance Area. A total of 182 monitoring wells were identified for groundwater level measurements for the Third Quarter 2012 and Fourth Quarter 2012 monitoring events. For the Third Quarter 2012 monitoring events, four wells were dry (OW-05, OW-06, OW-07, and VRW-01). For the Fourth Quarter 2012 monitoring events, five wells were dry (OW-05, OW-06, OW-07, P-06S, and VRW-01). EW-01 and EW-02 are former extraction wells with pumps and associated wiring and piping still in place. Whereas these down-hole items make water level measurements difficult, the wells may be needed for future site remediation and/or monitoring activities, and therefore Tetra Tech continues to include them in the monitoring well network. However, if water level measurements cannot be collected from either of these wells, several monitoring wells that are located in close proximity to these former extraction wells can be used for contouring and temporal trends. We will continue to monitor water level elevations from these two former extraction wells when possible. Monitoring wells that have previously been identified as artesian wells are fitted with pressure caps, to prevent groundwater flow onto the ground surface, and pressure gauges for measurement of shut-in head for calculation of static water level. Groundwater elevations for the Third Quarter 2012 and Fourth Quarter 2012 monitoring events from wells screened in the alluvium and weathered Mount Eden formation are shown on Figures 3-1 and 3-2, respectively. A tabulated summary of groundwater elevations for all the wells measured during the Third Quarter 2012 and Fourth Quarter 2012 monitoring events is presented in Table 3-1. Hydrographs for individual wells and for well groups are presented in Appendix D.



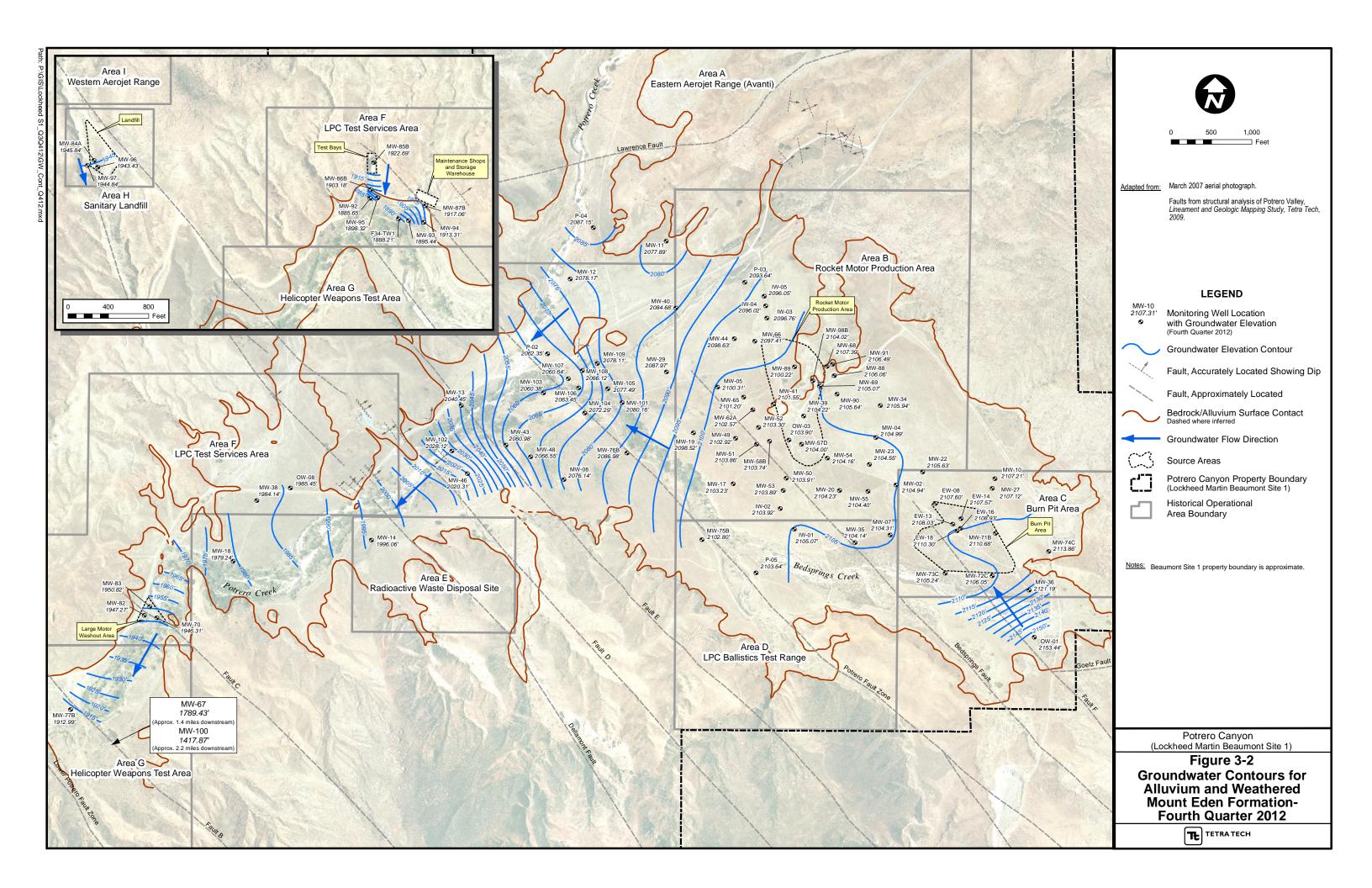


Table 3-1 Groundwater Elevation - Third Quarter 2012 and Fourth Quarter 2012

					Third	Quarter 2012		Fourth Quarter 2012			
Well ID	Site Area	Formation Screened	Measuring Point Elevation (feet msl)	Date Measured	Depth to Water (feet BTOC)	Groundwater Elevation (feet msl)	Groundwater Elevation Change from Second Quarter 2012	Date Measured	Depth to Water (feet BTOC)	Groundwater Elevation (feet msl)	Groundwater Elevation Change from Third Quarter 2012
EW-01 EW-02	RMPA RMPA	QAL QAL	2142.62 2126.15	08/14/12 08/14/12	37.45 23.72	2105.17 2102.43	-1.31 -1.05	11/13/12 11/13/12	38.81 24.80	2103.81 2101.35	-1.36 -1.08
EW-08	BPA	MEF	2178.40	08/14/12	69.60	2108.82	-1.25	11/14/12	70.82	2107.60	-1.22
EW-09 EW-10	BPA BPA	MEF MEF	2179.67 2180.19	08/14/12 08/14/12	70.95 71.30	2108.72 2108.93	-1.26 -1.61	11/14/12 11/14/12	72.38 72.62	2107.29 2107.61	-1.43 -1.32
EW-10 EW-11	BPA	MEF	2180.19	08/14/12	72.00	2108.93	-1.61	11/14/12	73.50	2107.61	-1.32 -1.50
EW-12	BPA	MEF	2183.28	08/14/12	74.34	2108.82	-1.25	11/14/12	75.62	2107.54	-1.28
EW-13 EW-14	BPA BPA	MEF QAL/MEF	2185.57 2184.59	08/14/12 08/14/12	76.10 75.66	2109.47 2108.94	-1.20 -1.11	11/14/12 11/14/12	77.54 77.03	2108.03 2107.57	-1.44 -1.37
EW-14 EW-15	BPA	MEF	2184.10	08/14/12	73.83	2110.00	-1.11	11/14/12	75.21	2107.57	-1.38
EW-16	BPA	MEF	2185.52	08/14/12	75.25	2110.28	-1.25	11/14/12	76.60	2108.93	-1.35
EW-17 EW-18	BPA BPA	MEF MEF	2179.04 2184.98	08/14/12 08/14/12	71.99 73.35	2107.05 2111.70	-1.42 -1.28	11/14/12 11/14/12	73.40 74.75	2105.64 2110.30	-1.41 -1.40
EW-18	MCEA	QAL	2033.89	08/14/12	36.21	1997.68	-3.39	11/14/12	39.10	1994.79	-2.89
EW-20	BPA	MEF	2187.45	08/14/12	75.88	2111.57	-1.45	11/14/12	77.37	2110.08	-1.49
F33-TW2 F33-TW6	NPCA NPCA	QAL QAL	1959.75 1950.62	08/13/12 08/13/12	7.28 7.98	1952.47 1942.64	-1.30 -0.39	11/13/12 11/13/12	6.91 7.80	1952.84 1942.82	0.37 0.18
F33-TW7	NPCA	QAL	NA	08/13/12	9.83	NA	-0.39 NA	11/13/12	9.61	1942.82	NA
F34-TW1	MCEA	QAL	1894.08	08/13/12	6.21	1887.87	-0.79	11/13/12	5.87	1888.21	0.34
IW-01	RMPA	QAL	2160.73	08/14/12	55.20	2105.53	-1.32	11/13/12	55.66	2105.07	-0.46
IW-02 IW-03	RMPA RMPA	QAL QAL	2155.01 2132.86	08/14/12 08/14/12	49.65 35.49	2105.36 2097.37	-1.34 -0.47	11/13/12 11/13/12	51.09 36.10	2103.92 2096.76	-1.44 -0.61
IW-04	RMPA	QAL	2135.09	08/14/12	39.02	2096.07	-1.16	11/13/12	39.07	2096.02	-0.05
IW-05	RMPA	QAL	2136.94	08/14/12	40.30	2096.64	-0.41	11/13/12	40.89	2096.05	-0.59
MW-01 MW-02	RMPA RMPA	MEF MEF	2176.98 2170.10	08/13/12 08/14/12	71.29 63.79	2105.69 2106.31	-1.38 -1.33	11/14/12 11/14/12	72.81 65.16	2104.17 2104.94	-1.52 -1.37
MW-02 MW-03	RMPA	MEF	2170.10	08/14/12	124.27	2045.09	-0.82	11/14/12	125.30	2044.06	-1.03
MW-04	RMPA	QAL	2160.02	08/14/12	53.69	2106.33	-1.35	11/14/12	55.03	2104.99	-1.34
MW-05	RMPA	QAL	2121.40 2121.76	08/14/12 08/14/12	20.37 23.15	2101.03	-0.69	11/13/12	21.09 24.13	2100.31 2097.63	-0.72 -0.98
MW-06 MW-07	RMPA BPA	QAL QAL	2121.76	08/14/12	70.70	2098.61 2105.82	-0.89 -1.38	11/13/12 11/14/12	72.21	2097.63	-0.98 -1.51
MW-08	NPCA	QAL	2090.53	08/15/12	13.92	2076.61	-0.99	11/14/12	14.39	2076.14	-0.47
MW-09	NPCA	QAL	2089.16	08/15/12	0.30	2088.86	-1.30	11/14/12	1.61	2087.55	-1.31
MW-10 MW-11	RMPA NPCA	QAL QAL	2179.40 2122.61	08/13/12 08/13/12	70.78 44.32	2108.62 2078.29	-1.08 -0.08	11/14/12 11/13/12	72.19 44.72	2107.21 2077.89	-1.41 -0.40
MW-12	NPCA	QAL	2098.49	08/13/12	18.27	2080.22	-4.04	11/13/12	20.32	2078.17	-2.05
MW-13	NPCA	QAL	2057.89	08/13/12	16.35	2041.54	-4.53	11/13/12	17.44	2040.45	-1.09
MW-14 MW-15	MCEA MCEA	QAL QAL	2029.67 2009.76	08/15/12 08/13/12	30.43 28.53	1999.24 1981.23	-3.36 -1.88	11/14/12 11/13/12	33.61 29.63	1996.06 1980.13	-3.18 -1.10
MW-17	RMPA	QAL	2140.40	08/14/12	35.81	2104.59	-1.33	11/13/12	37.17	2103.23	-1.36
MW-18	MCEA	QAL	2008.69	08/13/12	28.43	1980.26	-1.80	11/13/12	29.45	1979.24	-1.02
MW-19 MW-20	NPCA RMPA	QAL QAL	2118.49 2162.03	08/14/12 08/14/12	19.08 56.38	2099.41 2105.65	-0.94 -1.37	11/13/12 11/14/12	19.97 57.80	2098.52 2104.23	-0.89 -1.42
MW-22	RMPA	QAL	2173.48	08/14/12	66.51	2105.05	-1.28	11/14/12	67.85	2104.23	-1.34
MW-23	RMPA	QAL	2165.02	08/14/12	59.08	2105.94	-1.38	11/14/12	60.47	2104.55	-1.39
MW-26 MW-27	BPA BPA	MEF	2183.81 2182.73	08/14/12 08/13/12	75.44 74.22	2108.37 2108.51	-1.47 -1.10	11/14/12 11/14/12	76.94 75.61	2106.87 2107.12	-1.50 -1.39
MW-27 MW-28	RMPA	QAL QAL	2162.73	08/13/12	55.31	2105.53	-1.36	11/14/12	56.73	2104.11	-1.42
MW-29	NPCA	MEF	2115.09	08/14/12	26.69	2088.40	-0.48	11/13/12	27.12	2087.97	-0.43
MW-30	RMPA	QAL	2165.01	08/14/12	58.22	2106.79	-1.27	11/14/12	59.56	2105.45	-1.34
MW-31 MW-32	BPA RMPA	Granite Granite	2186.52 2176.61	08/14/12 08/13/12	90.80 84.25	2095.72 2092.36	-1.21 -1.28	11/14/12 11/14/12	92.23 85.69	2094.29 2090.92	-1.43 -1.44
MW-34	RMPA	QAL	2153.80	08/14/12	46.65	2107.15	-1.19	11/14/12	47.86	2105.94	-1.21
MW-35	RMPA	QAL	2170.98	08/13/12	65.33	2105.65	-1.35	11/14/12	66.84	2104.14	-1.51
MW-36 MW-38	UG MCEA	QAL MEF	2205.18 2030.29	08/13/12 08/13/12	81.50 44.63	2123.68 1985.66	-2.08 -1.25	11/14/12 11/13/12	83.99 46.15	2121.19 1984.14	-2.49 -1.52
MW-39	RMPA	QAL	2144.18	08/14/12	38.63	2105.55	-1.36	11/14/12	39.96	2104.22	-1.33
MW-40	NPCA	MEF	2126.39	08/13/12	41.18	2085.21	-0.40	11/13/12	41.71	2084.68	-0.53
MW-41 MW-43	RMPA NPCA	MEF QAL	2133.95 2068.58	08/15/12 08/15/12	31.36 7.20	2102.59 2061.38	-1.06 -2.10	11/14/12 11/14/12	32.40 7.60	2101.55 2060.98	-1.04 -0.40
MW-43 MW-44	NPCA	QAL	2128.69	08/13/12	29.34	2099.35	-0.64	11/14/12	30.06	2098.63	-0.40
MW-45	MCEA	QAL	2068.18	08/15/12	4.0 PSI	2077.42	-0.69	11/14/12	3.6 PSI	2076.49	-0.92
MW-46 MW-47	MCEA NPCA	QAL QAL	2072.17 2076.67	08/15/12 08/15/12	50.42 3.4 PSI	2021.75 2084.52	-1.66 -1.39	11/14/12 11/14/12	51.86 3.0 PSI	2020.31 2083.60	-1.44 -0.92
MW-47 MW-48	NPCA NPCA	QAL	2076.67	08/15/12	10.23	2084.52	-1.87	11/14/12	9.89	2083.60	0.34
MW-49	RMPA	QAL	2130.92	08/14/12	26.70	2104.22	-1.27	11/13/12	28.00	2102.92	-1.30
MW-50 MW-51	RMPA RMPA	QAL QAL	2151.43 2138.36	08/14/12 08/14/12	46.15 33.25	2105.28 2105.11	-1.33 -1.21	11/13/12 11/13/12	47.52 34.50	2103.91 2103.86	-1.37 -1.25
MW-51 MW-52	RMPA	QAL	2138.36	08/14/12	33.25	2105.11	-1.25 -1.25	11/13/12	32.88	2103.86	-1.25 -1.31
MW-53	RMPA	QAL	2153.29	08/14/12	47.99	2105.30	-1.33	11/13/12	49.40	2103.89	-1.41
MW-54	RMPA	QAL	2153.44	08/14/12	47.88	2105.56	-1.39	11/14/12	49.28	2104.16	-1.40
MW-55 MW-56A	RMPA RMPA	QAL MEF	2166.66 2143.09	08/14/12 08/14/12	60.82 49.94	2105.84 2093.15	-1.33 -1.20	11/14/12 11/13/12	62.26 51.19	2104.40 2091.90	-1.44 -1.25
MW-56B	RMPA	QAL	2142.58	08/14/12	37.37	2105.21	-1.34	11/13/12	38.75	2103.83	-1.38
MW-56C	RMPA	QAL	2142.77	08/14/12	37.65	2105.12	-1.29	11/13/12	39.01	2103.76	-1.36
MW-56D MW-57A	RMPA RMPA	QAL QAL	2142.48 2145.98	08/14/12 08/14/12	37.24 40.56	2105.24 2105.42	-1.34 -1.31	11/13/12 11/13/12	38.61 41.95	2103.87 2104.03	-1.37 -1.39
MW-57B	RMPA	QAL	2146.19	08/14/12	40.76	2105.43	-1.31	11/13/12	42.15	2104.04	-1.39
MW-57C	RMPA	QAL	2146.02	08/14/12	40.58	2105.44	-1.32	11/13/12	41.98	2104.04	-1.40
MW-57D MW-58A	RMPA RMPA	QAL QAL	2146.10 2140.73	08/14/12 08/14/12	40.72 35.85	2105.38 2104.88	-1.31 -1.30	11/13/12 11/13/12	42.10 37.22	2104.00 2103.51	-1.38 -1.37
MW-58B	RMPA	QAL	2140.73	08/14/12	35.70	2105.08	-1.29	11/13/12	37.04	2103.74	-1.34
MW-58C	RMPA	QAL	2141.02	08/14/12	36.06	2104.96	-1.27	11/13/12	37.41	2103.61	-1.35
MW-58D	RMPA	QAL	2140.94	08/14/12	36.07	2104.87	-1.31	11/13/12	37.45	2103.49	-1.38
MW-59A MW-59B	BPA BPA	MEF MEF	2180.14 2180.39	08/14/12 08/14/12	76.72 72.11	2103.42 2108.28	-1.27 -1.26	11/14/12 11/14/12	78.16 73.55	2101.98 2106.84	-1.44 -1.44
MW-59C	BPA	MEF	2179.93	08/14/12	73.90	2106.03	-1.29	11/14/12	75.36	2104.57	-1.46
MW-59D	BPA	MEF	2180.53	08/14/12	73.88	2106.65	-1.41	11/14/12	75.31	2105.22	-1.43
MW-60A MW-60B	BPA BPA	MEF MEF	2182.59 2182.77	08/14/12 08/14/12	76.32 74.85	2106.27 2107.92	-1.18 -1.21	11/14/12 11/14/12	77.70 76.21	2104.89 2106.56	-1.38 -1.36
Notes:	BPA -	Burn Pit Are	a		DG -	Downgradient		"_"	Formation screen	ned not defined	1.50
	MCEA -		nyon Entrance		BTOC -	Below top of cas	sing	QAL -	Quaternary alluv	ium	
	NPCA - Northern Potrero Creek Area msl RMPA - Rocket Motor Production Area NA					Mean sea level Not available		QAL/MEF - MEF -	Quaternary alluv Mount Eden forn	ium/Mount Eden fo	ormation

RMPA - Rocket Motor Production Area
UG - Upgradient NA - Not available PSI - Pounds per square inch MEF - Mount Eden formation

Table 3-1 Groundwater Elevation - Third Quarter 2012 and Fourth Quarter 2012 (continued)

					Third	Quarter 2012		Fourth Quarter 2012			
Well ID	Site Area	Formation Screened	Measuring Point Elevation (feet msl)	Date Measured	Depth to Water (feet BTOC)	Groundwater Elevation (feet msl)	Groundwater Elevation Change from Second Quarter 2012	Date Measured	Depth to Water (feet BTOC)	Groundwater Elevation (feet msl)	Groundwater Elevation Change from Third Quarter 2012
MW-61A	BPA	MEF	2186.95	08/14/12	84.09	2102.86	-4.14	11/14/12	85.58	2101.37	-1.49
MW-61B MW-61C	BPA BPA	MEF MEF	2186.77 2186.84	08/14/12 08/14/12	75.79 81.76	2110.98 2105.08	-1.30 -1.27	11/14/12 11/14/12	77.16 83.23	2109.61 2103.61	-1.37 -1.47
MW-61D	BPA	MEF	2186.83	08/14/12	79.11	2107.72	-1.27	11/14/12	80.54	2106.29	-1.43
MW-62A	RMPA	QAL	2131.32	08/14/12	27.59	2103.73	-1.07	11/13/12	28.75	2102.57	-1.16
MW-62B MW-63	RMPA RMPA	QAL QAL	2131.49 2156.20	08/14/12 08/14/12	27.67 50.58	2103.82 2105.62	-1.21 -1.38	11/13/12 11/14/12	28.91 52.00	2102.58 2104.20	-1.24 -1.42
MW-64	RMPA	QAL	2128.41	08/14/12	26.08	2102.33	-0.88	11/13/12	27.02	2101.39	-0.94
MW-65	RMPA	QAL	2128.92	08/14/12	26.76	2102.16	-0.89	11/13/12	27.72	2101.20	-0.96
MW-66 MW-67	RMPA MCEA	QAL QAL	2130.43 1799.54	08/14/12 08/13/12	32.37 7.32	2098.06 1792.22	-0.55 -2.57	11/13/12 11/13/12	33.02 10.11	2097.41 1789.43	-0.65 -2.79
MW-68	RMPA	QAL	2144.69	08/13/12	36.77	2107.92	-0.60	11/13/12	37.30	2107.39	-0.53
MW-69	RMPA	QAL	2143.26	08/15/12	37.25	2106.01	-0.78	11/14/12	38.19	2105.07	-0.94
MW-70	MCEA BPA	QAL	1976.15	08/13/12	29.97	1946.18	-1.00	11/13/12	29.84	1946.31	0.13
MW-71A MW-71B	BPA BPA	Granite QAL/MEF	2193.77 2194.01	08/14/12 08/14/12	157.25 82.54	2036.52 2111.47	-0.53 -0.69	11/14/12 11/14/12	158.07 83.33	2035.70 2110.68	-0.82 -0.79
MW-71C	BPA	MEF	2193.87	08/14/12	84.65	2109.22	-1.12	11/14/12	85.87	2108.00	-1.22
MW-72A	BPA	Granite	2199.06	08/13/12	96.78	2102.28	-2.18	11/14/12	98.90	2100.16	-2.12
MW-72B MW-72C	BPA BPA	MEF QAL	2199.22 2199.35	08/13/12 08/13/12	91.23 91.31	2107.99 2108.04	-1.87 -1.87	11/14/12 11/14/12	93.21 93.30	2106.01 2106.05	-1.98 -1.99
MW-73A	BPA	MEF	2189.39	08/13/12	110.16	2079.23	-1.38	11/14/12	111.57	2077.82	-1.41
MW-73B	BPA	MEF	2189.48	08/13/12	94.83	2094.65	-1.43	11/14/12	96.34	2093.14	-1.51
MW-73C MW-74A	BPA UG	QAL Granite	2189.65 2199.66	08/13/12 08/13/12	82.97 158.46	2106.68 2041.20	-1.14 -0.31	11/14/12 11/14/12	84.41 159.00	2105.24 2040.66	-1.44 -0.54
MW-74A MW-74B	UG	Granite	2199.00	08/13/12	115.95	2083.86	-1.37	11/14/12	115.85	2083.96	0.10
MW-74C	UG	MEF	2199.96	08/13/12	85.48	2114.48	-0.54	11/14/12	86.10	2113.86	-0.62
MW-75A	RMPA	MEF	2149.44	08/15/12	54.23	2095.21	-1.28	11/14/12	55.58	2093.86	-1.35
MW-75B MW-75C	RMPA RMPA	QAL QAL	2149.51 2150.02	08/15/12 08/15/12	45.25 45.83	2104.26 2104.19	-1.37 -1.34	11/14/12 11/14/12	46.71 47.25	2102.80 2102.77	-1.46 -1.42
MW-76A	NPCA	MEF	2105.91	08/15/12	22.90	2083.01	-1.32	11/14/12	24.23	2081.68	-1.33
MW-76B	NPCA	QAL	2105.40	08/15/12	17.63	2087.77	-1.71	11/14/12	18.42	2086.98	-0.79
MW-76C MW-77A	NPCA MCEA	QAL MEF	2106.29 1930.62	08/15/12 08/13/12	8.03 14.07	2098.26 1916.55	-1.27 -1.82	11/14/12 11/13/12	9.27 14.78	2097.02 1915.84	-1.24 -0.71
MW-77B	MCEA	MEF	1930.88	08/13/12	17.65	1913.23	-0.74	11/13/12	17.89	1912.99	-0.71
MW-78	BPA	MEF	2182.63	08/14/12	86.43	2096.20	-1.23	11/14/12	87.84	2094.79	-1.41
MW-79A	RMPA	MEF	2142.00	08/14/12 08/14/12	41.03	2100.97	-1.33	11/13/12	42.37	2099.63	-1.34
MW-79C MW-80	RMPA NPCA	QAL MEF	2142.07 2070.47	08/14/12	37.93 0.45 PSI	2104.14 2071.51	-1.34 -1.89	11/13/12 11/14/12	39.30 1.94	2102.77 2068.53	-1.37 -2.98
MW-81	MCEA	MEF	2010.72	08/13/12	29.82	1980.90	-1.85	11/13/12	30.92	1979.80	-1.10
MW-82	NPCA	QAL	1974.17	08/13/12	26.89	1947.28	-0.69	11/13/12	26.90	1947.27	-0.01
MW-83 MW-84A	NPCA MCEA	QAL MEF	1976.93 2,010.02	08/13/12 08/13/12	25.54 63.78	1951.39 1946.24	-1.38 -0.25	11/13/12 11/13/12	26.11 64.18	1950.82 1945.84	-0.57 -0.40
MW-84B	MCEA	MEF	2,011.19	08/13/12	65.86	1945.33	-0.18	11/13/12	66.22	1944.97	-0.36
MW-85A	MCEA	MEF	1,929.31	08/13/12	7.50	1921.81	-1.07	11/13/12	8.04	1921.27	-0.54
MW-85B MW-86A	MCEA MCEA	MEF MEF	1,928.74 1,923.21	08/14/12 08/13/12	5.92 16.66	1922.82 1906.55	-2.35 -0.99	11/13/12 11/13/12	6.05 17.07	1922.69 1906.14	-0.13 -0.41
MW-86B	MCEA	QAL/MEF	1,923.21	08/13/12	19.81	1903.40	-0.72	11/13/12	20.03	1903.18	-0.22
MW-87A	MCEA	MEF	1,938.92	08/13/12	22.10	1916.82	-0.84	11/13/12	22.71	1916.21	-0.61
MW-87B MW-88	MCEA RMPA	MEF QAL	1,938.82 2,141.97	08/13/12 08/14/12	20.89 35.00	1917.93 2106.97	-0.84 -0.87	11/13/12 11/14/12	21.76 35.91	1917.06 2106.06	-0.87 -0.91
MW-89	RMPA	QAL	2,130.82	08/14/12	29.78	2100.97	-0.74	11/14/12	30.60	2100.00	-0.82
MW-90	RMPA	QAL	2,147.71	08/14/12	41.02	2106.69	-1.02	11/14/12	42.07	2105.64	-1.05
MW-91 MW-92	RMPA MCEA	MEF MEF	2,144.85 1,919.83	08/14/12 08/13/12	37.74 33.85	2107.11 1885.98	-0.71 -1.30	11/14/12 11/13/12	38.36 34.18	2106.49 1885.65	-0.62 -0.33
MW-92 MW-93	MCEA	MEF	1,919.83	08/13/12	35.84	1895.63	-0.99	11/13/12	36.03	1895.44	-0.33
MW-94	MCEA	MEF	1,936.55	08/13/12	22.83	1913.72	-0.56	11/13/12	23.24	1913.31	-0.41
MW-95	MCEA	MEF	1,920.80	08/13/12	22.27	1898.53	-0.62	11/13/12	22.48	1898.32	-0.21
MW-96 MW-97	MCEA MCEA	MEF MEF	1998.63 1996.47	08/13/12 08/13/12	54.66 51.00	1943.97 1945.47	-0.29 -1.02	11/13/12 11/13/12	55.20 51.63	1943.43 1944.84	-0.54 -0.63
MW-98A	RMPA	MEF	2141.68	08/15/12	44.19	2097.49	-0.96	11/14/12	45.30	2096.38	-1.11
MW-98B	RMPA	MEF	2141.73	08/15/12	37.10	2104.63	-0.67	11/14/12	37.71	2104.02	-0.61
MW-99 MW-100	RMPA DG	MEF Granite	2144.63 1525.79	08/14/12 08/14/12	55.53 103.81	2089.10 1421.84	-0.52 -3.43	11/14/12 11/13/12	56.19 107.78	2088.44 1417.87	-0.66 -3.97
MW-100	NPCA	QAL	2095.90	08/14/12	15.44	2080.46	-3.43	11/13/12	15.74	2080.16	-0.30
MW-102	MCEA	QAL	2067.21	08/15/12	37.61	2029.60	-3.59	11/14/12	39.09	2028.12	-1.48
MW-103 MW-104	NPCA NPCA	QAL QAL	2075.88 2087.47	08/15/12 08/15/12	14.66 15.42	2061.22 2072.05	-0.69 -1.57	11/14/12 11/14/12	15.50 15.18	2060.38 2072.29	-0.84 0.24
MW-104 MW-105	NPCA NPCA	QAL	2087.47	08/15/12	15.42	2072.03	-1.43	11/14/12	14.74	2072.29	0.52
MW-106	NPCA	QAL	2085.25	08/15/12	21.40	2063.85	-3.38	11/14/12	21.80	2063.45	-0.40
MW-107 MW-108	NPCA NPCA	QAL QA/MEF	2084.84 2087.22	08/15/12 08/15/12	23.21 20.55	2061.63 2066.67	-3.10 -3.74	11/14/12 11/14/12	24.20 21.10	2060.64 2066.12	-0.99 -0.55
MW-108 MW-109	NPCA NPCA	QA/MEF QA/MEF	2087.22	08/15/12	14.60	2066.67	-3.74 -1.58	11/14/12	14.75	2066.12	-0.55 -0.15
MW-110	BPA	QAL	2188.54	08/14/12	95.53	2093.01	-1.70	11/14/12	97.34	2091.20	-1.81
OW-01	BPA	QAL	2204.62	08/13/12	49.99	2154.63	-1.23	11/14/12	51.18	2153.44	-1.19
OW-02 OW-03	NPCA RMPA	QAL QAL	2078.97 2143.65	08/15/12 08/14/12	2.54 38.39	2076.43 2105.26	-0.25 -1.33	11/14/12 11/13/12	3.13 39.75	2075.84 2103.90	-0.59 -1.36
OW-05	NPCA	QAL	2160.85	08/13/12	Dry	Dry Well	NA	11/13/12	Dry	Dry Well	NA
OW-06	MCEA	QAL	2084.67	08/13/12	Dry	Dry Well	NA	11/13/12	Dry	Dry Well	NA
OW-07 OW-08	MCEA MCEA	QAL QAL	2108.06 2036.33	08/13/12 08/13/12	Dry 49.19	Dry Well 1987.14	NA -1.55	11/13/12 11/13/12	Dry 50.88	Dry Well 1985.45	NA -1.69
P-02	NPCA	QAL	2030.33	08/13/12	17.04	2064.11	-2.76	11/13/12	18.80	2062.35	-1.76
P-03	NPCA	QAL	2140.25	08/14/12	46.10	2094.15	-0.30	11/13/12	46.61	2093.64	-0.51
P-04 P-05	NPCA RMPA	QAL	2112.63 2162.20	08/13/12	22.97	2089.66 2105.13	-3.96 1.38	11/13/12 11/14/12	25.48	2087.15	-2.51
P-05 P-06S	MCEA	QAL QAL	2162.20 2034.44	08/15/12 08/15/12	57.07 36.53	1997.91	-1.38 -4.14	11/14/12	58.56 Dry	2103.64 Dry Well	-1.49 NA
P-06D	MCEA	QAL	2034.41	08/15/12	35.49	1998.92	-2.30	11/14/12	39.48	1994.93	-3.99
P-07	MCEA	QAL	2034.60	08/15/12	37.11	1997.49	-3.33	11/14/12	39.99	1994.61	-2.88
P-08 P-09	MCEA BPA	QAL MEF	2030.87 2187.38	08/15/12 08/14/12	32.82 76.10	1998.05 2111.28	-3.37 -1.43	11/14/12 11/14/12	35.75 77.53	1995.12 2109.85	-2.93 -1.43
VRW-01	BPA	QAL	2187.35	08/14/12	Dry	Dry Well	-1.45 NA	11/14/12	Dry	Dry Well	NA
VRW-02	BPA	QAL	2181.66	08/14/12	75.56	2106.10	-1.43	11/14/12	77.06	2104.60	-1.50
VRW-03 Notes:	BPA BPA -	MEF Burn Pit Area	2184.32	08/14/12	70.67 DG -	2113.65 Downgradient	-0.91	11/14/12	71.66 Formation screen	2112.66	-0.99
riotes:	MCEA -		a nyon Entrance	Area	BTOC -	Below top of cas	sing	OAL -	Ouaternary alluvi		

MCEA - Massacre Canyon Entrance Area NPCA - Northern Potrero Creek Area RMPA - Rocket Motor Production Area UG - Upgradient

BTOC - Below top of casing msl - Mean sea level NA -Not available

PSI -

QAL - Quaternary alluvium

QAL/MEF - Quaternary alluvium/Mount Eden formation

MEF - Mount Eden formation

Pounds per square inch

During Third Quarter 2012, the Beaumont National Weather Service (NWS) station reported approximately 0.30 inches of precipitation, and the average site-wide groundwater elevation decreased approximately 1.39 feet. During Fourth Quarter 2012, the Beaumont NWS station reported approximately 3.62 inches of precipitation and the average site-wide groundwater elevation decreased approximately 1.15 feet. Generally the groundwater elevations in site wells show a one-season lag before responding to seasonal precipitation. Table 3-2 presents the range and average change in groundwater elevation by area. Figures 3-3 and 3-4, respectively, present elevation differences between the Second Quarter 2012 and Third Quarter 2012, and between the Third Quarter 2012 and Fourth Quarter 2012 groundwater monitoring events.

Table 3-2 Groundwater Elevation Change - Third Quarter 2012 and Fourth Quarter 2012

Site Area	Range of Ground Change - Third Q	dwater Elevation Juarter 2012 (feet)	Average Change By Area (feet)	Range of Groundwa Change - Fourth Qua	Average Change By Area (feet)	
BPA	-4.14	-0.53	-1.38	-2.12	-0.79	-1.42
MCEA	-4.14	-0.18	-1.61	-3.99	0.34	-1.10
NPCA	-4.53	-0.08	-1.67	-2.98	0.52	-0.73
RMPA	-1.39	-0.41	-1.15	-1.52	-0.05	-1.18

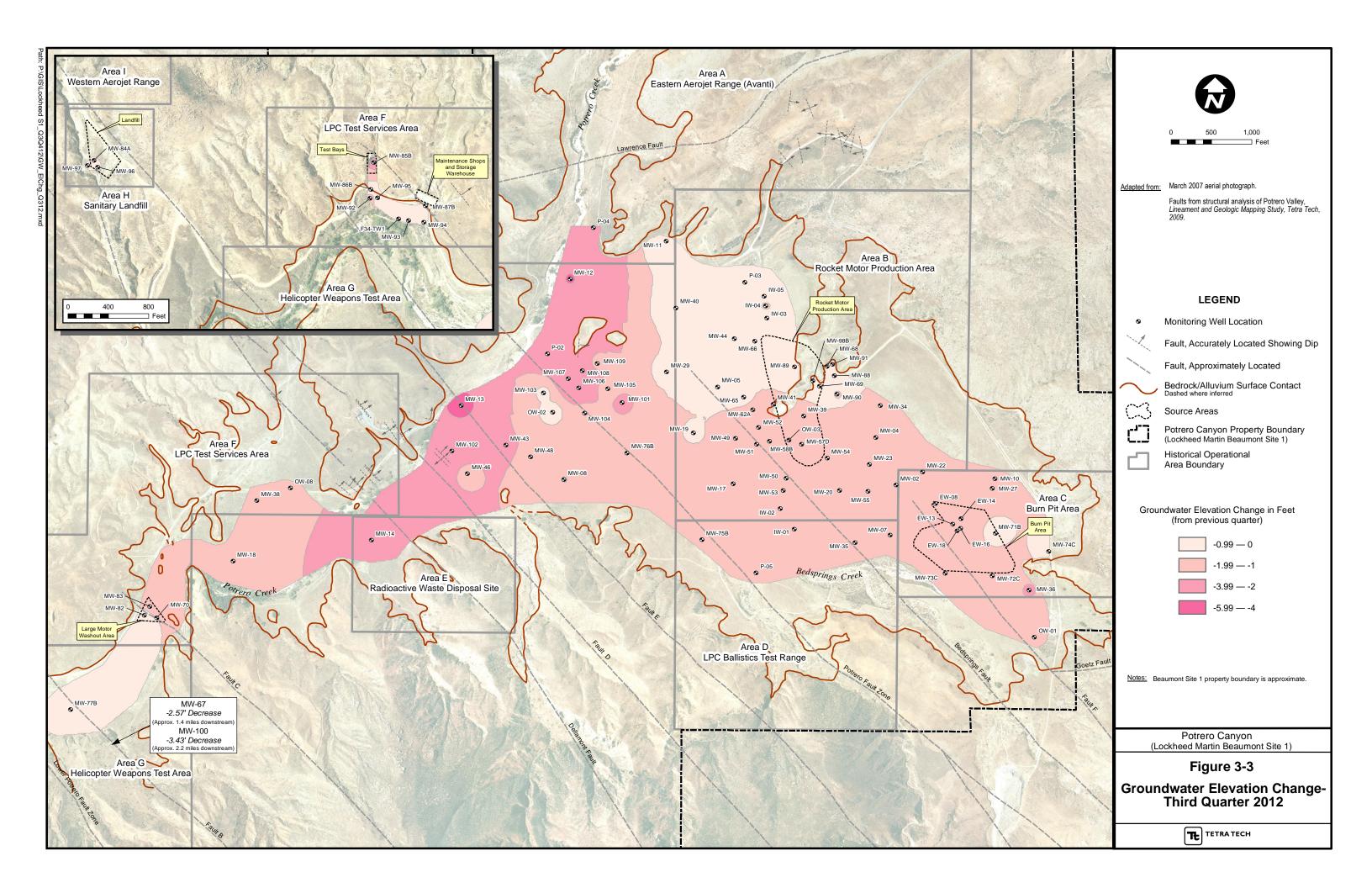
Notes:

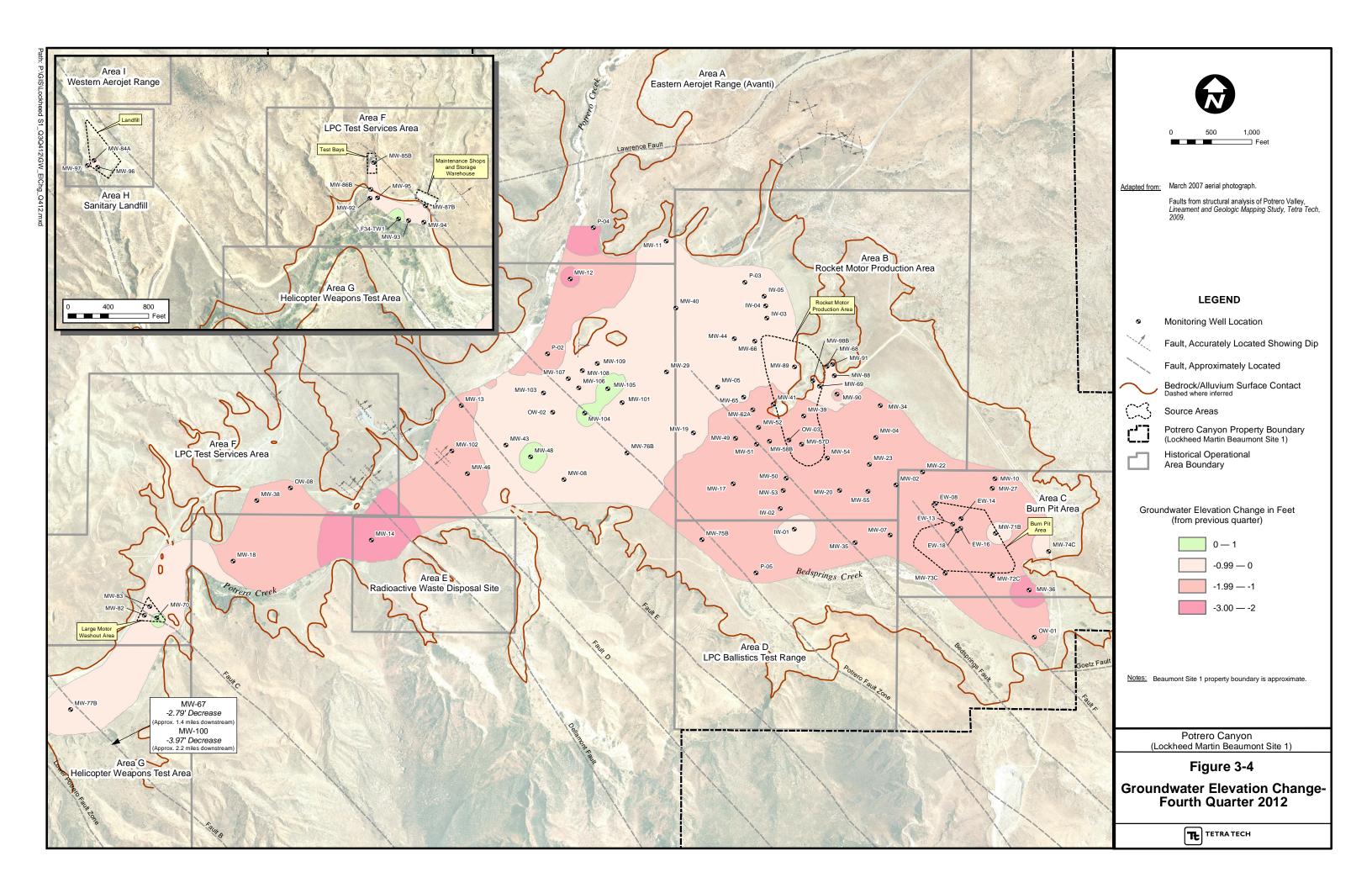
BPA - Burn Pit Area

MCEA - Massacre Canyon Entrance Area

NPCA - Northern Potrero Creek Area RMPA - Rocket Motor Production Area

Groundwater elevations and seasonal responses to changes in recharge for select shallow and deeper wells are shown on Figures 3-5 through 3-7. The selected wells represent a groundwater flow path from upgradient of the former Burn Pit Area (BPA), through the former BPA, through the former Rocket Motor Production Area (RMPA), and southwestward (downgradient) through the Northern Potrero Creek Area (NPCA) and Massacre Canyon Entrance Area (MCEA). Groundwater elevations in shallow wells (alluvium and shallow Mount Eden formation (MEF)) upgradient of the BPA and at the BPA show a rapid and significant response to rainfall, with a more dampened response observed farther out in the valley through the RMPA, NPCA, and MCEA (Figures 3-5 and 3-7). The deeper MEF and granitic/metasedimentary bedrock wells show a response very similar to the shallow wells during the periods of increased precipitation (Figure 3-6).





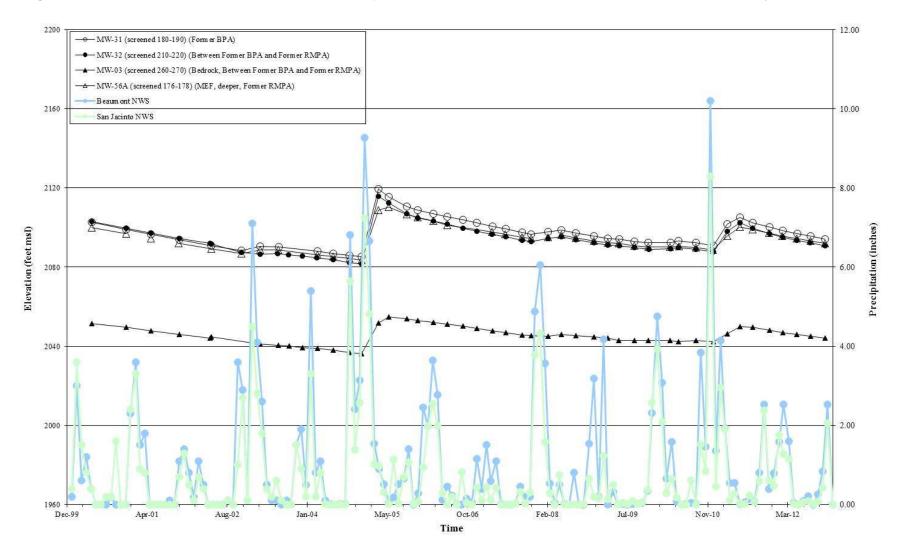
MW-36 (screened 85-105) (Upgradient of Form er BPA) * MW-61D (screened 114-116) (Former BPA) △ MW-56C (screened 48-58) (Former RMPA) → MW-43 (screened 8.5-23.5) (Northern Potrero Creek Area) -MW-18 (screened 30-50) (QAL, shallower, Massacre Canyon Entrance Area) 10 — MW-14 (screened 29-49) (Upgradient of Massacre Canyon Entrance Area) Beaum ont NWS San Jacinto NWS 2120 Precipitation (inches) Elevation (feet msl) 2040 2000 BE BE BB Dec-99 Apr-01 Aug-02 Jan-04 May-05 Oct-06 Feb-08 Jul-09 Nov-10 Mar-12

Figure 3-5 Groundwater Elevations vs. Time - Selected Alluvial and Shallow Mount Eden Formation Wells

Notes: BPA – Burn Pit Area msl – mean sea level NWS - National Weather Service QAL – Quaternary alluvium RMPA – Rocket Motor Production Area

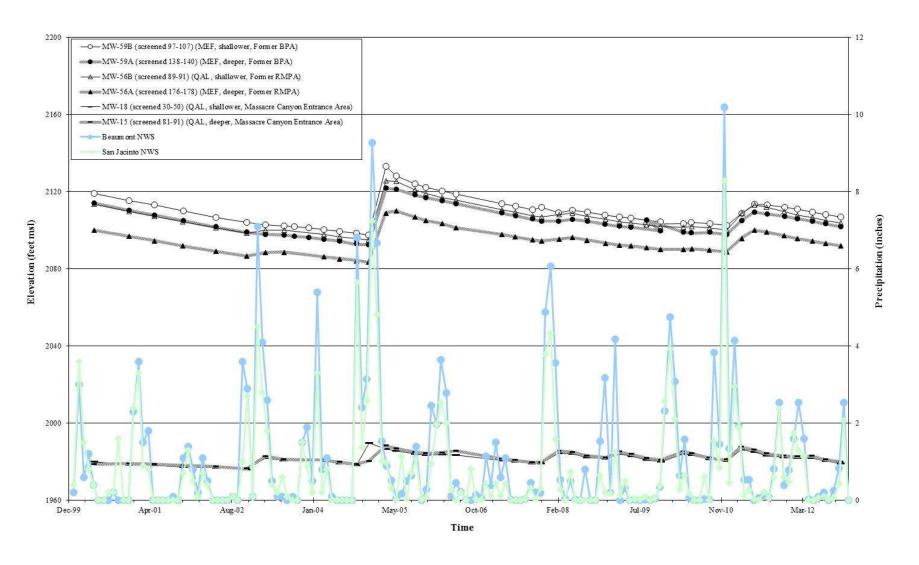
Time

Figure 3-6 Groundwater Elevations vs. Time - Deeper Mount Eden Formation and Granitic/Metasedimentary Bedrock Wells



Notes: BPA – Burn Pit Area msl – mean sea level MEF – Mount Eden formation NWS - National Weather Service RMPA – Rocket Motor Production Area

Figure 3-7 Groundwater Elevations Comparison - Selected Shallower and Deeper Screened Wells in the Alluvium and Shallow Mount Eden Formation



Notes: BPA – Burn Pit Area msl – mean sea level MEF – Mount Eden formation NWS - National Weather Service QAL – Quaternary alluvium RMPA – Rocket Motor Production Area

3.2 SURFACE WATER FLOW

During Third Quarter 2012 and Fourth Quarter 2012, Tetra Tech personnel walked the Potrero and Bedsprings creek riparian corridors to determine the presence, nature, and quantity of surface water in the creek beds. The locations where surface water was encountered were plotted, and a determination was made whether the water was flowing or stagnant. Where flowing water was encountered, the flow rate was determined using a modified version of the *United States Environmental Protection Agency Volunteer Stream Monitoring: A Methods Manual* (USEPA, 1997).

Four fixed stream locations, SF-1 through SF-4, were previously chosen for stream flow measurements. SF-1 is located near Gilman Hot Springs at the southeast border of the site, SF-2 is near MW-67, SF-3 is near MW-15 and MW-18, and SF-4 is near MW-101.

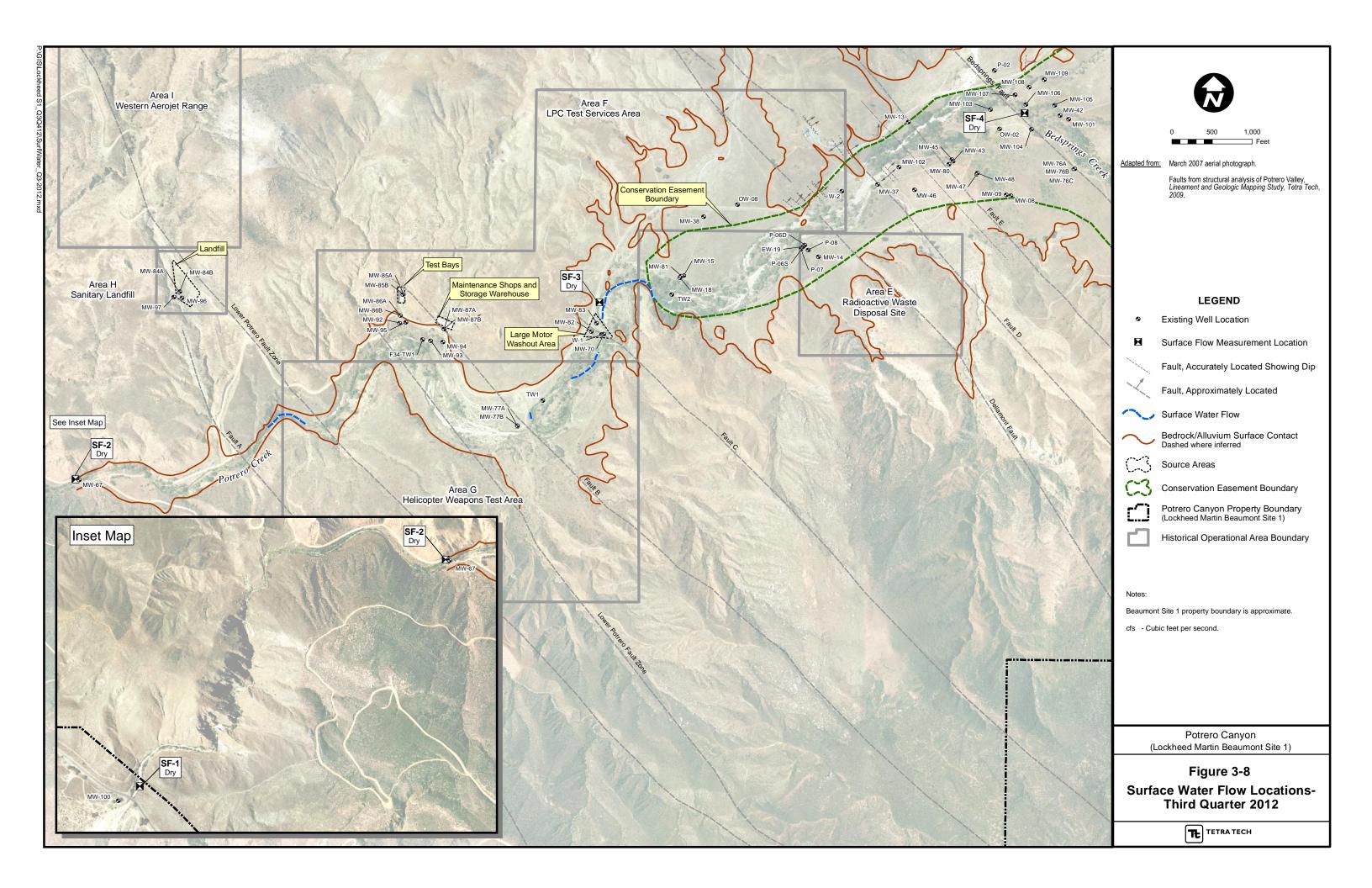
At each location a section of stream that is relatively straight for at least 20 feet was chosen for measurement. This 20-foot section was marked and width measurements were taken at various points to determine the average width. Depth measurements were collected at five points along the width of the stream to determine the average depth of the stream. The average width and depth measurements were multiplied together to obtain an average cross-sectional area. Velocity was measured by releasing a float upgradient and recording the time it took to float through the 20-foot marked section.

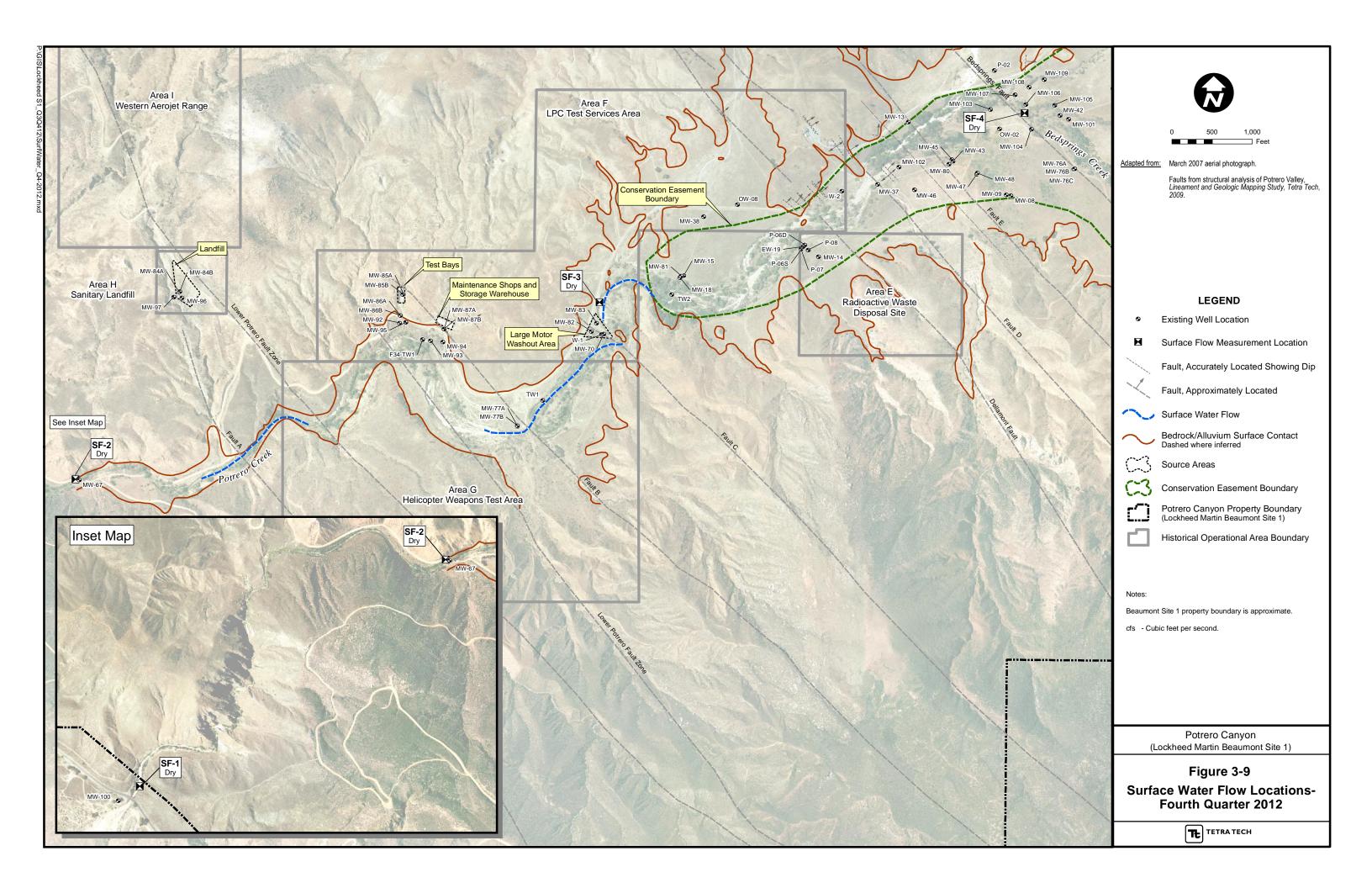
Three velocity measurements were taken and averaged. The length of the measured section was divided by the average velocity, and the answer was multiplied by a factor of 0.9 to correct for friction between the water and stream bed. The average cross-sectional area was then multiplied by the corrected average surface velocity to obtain the average cubic feet per second for water flowing through that section of the stream.

A summary of the surface water flow rates is presented in Table 3-3. The measurement locations and the locations where surface water was encountered are shown on Figures 3-8 and 3-9.

Table 3-3 Surface Water Flow Rates

Location ID	Description of Location	Date Measured	Length of Measured Section (ft)	Width of Measured Section (ft)	Depth of Measured Section (ft)	Float Travel Time (seconds)	Cross Sectional Area (ft²)	Surface Velocity (ft /sec)	Stream Flow Rate (cfs)	Site Stream Flow Rate (cfs)
			Tì	nird Quarter (Au	igust) 2012					
SF-1	Near Gilman Hot Springs Road	08/17/12	Dry	Dry	Dry	Dry	Dry	Dry	Dry	
SF-2	Near MW-67	08/17/12	Dry	Dry	Dry	Dry	Dry	Dry	Dry	Dry
SF-3	Near MW-15 and 18	08/17/12	Dry	Dry	Dry	Dry	Dry	Dry	Dry	Diy
SF-4	Near MW-42	08/17/12	Dry	Dry	Dry	Dry	Dry	Dry	Dry	
			F	ourth Quarter (Nov) 2012					
SF-1	Near Gilman Hot Springs Road	11/30/12	Dry	Dry	Dry	Dry	Dry	Dry	Dry	
SF-2	Near MW-67	11/30/12	Dry	Dry	Dry	Dry	Dry	Dry	Dry	D
SF-3	Near MW-15 and 18	11/30/12	Dry	Dry	Dry	Dry	Dry	Dry	Dry	Dry
SF-4	Near MW-42	11/30/12	Dry	Dry	Dry	Dry	Dry	Dry	Dry	1
Notes:	Measurements are averaged. cfs - cubic feet per second									





3.3 GROUNDWATER FLOW

Groundwater flow directions from Third Quarter 2012 and Fourth Quarter 2012 (Figures 3-1 and 3-2, respectively) were similar to previously observed patterns for a dry period (Appendix A, Figure 2-14). Generally, groundwater flowed northwest from the southeastern limits of the valley (near the former BPA) beneath the former RMPA, toward Potrero Creek where groundwater flow then changed direction and began heading southwest, parallel to the flow of Potrero Creek, into Massacre Canyon.

3.3.1 Horizontal and Vertical Groundwater Gradients

Horizontal groundwater gradients are calculated using a segmented path from well to well that approximates the overall site flowline. The horizontal gradient is a measure of the change in the hydraulic head over a change in distance between wells (the slope of the water table). The overall horizontal groundwater gradient (approximating a flowline from MW-36, upgradient of the BPA, through the RMPA and NPCA to MW-18, in the MCEA) remained constant at 0.013 feet/foot between Third Quarter 2012 and Fourth Quarter 2012. Horizontal gradients are relatively high upgradient of the BPA where recharge from Bedsprings Creek and the adjacent mountain areas enters the main valley. The gradients significantly decrease downgradient of the BPA in the main valley, and then begin to increase again as groundwater flows from the main valley into the canyon just below the confluence of Bedsprings and Potrero creeks.

Vertical groundwater gradients are calculated from individual clusters of wells. Well clusters are used to measure the difference in static water level at different depths in the aquifer. The vertical gradient is a comparison of static water level between wells at different depths in the aquifer, and is an indication of the vertical flow (downward - negative gradient; upward - positive gradient) of groundwater. The vertical groundwater gradients at the site are generally negative in the BPA, RMPA, and NPCA, indicating areas of recharge, and positive in the MCEA, indicating an area of discharge.

Table 3-4 presents a summary of horizontal and vertical groundwater gradients. Appendix F provides a complete listing of historical horizontal and vertical groundwater gradients and associated calculations.

Table 3-4 Summary of Horizontal and Vertical Groundwater Gradients

Horizontal Groundwater Gradients (feet/foot), approximating a flowline from MW-36 to MW-18 and subsections												
Location: Date	Overall MW-36 to MW-18	BPA MW-36 to MW-2	RMPA MW-2 to MW-5	NPCA MW-5 to MW-46	MCEA MW-46 to MW-18							
Previous - Second Quarter (May) 2012	0.013	0.009	0.002	0.021	0.014							
Third Quarter (August) 2012	0.013	0.008	0.002	0.022	0.014							
Fourth Quarter (November) 2012	0.013	0.008	0.002	0.022	0.013							

Vertical Groundwater	Gradients	(feet / foot)
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Location:	BPA	RMPA	NPCA	MCEA	MCEA
shallow screen	MW-59B (MEF)	MW-56B (QAL	MW-75B (QAL)	MW-18 (QAL)	MW-77B (MEF)
Date deep screen	MW-59A (MEF)	MW-56A (MEF)	MW-75A (MEF)	MW-15 (QAL)	MW-77A (MEF)
Previous - Second Quarter (May) 2012	-0.13	-0.14	-0.07	0.02	0.05
Third Quarter (August) 2012	-0.13	-0.14	-0.07	0.02	0.04
Fourth Quarter (November) 2012	-0.13	-0.14	-0.07	0.02	0.03

Notes:

BPA - Burn Pit Area RMPA - Rocket Motor Production Area NPCA - Northern Potrero Creek Area

MCEA - Massacre Canyon Entrance Area

QAL - Quaternary alluvium

MEF - Mount Eden formation

3.4 ANALYTICAL DATA SUMMARY

Summaries of validated laboratory analytical results for organic (volatile organic compounds [VOCs] and 1,4-dioxane) and inorganic (perchlorate, natural attenuation, and general minerals parameters) analytes detected above their respective method detection limits (MDLs) are presented in Table 3-5, Third Quarter 2012 monitoring event, and Tables 3-6 and 3-7, Fourth Quarter 2012 monitoring event. Appendix G provides a complete list of analytes tested, along with validated sample results by analytical method.

Sample results detected above the published California Department of Public Health maximum contaminant level (MCL) or the California Department of Public Health drinking water notification level (DWNL) are bolded in Tables 3-5, 3-6 and 3-7. Appendix H provides laboratory analytical data packages, which include environmental, field quality control (QC), and laboratory QC results, and Appendix I contains consolidated analytical data summary tables. Tables 3-8 and 3-9 present summary statistics of the organic and inorganic analytes detected during the Third Quarter 2012 and Fourth Quarter 2012 monitoring events, respectively.

Table 3-5 Summary of Validated Detected Organic and Inorganic Analytes - Third Quarter 2012

									-			_										
Sample Location	Sample Date	Per- chlorate	1,4- Dioxane	Acetone	Bromo- dichloro- methane	Benzene	Chloro- ethane	Carbon Tetra- chloride	Chloro- form	1,1- Dichloro- ethane	1,2- Dichloro- ethane	1,1- Dichloro- ethene	cis-1,2- Dichloro- ethene	trans-1,2- Dichloro- ethene	Methyl tert-butyl ether	Methylene Chloride	Toluene	1,1,1- Trichloro- ethane	1,1,2- Trichloro- ethane	Trichloro- ethene	Tetrachloro- ethene	Vinyl Chloride
										All re	esults reported in	μg/L unless oth	erwise stated									
MW-110	8/22/2012	7.7	< 0.10	< 5.0	< 0.11	< 0.14	< 0.35	< 0.15	< 0.46	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.43	< 0.15	< 0.22	< 0.12	< 0.31	< 0.25	< 0.23	< 0.13
MW-111A	8/21/2012	2,500	2,800 Je	< 5.0	< 0.11	1.4	< 0.35	< 0.15	5.6	48	130	1,800	18	0.96	< 0.43	1.2 Jq	4.8	0.86	39	610	1.8	0.34 Jq
MW-111B	8/21/2012	25,000	7,300 Je	< 5.0	< 0.11	5.3	< 0.35	1.6	35	230	550	9,700	95	4.7	< 0.43	4.0	4.2	7.6	140	4,800	17	0.31 Jq
MW-111C	8/31/2012	25,000	6,800	< 5.0	< 0.11	5.4	< 0.35	5.6	38	240	600	12,000	100	5.6	< 0.43	3.6	3.1	7.9	160	5,400	22	0.27 Jq
MW-111D	8/21/2012	23,000	6,600 Je	< 5.0	< 0.11	5.6	< 0.35	5.4	39	230	560	10,000	85	5.0	< 0.43	4.0	2.8	7.9	150	5,400	21	0.25 Jq
MW-111E	8/21/2012	36,000	6,100 Je	< 5.0	0.16 Jq	7.1	< 0.35	8.8	53	330	670	16,000	120 Jq	7.3	< 0.43	3.6	2.9	7.6	160 Jq	7,000	25	0.31 Jq
MW-112A	8/20/2012	4,300	63	< 5.0	< 0.11	0.29 Jq	< 0.35	0.96	2.3	11	16	270	1.7	0.28 Jq	0.82 Jq	< 0.15	0.23 Jq	0.23 Jq	1.9	250	1.0	< 0.13
MW-112B	8/20/2012	4,500	70	7.4 Jq	< 0.11	0.49 Jq	0.44 Jq	1.4	2.5	17	21	410	2.6	0.37 Jq	0.63 Jq	0.20 Jq	0.32 Jq	0.38 Jq	2.4	350	1.1	< 0.13
MW-112C	8/20/2012	3,000	59	5.9 Jq	<0.11	0.67	1.0	0.16 Jq	0.81	6.3	12	120	1.4	0.14 Jq	0.48 Jq	0.47 Jq	0.49 Jq	<0.12	0.85	87	< 0.23	<0.13
MDL	(μg/L)	0.071	0.10	5	0.11	0.14	0.35	0.15	0.46	0.098	0.21	0.12	0.18	0.10	0.43	0.15	0.22	0.12	0.31	0.25	0.23	0.13
MCL/DW:	NL (μg/L)	6	1 (1)	-	-	1	-	0.5	-	5	0.5	6	6	10	13	5	150	200	5	5	5	0.5

Notes: Only analytes positively detected are presented in this table. For a complete list, refer to the laboratory data packages in Appendix H.

 μ g/L - Micrograms per liter

MDL - Method detection limit

DWNL - California Department of Public Health drinking water notification level

MCL - California Department of Public Health maximum contaminant level

(1) - DWN

"-" - MCL or DWNL not available.

Bold - MCL or DWNL exceeded.

- <# Analyte not detected; method detection limit concentration is shown.
- J The analyte was positively identified, but the analyte concentration is an estimated value.
- e A holding time violation occurred.
- \boldsymbol{q} $\;$ The analyte detection was below the practical quantitation limit (PQL).

Table 3-6 Summary of Validated Detected Organic Analytes - Fourth Quarter 2012

Sample	Sample	1,4-	Bromo- dichloro-	D	Chloro-	Chloro-	Carbon Tetra-	Chloro-	1,1- Dichloro-	1,2- Dichloro-	1,1- Dichloro-	cis-1,2- Dichloro-	trans-1,2- Dichloro-	1,2- Dichloro-	Methyl tert-butyl	Methylene	Т-1	1,1,1- Trichloro-	1,1,2- Trichloro-	Tri- chloro-	Tetra- chloro-	Vinyl
Location	Date	Dioxane	methane	Benzene	benzene	ethane	chloride	form	ethane	ethane All results	ethene reported in µg/I	ethene	e stated	propane	ether	Chloride	Toluene	ethane	ethane	ethene	ethene	Chloride
F34-TW1	11/26/12	6.6	<0.11	< 0.14	< 0.23	< 0.35	< 0.15	< 0.46	< 0.098	<0.21	0.44 Jq	<0.18	<0.10	<0.19	< 0.43	< 0.15	< 0.22	<0.12	<0.31	1.6	< 0.23	<0.13
IW-04	11/19/12	21	< 0.11	0.19 Jq	< 0.23	< 0.35	< 0.15	< 0.46	0.20 Jq	< 0.21	9.6	0.93	0.40 Jq	<0.19	< 0.43	0.16 Jq	< 0.22	<0.12	< 0.31	14	< 0.23	0.34 Jq
MW-05	11/28/12	34	< 0.11	< 0.14	< 0.23	< 0.35	< 0.15	3.3	3.4	0.53	130	0.49 Jq	< 0.10	<0.19	< 0.43	< 0.15	< 0.22	0.34 Jq	0.33 Jq	150	< 0.23	<0.13
MW-08	11/26/12	< 0.10	< 0.11	< 0.14	< 0.23	< 0.35	< 0.15	< 0.46	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	<0.19	< 0.43	< 0.15	< 0.22	<0.12	< 0.31	< 0.25	< 0.23	< 0.13
MW-13	11/26/12	< 0.10	< 0.11	< 0.14	< 0.23	< 0.35	< 0.15	< 0.46	< 0.098	< 0.21	<0.12	<0.18	< 0.10	<0.19	< 0.43	< 0.15	< 0.22	< 0.12	< 0.31	< 0.25	< 0.23	< 0.13
MW-15	11/16/12	6.2	<0.11	< 0.14	<0.23	< 0.35	< 0.15	< 0.46	0.32 Jq	< 0.21	1.7	0.29 Jq	< 0.10	<0.19	<0.43	< 0.15	< 0.22	< 0.12	< 0.31	0.93	< 0.23	< 0.13
MW-18	11/16/12	4.5	<0.11	< 0.14	<0.23	< 0.35	< 0.15	< 0.46	0.18 Jq	<0.21	1.3		< 0.10	<0.19	<0.43	< 0.15	< 0.22	<0.12	< 0.31	1.0	< 0.23	< 0.13
MW-43	11/26/12	6.9	<0.11	< 0.14	<0.23	< 0.35	< 0.15	< 0.46	0.42 Jq	<0.21	5.0	1	+	<0.19	<0.43	< 0.15	< 0.22	<0.12	<0.31	4.7	< 0.23	<0.13
MW-48	11/28/12	0.71	<0.11	<0.14	<0.23	<0.35	<0.15	<0.46	<0.098	<0.21	<0.12	<0.18	<0.10	<0.19	<0.43	<0.15	<0.22	<0.12	<0.31	<0.25	<0.23	<0.13
MW-60A	11/29/12	110	<0.11	0.26 Jq	<0.23	<0.35	0.47 Jq	2.5	5.1	7.0	420	2.3	· · · · · · · · · · · · · · · · · · ·	<0.19	<0.43	<0.15	<0.22	0.25 Jq	1.5		0.46 Jq	<0.13
MW-60B	11/29/12	15	<0.11	<0.14	<0.23	<0.35	<0.15	<0.46	0.66	0.89	52	0.22 Jq	<0.10	<0.19	<0.43	<0.15	<0.22	<0.12	<0.31	22	<0.23	<0.13
MW-67	11/16/12	0.92	<0.11	<0.14	<0.23	<0.35	<0.15	<0.46	<0.098	<0.21	<0.12	<0.18	<0.10	<0.19	<0.43	<0.15	<0.22	<0.12	<0.31	<0.25	<0.23	<0.13
MW-68 MW-70	11/28/12 11/16/12	21 2.9	<0.11	<0.14	<0.23	<0.35 <0.35	<0.15 <0.15	<0.46	0.17 Jq <0.098	<0.21	<0.12	<0.18	<0.10	<0.19	<0.43	<0.15	<0.22	<0.12	<0.31	<0.25	<0.23	<0.13
MW-76B	11/10/12	0.27	<0.11	<0.14	<0.23	<0.35	<0.15	<0.46	<0.098	<0.21	<0.12	<0.18	<0.10	<0.19	<0.43	<0.15	<0.22	<0.12	<0.31	<0.25	<0.23	<0.13
MW-88	11/19/12	0.19 Jq	<0.11	<0.14	<0.23	<0.35	<0.15	<0.46	<0.098	<0.21	<0.12	<0.18	<0.10	<0.19	<0.43	<0.15	<0.22	<0.12	<0.31	<0.25	<0.23	<0.13
MW-93	11/19/12	22	<0.11	<0.14	<0.23	<0.35	<0.15	<0.46	0.17 Jq	<0.21		<0.18	<0.10	<0.19	<0.43	<0.15	<0.22	<0.12	<0.31	3.2	<0.23	<0.13
MW-98B	11/28/12	11	<0.11	<0.14	<0.23	< 0.35	<0.15	1.4	0.33 Jq	<0.21		<0.18	<0.10	<0.19	<0.43	<0.15	<0.22	<0.12	<0.31	26	<0.23	<0.13
MW-100	11/16/12	0.23	< 0.11	< 0.14	< 0.23	< 0.35	< 0.15	< 0.46	<0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.19	< 0.43	< 0.15	< 0.22	< 0.12	< 0.31	< 0.25	< 0.23	< 0.13
MW-101	11/28/12	21	< 0.11	< 0.14	< 0.23	< 0.35	< 0.15	< 0.46	1.3	0.41 Jq	45	43	1.5	<0.19	< 0.43	< 0.15	< 0.22	< 0.12	< 0.31	32	< 0.23	1.8
MW-102	11/28/12	18	<0.11	< 0.14	< 0.23	< 0.35	< 0.15	< 0.46	1.3	< 0.21	17	26	1.8	<0.19	< 0.43	< 0.15	< 0.22	< 0.12	< 0.31	15	< 0.23	2.4
MW-103	11/27/12	14	< 0.11	< 0.14	< 0.23	< 0.35	< 0.15	< 0.46	0.77	< 0.21	3.7	4.9	< 0.10	< 0.19	< 0.43	< 0.15	< 0.22	< 0.12	< 0.31	11	< 0.23	< 0.13
MW-104	11/27/12	27	< 0.11	< 0.14	< 0.23	< 0.35	< 0.15	< 0.46	5.9	0.26 Jq	55	3.7	0.25 Jq	<0.19	< 0.43	< 0.15	< 0.22	<0.12	<0.31	4.5	< 0.23	15
MW-105	11/27/12	30	< 0.11	< 0.14	< 0.23	< 0.35	< 0.15	< 0.46	4.8	0.55	74	7.0	1.7	<0.19	< 0.43	< 0.15	< 0.22	< 0.12	< 0.31	69	< 0.23	2.0
MW-106	11/27/12	28	<0.11	< 0.14	< 0.23	< 0.35	< 0.15	< 0.46	2.7	0.37 Jq	28	2.8	1.3	< 0.19	< 0.43	< 0.15	< 0.22	< 0.12	< 0.31	27	< 0.23	0.31 Jq
MW-107	11/27/12	16	<0.11	< 0.14	<0.23	< 0.35	< 0.15	< 0.46	1.1	<0.21	10	3.3		<0.19	< 0.43	< 0.15	< 0.22	<0.12	<0.31	11	< 0.23	< 0.13
MW-109	11/27/12	26	<0.11	<0.14	<0.23	<0.35	<0.15	<0.46	3.0	0.52	55		· · · · · · · · · · · · · · · · · · ·	<0.19	<0.43	<0.15	<0.22	<0.12	<0.31	64	<0.23	0.78
MW-110	11/19/12	<0.10	<0.11	<0.14	<0.23	<0.35	<0.15	<0.46	<0.098	<0.21	<0.12	<0.18	<0.10	<0.19	<0.43	<0.15	<0.22	<0.12	<0.31	<0.25	<0.23	<0.13
MW-111A	11/20/12	1,900	<0.11	1.3	<0.23	<0.35	<0.15	5.5	<0.098	130	2,000	19	+	<0.19	<0.43	1.0 Jq	3.3	0.80	34	+	2.1	0.37 Jq
MW-111B	11/20/12	5,800	<0.11	5.6	0.24 Jq	<0.35	0.71	39	240	590	10,000	110	+	0.28 Jq	<0.43	3.5	3.8	6.4	190	4,400	18	0.40 Jq
MW-111D MW-111E	11/20/12 11/20/12	7,900 4,800	<0.11 0.14 Jq	6.1	<0.23	<0.35 <0.35	5.8	44 60	310 430	680 840	14,000 20,000	120 170	6.0 8.5	0.27 Jq 0.35 Jq	<0.43	3.4 2.9 Jq	1.9	6.8	150 170	6,500 9,200	25 32	0.39 Jq 0.51
MW-111E MW-112A	11/20/12	64	<0.11	7.8 0.23 Jq	<0.23	<0.35	0.77	2.1	9.7	14	260	1.5		<0.19	0.75 Jq	<0.15	<0.22	0.18 Jq	1.7	270	0.95	<0.13
MW-112A MW-112B	11/20/12	72	<0.11	0.23 Jq 0.44 Jq	<0.23	0.53	1.2	2.5	17	20	430	2.8	· · · · · · · · · · · · · · · · · · ·	<0.19	0.73 Jq 0.58 Jq	0.23 Jq	0.24 Jq	0.30 Jq	2.2		1.2	<0.13
MW-112D	11/20/12	66	<0.11	0.53	<0.23	0.98	0.28 Jq	1.2	8.9	14	170	2.1		<0.19	0.58 Jq 0.53 Jq	0.46 Jq	0.24 Jq	<0.12	1.1		0.27 Jq	<0.13
OW-02	11/28/12	10	<0.11	<0.14	<0.23	<0.35	<0.15	<0.46	0.45 Jg	<0.21		<0.18	<0.10	<0.19	<0.43	<0.15	<0.22	0.18 Ja	<0.31	17	<0.23	<0.13
P-02	11/26/12		<0.11	<0.14	<0.23	<0.35	<0.15	<0.46	<0.098	<0.21	<0.12	<0.18	<0.10	<0.19	<0.43	<0.15	<0.22	<0.12	<0.31	<0.25	<0.23	<0.13
SW-02	11/15/12		<0.11	<0.14	<0.23	< 0.35	<0.15	< 0.46	0.40 Jq	<0.21	9.8			<0.19	<0.43	<0.15	0.39 Jq	0.16 Jq	<0.31	13	< 0.23	0.38 Jq
SW-03	11/15/12		<0.11	<0.14	<0.23	<0.35	<0.15	<0.46	<0.098	<0.21	1.1	0.42 Jq		<0.19	<0.43	<0.15		<0.12	<0.31	1.1	<0.23	<0.13
SW-04	11/15/12		< 0.11	< 0.14	< 0.23	< 0.35	< 0.15	< 0.46	< 0.098	< 0.21	1.0			<0.19	<0.43	< 0.15		<0.12	<0.31	1.2	< 0.23	<0.13
SW-09	11/15/12			< 0.14	< 0.23	< 0.35	< 0.15	< 0.46	< 0.098	< 0.21	< 0.12	<0.18	< 0.10	<0.19	< 0.43	< 0.15	< 0.22	< 0.12	< 0.31	< 0.25	< 0.23	<0.13
SW-18	11/15/12	3.8	< 0.11	< 0.14	< 0.23	< 0.35	< 0.15	< 0.46	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	<0.19	< 0.43	< 0.15	< 0.22	< 0.12	< 0.31	< 0.25	< 0.23	< 0.13
MDL (µ	ug/L)	0.10	0.11	0.14	0.23	0.35	0.15	0.46	0.098	0.21	0.12	0.18	0.10	0.19	0.43	0.15	0.22	0.12	0.31	0.25	0.23	0.13
MCL/DWN		1 (1)	-	1	-	-	0.5	-	5	0.5	6	6	10	5	13	5	150	200	5	5	5	0.5
		es positively d	latactad are prese	nted in this t	table For a cor	mplete list ref	er to the laborato	rv data nacka	ges in Annendiy	П												

Notes: Only analytes positively detected are presented in this table. For a complete list, refer to the laboratory data packages in Appendix H.

μg/L - Micrograms per liter

mg/L - Milligrams per liter

MDL - Method detection limit

DWNL - California Department of Public Health drinking water notification level

MCL - California Department of Public Health maximum contaminant level

(1) - DWNL

"-" - MCL or DWNL not available.
Bold - MCL or DWNL exceeded.

<#- Analyte not detected; method detection limit concentration is shown.

J - The analyte was positively identified, but the analyte concentration is an estimated value.

q - The analyte detection was below the practical quantitation limit (PQL).

Table 3-7 Summary of Validated Detected Inorganic Analytes - Fourth Quarter 2012

		Per-	Dissolved Organic	Alkalinity,			Nitrate			Dissolved
Sample	Sample	chlorate	Carbon -	Total -	Bicarbonate	Chloride	as N -	Sulfate	Iron -	Manganese
Location F34-TW1	Date 11/26/12	- μg/L <0.071	mg/L NA	mg/L NA	-mg/L NA	-mg/L NA	mg/L NA	-mg/L NA	ug/L NA	-ug/L NA
IW-04	11/26/12	0.27	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA
MW-05	11/19/12	2,900	0.36 Jq	120	150	9.5	7.9	11	<2.3	1.4 Jq
MW-08	11/26/12	<0.071	0.30 Jq 0.27 Jq	96	120	5.1	<0.11	11	1.400	1.4 34
MW-13	11/26/12	0.36	0.27 34	310	370	9.6	1.4	79	4.8 Jq	<0.24
MW-15	11/26/12	<0.071	NA	NA	NA	NA	NA	NA	NA	NA
MW-18	11/16/12	3.0	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA
MW-43	11/16/12	54	0.29 Jq	83	100	5.8	0.77	8.7	96	4.5 Jq
MW-48	11/28/12	< 0.071	1.1	170	210	7.9	<0.11	6.3	1,900	250
MW-46 MW-60A	11/28/12	6,600	NA	NA	NA	NA	NA	NA	1,900 NA	NA
MW-60B	11/29/12	1,800	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA
MW-67	11/16/12	<0.071	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA
MW-68	11/10/12	17,000	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA
MW-70	11/26/12	0.22	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA
MW-76B	11/10/12	<0.071	0.31 Jq	97	120	5.6	<0.11	8.4	220	11
MW-88	11/28/12	610	NA	NA	NA	NA	NA	NA	NA	NA
MW-93	11/19/12	5.5	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA
MW-98B	11/19/12	1.700	NA NA	NA NA	NA NA	NA NA	NA	NA	NA NA	NA NA
MW-100	11/26/12	<0.071	NA NA	NA NA	NA NA	NA NA	NA NA	NA	NA NA	NA NA
MW-100	11/10/12	<0.071	0.39 Jq	77	94	7.1	<0.11	28	1,700	110
MW-101	11/28/12	<0.071	0.39 Jq 0.49 Jq	120	140	6.3	<0.11	16	80	56
MW-102	11/28/12	31	1.1	140	180	7.5	<0.11	6.9 Jd	2,100 Jcf	11
MW-103	11/27/12	<0.071	0.70	97	120	7.3	<0.11	4.0	400	98
MW-105	11/27/12	<0.071	0.70 0.64 Jq	130	160	11	<0.11	27	120	860
MW-106	11/27/12	<0.071	2.7	220	260	17	<0.11	63	5.1 Jq	120
MW-107	11/27/12	53	1.7	250	300	14	<0.11	26	12 Jq	62
MW-109	11/27/12	310	0.48 Jq	120	140	9.3	1.3	23	5.8 Jq	320
MW-110	11/19/12	14	NA	NA	NA NA	NA	NA	NA	NA	NA NA
MW-111A	11/20/12	2,600	NA	NA	NA	NA	NA	NA	NA	NA
MW-111B	11/20/12	36,000	NA	NA	NA	NA	NA	NA	NA	NA
MW-111D	11/20/12	38,000	NA	NA	NA	NA	NA	NA	NA	NA
MW-111E	11/20/12	49,000	NA	NA	NA	NA	NA	NA	NA	NA
MW-112A	11/20/12	5,800	NA	NA	NA	NA	NA	NA	NA	NA
MW-112B	11/20/12	6,800	NA	NA	NA	NA	NA	NA	NA	NA
MW-112C	11/20/12	4,700	NA	NA	NA	NA	NA	NA	NA	NA
OW-02	11/28/12	350	0.33 Jq	71	87	7.1	3.2	7.5	6.6 Jq	0.26 Jq
P-02	11/26/12	0.32	1.3	230	280	76	0.72	490	1,200	790
SW-02	11/15/12	100	NA	NA	NA	NA	NA	NA	NA NA	NA
SW-03	11/15/12	3.0	NA	NA	NA	NA	NA	NA	NA	NA
SW-04	11/15/12	3.6	NA	NA	NA	NA	NA	NA	NA	NA
SW-09	11/15/12	< 0.071	NA	NA	NA	NA	NA	NA	NA	NA
SW-18	11/15/12	< 0.071	NA	NA	NA	NA	NA	NA	NA	NA
	MDL	0.071	0.36	1.7	1.7	1.7	0.11	0.37	2.3	0.24
M	CL/DWNL	6	-	-	-	250	10	250	300	500 (1)
Notes		4				-1-4-1:-4£				3 7

Notes: Only analytes positively detected are presented in this table. For a complete list, refer to the laboratory data packages in Appendix H.

MDL - Method detection limit

MCL - California Department of Public Health Services maximum contaminant level

DWNL - California Department of Public Health Services drinking water notification level

(1) DWNL

 $\mu g/L$ - Micrograms per liter

mg/L - Milligrams per liter

Bold - MCL or DWNL exceeded.

NA - not analyzed

- <# Analyte not detected; method detection limit concentration is shown.
- J The analyte was positively identified, but the analyte concentration is an estimated value.
- c The Matrix Spike (MS) and/or Matrix Spike Duplicate (MSD) recoveries were outside control limits.
- f The duplicate Relative Percent Difference was outside the control limit.
- q The analyte detection was below the practical quantitation limit (PQL).

[&]quot;-" - MCL or DWNL not available.

Table 3-8 Summary Statistics of Validated Organic and Inorganic Analytes - Third Quarter 2012

Organic Analytes Detected	Total Number of Samples Analyzed	Total Number of Detections (1)	Number of Detections Exceeding MCL or DWNL (1)	MCL	/DWNL	Conce	imum ntration ected	Maxin Concent Detec	ration
1,4-Dioxane	9	8	8	1 (2)	μg/L	59	μg/L	7,300	μg/L
Acetone	9	2	0	-	μg/L	5.9	μg/L	7.4	μg/L
Bromodichloromethane	9	1	0	-	μg/L	0.16	μg/L	0.16	μg/L
Benzene	9	8	5	1	μg/L	0.29	μg/L	7.1	μg/L
Chloroethane	9	2	0	-	μg/L	0.44	μg/L	1.0	μg/L
Carbon tetrachloride	9	7	6	0.5	μg/L	0.16	μg/L	8.8	μg/L
Chloroform	9	8	0	-	μg/L	0.81	μg/L	53	μg/L
1,1-Dichloroethane	9	8	8	5	μg/L	6.3	μg/L	330	μg/L
1,2-Dichloroethane	9	8	8	0.5	μg/L	12	μg/L	670	μg/L
1,1-Dichloroethene	9	8	8	6	μg/L	120	μg/L	16,000	μg/L
cis-1,2-Dichloroethene	9	8	5	6	μg/L	1.4	μg/L	120	μg/L
trans-1,2-Dichloroethene	9	8	0	10	μg/L	0.14	μg/L	7.3	μg/L
Methyl-tert-butyl ether	9	3	0	13	μg/L	0.48	μg/L	0.82	μg/L
Methylene Chloride	9	7	0	5	μg/L	0.20	μg/L	1.20	μg/L
Toluene	9	8	0	150	μg/L	0.23	μg/L	4.8	μg/L
1,1,1-Trichloroethane	9	7	0	200	μg/L	0.23	μg/L	7.9	μg/L
1,1,2-Trichloroethane	9	8	5	5	μg/L	0.85	μg/L	160	μg/L
Trichloroethene	9	8	8	5	μg/L	87	μg/L	7,000	μg/L
Tetrachloroethene	9	7	4	5	μg/L	1.0	μg/L	25	μg/L
Vinyl chloride	9	5	0	0.5	μg/L	0.25	μg/L	0.34	μg/L
Inorganic Analytes Detected	Total Number of Samples Analyzed	Total Number of Detections (1)	Number of Detections Exceeding MCL or DWNL (1)	MCL/DWNL		Conce	imum ntration ected	Maxin Concent Detec	ration
Perchlorate	9	9	9	6	μg/L	7.7	μg/L	36,000	μg/L

Notes: DWNL - California Department of Public Health drinking water notification level

MCL - California Department of Public Health maximum contaminant level

" - " - MCL or DWNL not established.

(1) - Number of detections excludes sample duplicates, trip blanks and equipment blanks

(2) - DWNL

μg/L - Micrograms per liter

Table 3-9 Summary Statistics of Validated Organic and Inorganic Analytes - Fourth Quarter 2012

Organic Analytes Detected	Total Number of Samples Analyzed	Total Number of Detections (1)	Number of Detections Exceeding MCL or DWNL (1)	MCI	./DWNL	Concer	mum ntration ected	Maxii Concent Detec	tration
1,4-Dioxane	42	38	33	1 (2)	μg/L	0.19	μg/L	7,900	μg/L
Bromodichloromethane	42	1	0	-	μg/L	0.14	μg/L	0.14	μg/L
Benzene	42	9	4	1	μg/L	0.19	μg/L	7.8	μg/L
Chlorobenzene	42	1	0	-	μg/L	0.24	μg/L	0.24	μg/L
Chloroethane	42	2	0	-	μg/L	0.53	μg/L	0.98	μg/L
Carbon tetrachloride	42	7	5	0.5	μg/L	0.28	μg/L	11	μg/L
Chloroform	42	10	0	-	μg/L	1.2	μg/L	60	μg/L
1,1-Dichloroethane	42	26	8	5	μg/L	0.17	μg/L	430	μg/L
1,2-Dichloroethane	42	15	12	0.5	μg/L	0.26	μg/L	840	μg/L
1,1-Dichloroethene	42	30	22	6	μg/L	0.26	μg/L	20,000	μg/L
cis-1,2-Dichloroethene	42	24	7	6	μg/L	0.19	μg/L	170	μg/L
trans-1,2-Dichloroethene	42	16	0	10	μg/L	0.18	μg/L	8.5	μg/L
1,2-Dichloropropane	42	3	0	5	μg/L	0.27	μg/L	0.35	μg/L
Methyl-tert-butyl ether	42	3	0	13	μg/L	0.53	μg/L	0.75	μg/L
Methylene Chloride	42	7	0	5	μg/L	0.16	μg/L	3.5	μg/L
Toluene	42	9	0	150	μg/L	0.24	μg/L	6.2	μg/L
1,1,1-Trichloroethane	42	10	0	200	μg/L	0.16	μg/L	6.8	μg/L
1,1,2-Trichloroethane	42	9	4	5	μg/L	1.10	μg/L	190	μg/L
Trichloroethene	42	29	21	5	μg/L	0.93	μg/L	9,200	μg/L
Tetrachloroethene	42	8	3	5	μg/L	0.27	μg/L	32	μg/L
Vinyl chloride	42	12	6	0.5	μg/L	0.31	μg/L	15	μg/L
Dissolved Organic Carbon	16	16	0	-	mg/L	0.27	mg/L	2.7	mg/L
Methane	16	13	0	-	μg/L	0.20	μg/L	1,700	μg/L
Acetic Acid	16	6	0		mg/L	0.075	mg/L	0.12	mg/L
2-Hydroxypropanoic Acid	16	1	0	-	mg/L	0.22	mg/L	0.22	mg/L
Hexanoic Acid	16	1	0	-	mg/L	0.44	mg/L	0.44	mg/L
Inorganic Analytes Detected	Total Number of Samples Analyzed	Total Number of Detections (1)	Number of Detections Exceeding MCL or DWNL (1)	MCL	/ DWNL	Concer	mum ntration ected	Maxii Concen Detec	tration
Perchlorate	42	28	20	6	μg/L	0.22	μg/L	49,000	μg/L
Hydrogen	16	9	0	-	nM	0.60	nM	2.2	nM
Alkalinity, Total	16	16	0	-	mg/L	71	mg/L	310	mg/L
Bicarbonate	16	16	0	-	mg/L	87	mg/L	370	mg/L
Chloride	16	16	0	250	mg/L	5.1	mg/L	76	mg/L
Nitrate	16	6	0	10	mg/L	0.72	mg/L	7.9	mg/L
Sulfate	16	16	1	250	mg/L	4.0	mg/L	490	mg/L
Iron	16	15	6	300	μg/L	4.8	μg/L	2,100	μg/L
Dissolved Manganese	16	15	1	500	μg/L	0.26	μg/L	860	μg/L

Notes: DWNL - California Department of Public Health drinking water notification level

MCL - California Department of Public Health maximum contaminant level

" - " - MCL or DWNL not established.

(1) - Number of detections excludes sample duplicates, trip blanks and equipment blanks.

(2) - DWNL

 $\begin{array}{ll} mg/L - & Milligrams \ per \ liter \\ \mu g/L - & Micrograms \ per \ liter \end{array}$

nM - NanoMoles

3.4.1 Data Quality Review

The quality control samples were reviewed as described in the *Programmatic Sampling and Analysis Plan, Lockheed Martin Corporation, Beaumont Sites 1 and 2, Beaumont, California* (Tetra Tech, 2010). The data for the groundwater sampling activities were contained in analytical data packages generated by Microseeps Laboratories, Inc. and E.S. Babcock and Sons Laboratories, Inc. These data packages were reviewed using the latest versions of the United States Environmental Protection Agency Contract Laboratory Program National Functional Guidelines for Organic and Inorganic Superfund Data Review (USEPA, 2008 and 2010).

Preservation criteria, holding times, field blanks, laboratory control samples, method blanks, duplicate environmental samples, spiked samples, and surrogate and spike recovery data were reviewed. Within each environmental sample, the sample-specific quality control spike recoveries were examined. These data examinations included comparing statistically calculated control limits to percent recoveries of all spiked analytes and duplicate spiked analytes. Relative percent difference (RPD) control limits were compared to actual spiked (matrix spike/matrix spike duplicate) RPD results. Surrogate recoveries were examined for all organic compound analyses and compared to their control limits.

Environmental samples were analyzed by the following methods: Method ASTM 2320 for alkalinity, Method AM23GS for volatile fatty acids, Method AM20GAX for hydrogen, Method E300.0 for anions, EPA Method E332.0 for perchlorate, Method A5310B for total organic carbon, Method RSK-175 for methane, ethane, ethene, EPA Method SW8270C SIM for 1,4-dioxane, Method SW6010B for metals, and Methods E524.2 and SW8260B for VOCs.

Unless otherwise noted below, all data results met required criteria, are of known precision and accuracy, did not require qualification, and may be used as reported.

EPA Method SW8270C SIM for 1, 4-dioxane had holding time errors that caused 6.9 percent (4 samples out of 58 samples) of the total SW8270C SIM data to be qualified as estimated. The samples were extracted one day after the seven-day holding time requirement. The holding time errors are minor and the negative effect on the data is minor. The data qualified as estimated are usable for the intended purpose.

After conversations with the laboratory concerning holding time errors, the cause of the error has been determined to be the manner in which the laboratory determines the holding time.

- The laboratory consulted the EPA and determined that holding times are counted by days and not hours for methods that the holding time is stated in days. Therefore, by rounding days, it is possible to not have a holding time error for eight days out from sample receipt for EPA Method SW8270C SIM.
- Tetra Tech uses a 24-hour clock to determine holding times as the National Functional Guidelines indicate. Using a 24-clock for day determination does not allow for day rounding. Therefore, eight days out from sample receipt is a holding time error for EPA Method SW8270C SIM.

We will continue to interact with the laboratory in order to resolve this holding time situation and request that holding times be calculated using a 24 hour clock.

Method 300.0 for anions had laboratory control samples (LCS) recovery below the lower limit which resulted in 6.3 percent (2 out of 54) of the total anion data to be qualified as estimated. The LCS lower control limit is 90 percent. The recovery that leads to the qualified results was 88.8 percent. Because the LCS error is minor, the effect on the data is minor and the data may be used for the intended purpose.

Method SW6010B for metals had matrix spike recovery errors and RPD errors that qualified as estimated 5.5 percent (2 out of 36) of the total SW6010B data. The data may be used for the intended purpose.

3.5 CHEMICALS OF POTENTIAL CONCERN

The identification of chemicals of potential concern is an ongoing process that takes place annually as part of the second quarter sampling. The purposes for identifying chemicals of potential concern are to establish a list of analytes that best represents the extent and magnitude of affected groundwater, and to focus more detailed analysis on only those analytes. The analytes were organized and evaluated in two groups, organic and inorganic, and divided into primary and secondary chemicals of potential concern. Tables 3-5, 3-6, and 3-7 present summaries of the organic and inorganic analytes detected during the Third Quarter 2012 and Fourth Quarter 2012 monitoring events. Data that are "B" qualified because of their associations with either laboratory blank contamination or field cross-contamination are not included in the chemicals of potential concern evaluation.

The identification process for chemicals of potential concern does not eliminate analytes from testing, but does reduce the number of analytes that are evaluated and discussed during reporting. All of the secondary chemicals of potential concern will continue to be tested during future monitoring events, because of their association with other analytes that are listed as primary chemicals of potential concern. However, the secondary chemicals of potential concern are not discussed further because they are detected on a more limited or inconsistent basis, and/or are detected at concentrations below a regulatory threshold. The standard list of analytes for each method will continue to be tested for and screened annually to insure that the appropriate chemicals of potential concern are being identified and evaluated.

3.5.1 Identification of Chemicals of Potential Concern

Chemicals of potential concern have been selected to include compounds that are consistently detected in groundwater at concentrations above regulatory limits and that can be used to assess the extent of affected groundwater. Primary chemicals of potential concern are parent products such as trichloroethene (TCE) and 1,1,1-trichloroethane (1,1,1-TCA), and are always present with secondary chemicals of potential concern. Secondary chemicals of potential concern are breakdown products such as 1,1-dichloroethane (1,1-DCA) and 1,1-dichloroethene (1,1-DCE), and are detected at lower concentrations than their parent products. At this site 1,1-DCE, a breakdown product of 1,1,1-TCA, is detected at higher concentrations than 1,1,1-TCA, so 1,1-DCE is considered the primary chemical of potential concern, and 1,1,1-TCA is considered a secondary chemical of potential concern.

As discussed above, the chemicals of potential concern analysis is intended to streamline and focus the evaluation of the contaminant data collected during monitoring events. It is not intended to trivialize or dismiss the analytes screened out as part of the process. Therefore, to ensure that all analytes detected receive the proper attention, this chemicals of potential concern analysis is performed annually.

An annual evaluation of chemicals of potential concern based on the results of the Second Quarter 2012 water quality monitoring event was presented in the First and Second Quarter 2012 Semiannual Groundwater Monitoring Report (Tetra Tech, 2011). Based on the results of water quality monitoring and the screening of those results against the existing chemicals of potential concern, the MCLs, and DWNLs, no additional chemicals of potential concern were identified,

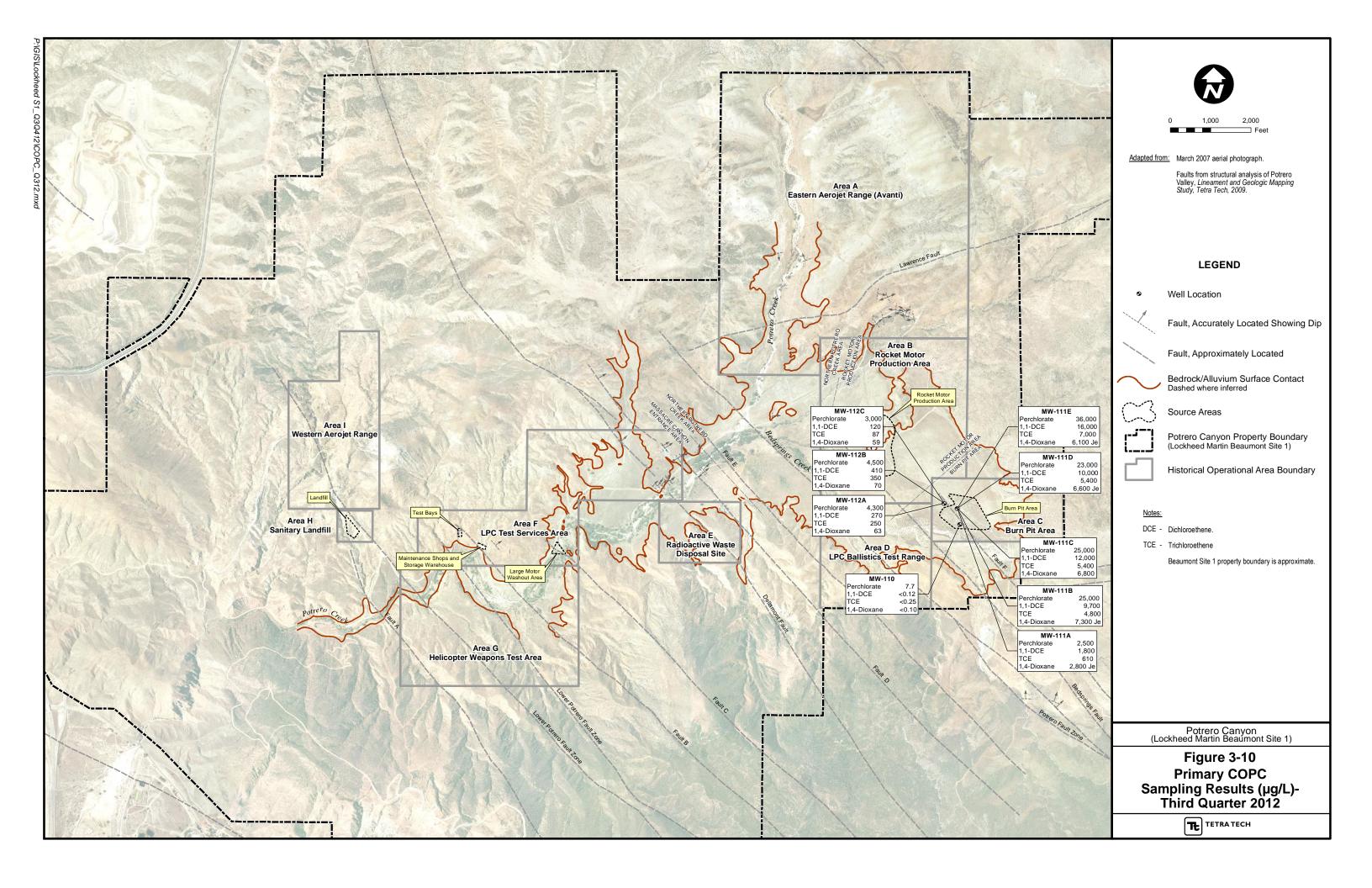
nor was there evidence for removing an analyte from the existing chemicals of potential concern list. Table 3-10 presents those groundwater analytes that have been identified as chemicals of potential concern. Time-series graphs of primary and secondary chemicals of potential concern are provided in Appendix E.

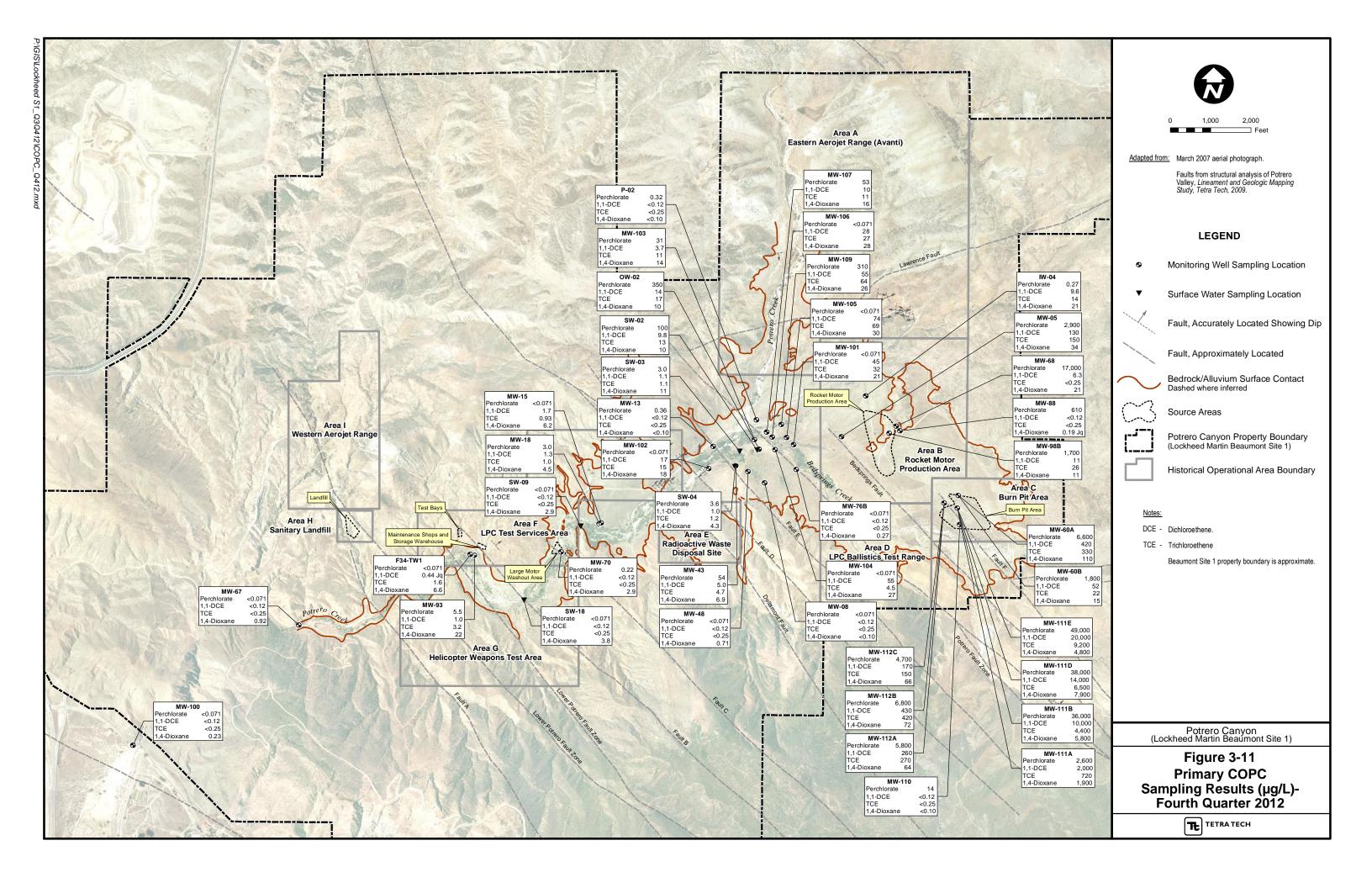
Table 3-10 Groundwater Chemicals of Potential Concern

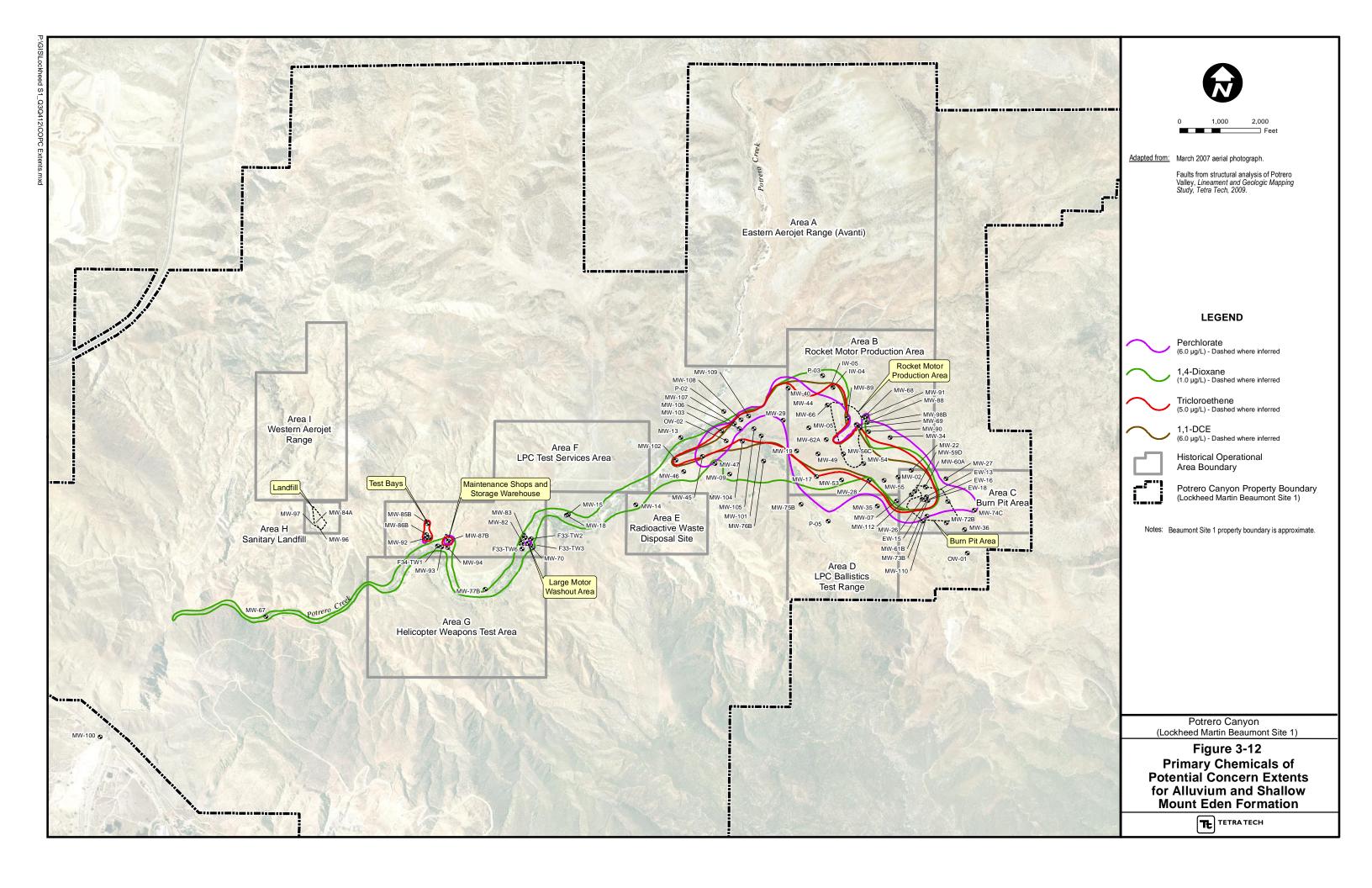
Analyte	Classification	Comments
Perchlorate	Primary	Parent product (propellant), widely detected at site
1,1-Dichloroethene	Primary	Breakdown product of 1,1,1-TCA, detected at higher concentrations than 1,1,1-TCA at site
Trichloroethene	Primary	Parent product (solvent), widely detected at site
1,4-Dioxane	Primary	Stabilizer in 1,1,1-TCA, widely detected at site
1,1-Dichloroethane	Secondary	Breakdown product of 1,1,1-TCA
1,2-Dichloroethane	Secondary	Breakdown product of 1,1,1-TCA
1,1,1-Trichloroethane	Secondary	Parent product (solvent), detected at lower concentrations than breakdown product (1,1-DCE) at site
1,1,2-Trichloroethane	Secondary	Isomeric impurity of 1,1,1-TCA
cis-1,2-Dichloroethene	Secondary	Breakdown product of TCE
Vinyl chloride	Secondary	Breakdown product of TCE and/or 1,1,1-TCA

3.6 DISTRIBUTION OF THE PRIMARY CHEMICALS OF POTENTIAL CONCERN

The Third Quarter 2012 and Fourth Quarter 2012 monitoring events are minor events. Only guard wells, wells with increasing contaminant trends, new wells, and surface water locations are sampled and tested during these events (Tetra Tech, 2003b). Only those wells and surface water locations sampled and tested during this event will be discussed. Figures 3-10 and 3-11 present the primary chemicals of potential concern sampling results for the wells sampled during the Third Quarter 2012 and Fourth Quarter 2012 monitoring events, respectively. A figure illustrating the extent of the primary chemicals of potential concern based on the recent data is presented in Figure 3-12.







3.6.1 Guard Wells

Guard wells are wells that are used as an early warning to protect private and municipal wells by detecting any upgradient contaminants. Guard wells are also used to monitor any migration of contaminants off-site.

Four monitoring wells, MW-15, MW-18, MW-67, and MW-100, were designated as guard wells during the semiannual event conducted during the second quarter of the year. Wells MW-15 and MW-18 are a clustered well pair. Well MW-18 is completed near the top of the alluvial aquifer and MW-15 is completed near the bottom of the alluvial aquifer. Well MW-67 is the farthest downgradient site well, and is located approximately 0.9 miles upgradient of the southern edge of the site. MW-100 is located off-site approximately 500 feet from the southern property boundary just south of Gilman Springs Road near the mouth of Potrero Creek. Table 3-11 presents a summary of the detected chemicals of potential concern reported in guard well samples collected during Fourth Quarter 2012, and previous monitoring events. In general, the chemicals of potential concern concentrations during this reporting period have remained stable in the guard wells.

Table 3-11 Summary of Detected Chemicals of Potential Concern in Guard Wells

Sample Location	Site Area	Sample Date	Per- chlorate	1,4- Dioxane	1,1-Dichloro- ethane	1,1-Dichloro- ethene	cis-1,2-Dichloro- ethene	Trichloro- ethene
	1				ug/L unless otherwi			T
		01/06/11	< 0.071	5.9	0.34 Jq	2.0	0.30 Jq	1.2
		06/07/11	< 0.071	6.8	0.30 Jq	1.9	0.29 Jq	1.1
MW-15	MCEA	12/09/11	< 0.071	5.8	0.25 Jq	1.6	0.28 Jq	1.0
		05/31/12	< 0.071	6.7	0.34 Jq	1.8	0.26 Jq	1.1
		11/16/12	< 0.071	6.2	0.32 Jq	1.7	0.29 Jq	0.9
		01/06/11	1.2	3.7	0.15 Jq	1.0	< 0.18	0.97
		06/07/11	1.3	4.3	0.15 Jq	0.93	< 0.18	0.76
MW-18	MCEA	12/09/11	0.72	3.7	0.14 Jq	0.88	<0.18	0.83
		05/31/12	2.1	3.8	0.14 Jq	1.1	< 0.18	1.1
		11/16/12	3.0	4.5	0.18 Jq	1.3	< 0.18	1.0
		01/07/11	0.55	< 0.10	< 0.098	0.15 Jq	< 0.18	< 0.25
		06/06/11	< 0.071	1.2	< 0.098	< 0.12	< 0.18	< 0.25
MW-67	MCEA	12/08/11	< 0.071	1.1	< 0.098	< 0.12	<0.18	< 0.25
		05/29/12	< 0.071	1.2	< 0.098	< 0.12	<0.18	< 0.25
		11/16/12	< 0.071	0.92	< 0.098	< 0.12	< 0.18	< 0.25
		01/04/11	< 0.071	0.17 Jq	< 0.098	< 0.12	<0.18	< 0.26
		06/06/11	< 0.071	0.15 Jq	< 0.098	<0.12	< 0.18	< 0.25
MW-100	DG	12/12/11	< 0.071	0.18 Jq	<0.098	<0.12	<0.18	< 0.25
		05/29/12	< 0.35	0.21	< 0.098	<0.12	<0.18	<0.25
		11/16/12	< 0.071	0.23	<0.098	<0.12	< 0.18	<0.25
MCI	MCL/DWNL (µg/L)			1 (1)	5	6	6	5

Notes:

MCEA - Massacre Canyon Entrance Area.

DG – Downgradient

MCL - California Department of Public Health maximum contaminant level

DWNL - California Department of Public Health drinking water notification level

(1) DWNL

μg/L - Micrograms per liter

Bold - MCL or DWNL exceeded.

- <# Analyte not detected; method detection limit concentration is shown.</p>
- \boldsymbol{J} The analyte was positively identified, but the analyte concentration is an estimated value.
- \boldsymbol{q} The analyte detection was below the practical quantitation limit (PQL).

3.6.2 **Increasing Trend Wells**

During the Second Quarter 2011 statistical trend analyses, (Tetra Tech, 2011), 21 monitoring

wells were designated as having increasing or probably increasing trends. Based on the magnitude

of the trend and the wells' locations, nine of these wells were included in the Fourth Quarter 2012

semiannual sampling event. The portion of the site where these wells are located, the location

identification, and the chemicals of potential concern that have the increasing trend are listed

below:

Two wells are located in the BPA.

MW-60A: perchlorate, TCE, 1,1-DCE, and 1,4-dioxane

MW-60B: TCE and 1,4-dioxane

Four wells are located in the RMPA.

IW-04: TCE and 1,1-DCE

MW-68: perchlorate, 1,1-DCE, and 1,4-dioxane

MW-88: perchlorate

MW-98B: TCE and 1,1-DCE

One well is located in the NPCA.

• MW-103: perchlorate

Two wells are located in the MCEA.

F34-TW1: 1,4-dioxane

MW-93: perchlorate

These increasing trend wells were chosen based on the Second Quarter 2011 trend analyses, and

may change pending approval of the First Quarter 2012 and Second Quarter 2012 Groundwater

Monitoring Report (Tetra Tech 2012c). Table 3-12 presents a summary of the detected chemicals

of potential concern reported in increasing trend well samples collected during Fourth Quarter

2012, and previous monitoring events.

Table 3-12 Summary of Detected Chemicals of Potential Concern in Increasing Trend Wells

Sample Location	Site Area	Sample Date	Perchlorate	1,4-Dioxane	1,1- Dichloroethene	Trichloroethene
-		All results rep	orted in µg/L unless ot	herwise stated		
		06/11/10	< 0.071	5.5	< 0.12	0.55
		06/06/11	2.1	9.6	0.23 Jq	0.76
F34-TW1	MCEA	12/08/11	< 0.071	11	0.13 Jq	0.85
		6/13/2012	< 0.35	4.1	0.28 Jq	0.86
		11/26/2012	< 0.071	6.6	0.44 Jq	1.6
		01/17/11	< 0.071	23	25	18
		06/14/11	< 0.071	25	35	23
IW-04	RMPA	12/08/11	< 0.071	21	17	23
		6/7/2012	< 0.071	23	14	21
		11/19/2012	0.27	21	9.6	14
		01/06/11	4,700	140	430	290
		06/21/11	4,900	130	460	280
MW-60A	BPA	12/12/11	4,700	130	380	280
		6/22/2012	4,900	150	370	270
		11/29/2012	6,600	110	420	330
		01/06/11	1,100	11	43	14
		06/21/11	1,300	8.8	41	12
MW-60B	BPA	12/12/11	900	8.8	34	13
		6/22/2012	1,100	10	39	15
		11/29/2012	1,800	15	52	22
		01/19/11	14,000	21	5.7	9.2
		06/21/11	13,000	25	6.1	< 0.25
MW-68	RMPA	12/12/11	6,000	11	2.6	< 0.25
		6/15/2012	9,100	18	4.4	< 0.25
		11/28/2012	17,000	21	6.3	< 0.25
		06/09/10	1,100	0.35	< 0.12	< 0.17
		06/17/11	6,100	0.23	< 0.12	< 0.25
MW-88	RMPA	12/12/11	740	0.23	< 0.12	< 0.25
		6/15/2012	770	0.20	< 0.12	< 0.25
		11/19/2012	610	0.19 Jq	< 0.12	< 0.25
		06/11/10	10 Jd	12	0.28 Jq	1.1
		06/07/11	8.4	4.4	< 0.12	< 0.25
MW-93	MCEA	12/12/11	6.7	14	0.56	1.7
		5/31/2012	4.9	19	1.1	3.1
		11/19/2012	5.5	22	1.0	3.2
		01/05/11	1,400	13	23	39
		06/17/11	1,600	5.2	12	17
MW-98B	RMPA	12/09/11	470	2,2	9.6	12
		6/5/2012	570	2.0	11	12
		11/28/2012	1,700	11	11	26
		06/09/11	160	11	3.6	5.9
		08/22/11	180	14	6.6	8.4
MW-103	NPCA	12/07/11	76	15	5.6	10
	1.1 011	6/12/2012	120	15	4.8	8.6
		11/27/2012	31	14	3.7	11
MCL/DWNL (µg/L)		1	6	1 (1)	6	5
Notes:			, ,	- \-/	<u> </u>	

Notes:

MCL - California Department of Public Health maximum contaminant level

DWNL - California Department of Public Health drinking water notification level.

(1) DWNL MCEA - Massacre Canyon Entrance Area.

µg/L - Micrograms per liter. NPCA - Northern Potrero Creek Area.

Bold - MCL or DWNL exceeded. RMPA - Rocket Motor Production Area.

BPA - Burn Pit Area.

- <# Analyte not detected; method detection limit concentration is shown.
- J The analyte was positively identified, but the analyte concentration is an estimated value.
- d The Laboratory Control Sample (LCS) recovery was outside control limits.
- q The analyte detection was below the practical quantitation limit (PQL).

3.6.3 New Wells

New monitoring wells MW-110, MW-111A, MW-111B, MW-111D, MW-111E, and MW-112A through MW-112C were sampled during Fourth Quarter 2012. These wells were installed in three angle boreholes in the BPA as part of the Site 1 Hydraulic Testing Investigation (Tetra Tech, 2012a). MW-110 was installed with a 2-inch diameter single completion pre-pack well across a zone of flowing sands. MW-111A through MW-111E, and MW-112A through MW-112C were installed with Water FLUTeTM multilevel monitoring systems in the MEF sandstone. MW-111C was unable to be sampled due to sediment clogging of the Water FLUTeTM port at this depth interval. Table 3-13 identifies the well designation and the screen or sampling port location. Complete well construction information appears in Appendix C.

Table 3-13 New Well Construction Information

Monitoring Port	Screen/Port	Screen/Port					
Well ID	Linear Depth	Vertical Depth					
	(Feet BGS)	(Feet BGS)					
MW-110	110 to 130	95.3 to 112.6					
MW-111A	180 to185	155.9 to 160.2					
MW-111B	161 to 166	139.4 to 143.8					
MW-111C	131 to 136	113.4 to 117.8					
MW-111D	121 to 126	104.8 to 109.1					
MW-111E	97 to 102	84.0 to 88.3					
MW-112A	125 to 130	108.3 to 112.6					
MW-112B	100 to 105	86.6 to 90.9					
MW-112C	87 to 92	75.3 to 79.7					

Notes:

BGS - Below ground surface

The addition of these wells helps to characterize the Mount Eden sandstone and contaminant concentrations with depth, and provides additional hydraulic data to support the evaluation of remedial alternatives at the site. It is expected to take a longer period than normal for wells MW-111A through MW-111E and wells MW-112A through MW-112C to fully equilibrate with the adjacent formation and produce representative groundwater samples, due to (1) the extremely low permeability in the BPA and (2) the fact that wells with the Water FLUTeTM system installed cannot be developed following traditional well development procedures. Additional purging cycles are being performed on the wells between sampling events to speed up the process of equilibration. A summary of the sample results from Fourth Quarter 2012 and previous sampling events appears in Table 3-14.

Table 3-14 Summary of Detected Chemicals of Potential Concern in New Wells

Sample Location	Sample Date	Per- chlorate	1,4- Dioxane	1,1- Dichloro- ethane	1,2- Dichloro- ethane	1,1- Dichloro- ethene	c-1,2- Dichloro- ethene	1,1,2- Trichloro- ethane	Trichloro- ethene
	T		All r	esults reported i	n μg/L unless o	therwise stated			
	02/27/12	2.0	0.24 Ba	< 0.098	<0.21	< 0.12	< 0.18	< 0.31	< 0.25
MW-110	06/25/12	7.0	< 0.10	< 0.098	<0.21	< 0.12	< 0.18	< 0.31	< 0.25
14144 110	08/22/12	7.7	< 0.10	< 0.098	< 0.21	< 0.12	< 0.18	< 0.31	< 0.25
	11/19/12	14	< 0.10	< 0.098	< 0.21	< 0.12	< 0.18	< 0.31	< 0.25
	02/28/12	35,000	3,800	170	470	4,800	74	110	3,300
MW-111A MW-111B MW-111C MW-111D MW-111E MW-112A	06/21/12	3,300	2,900	32	100	1,300	13	32	510
WIW-IIIA	08/21/12	2500	2,800 Je	48	130	1,800	18	39	610
	11/20/12	2,600	1,900	< 0.098	130	2,000	19	34	720
	02/28/12	17,000	6,100	140	390	5,700	55	120	3,200
MW-111B	06/21/12	20,000	7,000	170	440	7,600	67	120	3,500
	08/21/12	25,000	7,300 Je	230	550	9700	95	140	4,800
	11/20/12	36,000	5,800	240	590	10,000	110	190	4,400
	02/28/12	18,000	6,400	120	370	4,100	50	110	2,500
MW-111C	06/21/12	21,000	7,300	170	440	7,700	69	120	3,400
	08/31/12	25,000	6,800	240	600	12,000	100	160	5,400
	02/28/12	20,000	6,300	140	400	5,200	55	120	2,600
MW 111D	06/21/12	21,000	8,400	150	430	6,900	63	130	3,300
WIW-IIID	08/21/12	23,000	6,600 Je	230	560	10,000	85	150	5,400
	11/20/12	38,000	7,900	310	680	14,000	120	150	6,500
	02/28/12	7,700	4,800	45	170	1,400	20	57	670
MW 111E	06/21/12	35,000	4,900	270	560	12,000	110	110	5,700
MW-IIIE	08/21/12	36,000	6,100 Je	330	670	16,000	120 Jq	160 Jq	7,000
	11/20/12	49,000	4,800	430	840	20,000	170	170	9,200
	02/29/12	1,300	46	5.8	9.6	69	1.3	1.2	67
MW 112 A	06/21/12	4,300	66	9.2	13	220	1.4	1.7	220
IVI VV -1 1 ZA	08/20/12	4,300	63	11	16	270	1.7	1.9	250
	11/20/12	5,800	64	9.7	14	260	1.5	1.7	270
	02/29/12	4,800	71	15	18	310 Jf	2.4	2.4	340 Jf
MW 112D	06/21/12	4,900	66	16	18	370	2.3	2.2	340
1V1 VV -1 1∠B	08/20/12	4,500	70	17	21	410	2.6	2.4	350
	11/20/12	6,800	72	17	20	430	2.8	2.2	420
	02/29/12	4,300	59	10	14	210	1.6	1.6	230
MW-112C	06/21/12	2,500	62	8.5	13	140	1.5	1.0	100
1V1 VV -112C	08/20/12	3,000	59	6.3	12	120	1.4	0.85	87
	11/20/12	4,700	66	8.9	14	170	2.1	1.1	150
MCL/DWN	IL (µg/L)	6	1 (1)	5	5	6	6	5	5

Notes:

MCL - California Department of Public Health maximum contaminant level

DWNL - California Department of Public Health drinking water notification level

(1) DWNL

μg/L - micrograms per liter

Bold - MCL or DWNL exceeded.

- <# Analyte not detected; method detection limit concentration is shown.</p>
- $B\ -\ The\ result\ is < 5\ times\ the\ blank\ contamination.\ Cross\ contamination\ is\ suspected\ and\ the\ data\ are\ considered\ unusable.$
- J The analyte was positively identified, but the analyte concentration is an estimated value.
- a The analyte was found in the method blank.
- e A holding time violation occurred.
- \boldsymbol{f} The duplicate Relative Percent Difference was outside the control limit.
- q The analyte detection was below the practical quantitation Limit (PQL).

3.6.4 Surface Water

Surface water samples were scheduled for collection from 17 locations during the Fourth Quarter 2012 groundwater sampling event. Table 3-15 presents concentrations of chemicals of potential concern reported in surface water samples collected from this sampling event.

During Fourth Quarter 2012 surface water samples were collected from five locations (SW-02, SW-03, SW-04, SW-09, and SW-18) along the Potrero and Bedsprings creek drainages. The remaining 12 locations were dry at the time of sampling. Because surface water location SW-16 was dry, an attempt was made to collect a sample from the alternate location SW-17, but it was also dry and therefore was not sampled. The four primary chemicals of potential concern (1,4-dioxane, 1,1-DCE, TCE, and perchlorate) and four secondary chemicals of potential concern (1,1-DCA, cis-1,2-dichloroethene, 1,1,1-TCA, and vinyl chloride) were detected in surface water samples collected from locations SW-02, SW-03, and SW-04. These samples were collected from springs and/or manmade surface depressions fed by nearby springs located outside of the stream beds but near the intersection of Bedsprings and Potrero creeks.

1,4-Dioxane was the only chemical of potential concern detected in the surface water samples collected from locations SW-09 and SW-18. These samples were collected from water flowing in Potrero Creek and are topographically downgradient of the springs discussed in the previous paragraph. Figure 3-11 presents concentrations of chemicals of potential concern reported in surface water samples collected from the Fourth Quarter 2012 monitoring event.

Table 3-15 Summary of Detected Chemicals of Potential Concern in Surface Water - Fourth Quarter 2012

Sample Location	Sample Date	Per- chlorate	1,4- Dioxane	1,1- Dichloro- ethane	1,1- Dichloro- ethene	c-1,2- Dichloro- ethene	1,1,1- Trichloro- ethane	Trichloro ethene	Vinyl Chloride
	•		All results	reported in µg/	L unless otherv	vise stated			
SW-02	11/15/12	100	10	0.40 Jq	9.8	9.8 0.75		13	0.38 Jq
SW-03	11/15/12	3.0	11	< 0.098	1.1	0.42 Jq	< 0.12	1.1	< 0.13
SW-04	11/15/12	3.6	4.3	< 0.098	1.0	0.25 Jq	< 0.12	1.2	< 0.13
SW-09	11/15/12	< 0.071	2.9	< 0.098	< 0.12	< 0.18	< 0.12	< 0.25	< 0.13
SW-18	11/15/12	< 0.071	3.8	<0.098	<0.12	<0.18	<0.12	< 0.25	< 0.13
Method Detection Limit (μg/L)		0.071	0.10	0.098	0.12	0.18	0.12	0.25	0.13
MCL/DWNL (µg/L)		6	1(1)	5	6	6	200	5	0.5

Notes:

 $\mu g/L$ - Micrograms per liter

MCL - California Department of Public Health maximum contaminant level

DWNL - California Department of Public Health drinking water notification level

(1) DWNL

Bold - MCL or DWNL exceeded.

- <# Analyte not detected; method detection limit concentration is shown.</p>
- J The analyte was positively identified, but the analyte concentration is an estimated value.
- \boldsymbol{q} The analyte detection was below the practical quantitation limit (PQL).

3.7 CONTAMINANT ATTENUATION SAMPLING

A site-wide contaminant attenuation evaluation was completed in the Spring of 2012 (Tetra Tech, 2012b). The evaluation concluded that approximately 95% of the contaminant flux was being attenuated, and that contaminant attenuation was going to be an important component of any remedial strategy implemented at this site. The attenuation is primarily occurring in the riparian corridor located at the confluence of Bedsprings and Potrero creeks, with approximately 22% of the attenuation being due to evapotranspiration and 72% being due to biodegradation. The evaluation concluded that approximately 95% of the contaminant mass flux was being attenuated, primarily in the riparian corridor located in the NPCA and MCEA. Although the bulk of the contaminant attenuation is taking place in this area, the study also found that contaminant attenuation is also occurring at Features F-33, F-34, and F-39. The contaminant attenuation evaluation recommended the routine monitoring of the degradation processes and aquifer redox conditions in up to 22 monitoring wells. Following is a discussion of the results of the Fourth Quarter 2012 contaminant attenuation monitoring.

Monitoring wells scheduled for contaminant attenuation parameter sampling during the Fourth Quarter 2012 monitoring event included 15 monitoring wells (MW-05, MW-08, MW-13, MW-43, MW-48, MW-76B, MW-101, MW-103, MW-104, MW-105, MW-106, MW-107, MW-109, OW-02, and P-02) located in the NPCA and one monitoring well (MW-102) located in the MCEA.

Samples for laboratory analysis were collected for alkalinity, chloride, dissolved organic carbon (DOC), dissolved manganese, sulfate, nitrate, methane, ethane, ethene, hydrogen, and volatile organic compounds (VFAs). Dissolved oxygen (DO) and oxidation-reduction potential (ORP) were monitored with field instruments during purging and sampling, and ferrous iron and sulfide were analyzed using a field instrument prior to sample collection. Figure 3-13 shows the locations of the monitoring wells sampled for contaminant attenuation parameters during the Fourth Quarter 2012 monitoring event. Table 3-16 presents a summary of the field measurements and validated analytical results.

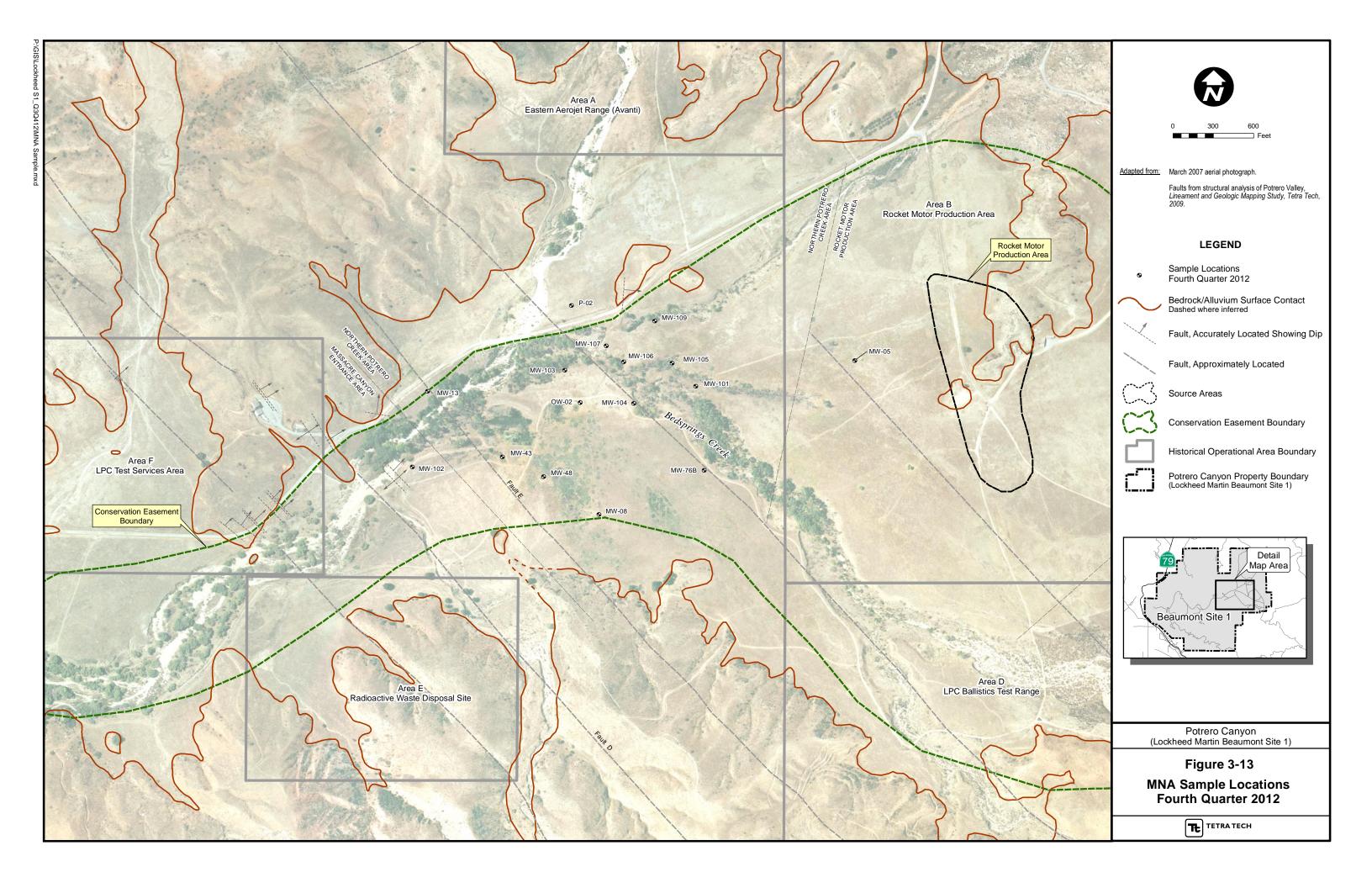


Table 3-16 Summary of Validated Detected Contaminant Attenuation Analytes and Field Measurements - Fourth Quarter 2012

			Field I	Parameters		Analytes											
Sample Location	Sample Date	DO - mg/L	ORP -mVs	Sulfide -mg/L (1)	Ferrous Iron - mg/L (1)	Per- chlorate -ug/L	Acetic Acid - mg/L	2- Hydroxy- propanoic Acid - mg/L	Hexanoic Acid - mg/L	Dissolved Organic Carbon - mg/L	Hydrogen -nM	Methane -ug/L	Nitrate (as N) -mg/L	Sulfate -mg/L	Chloride -mg/L	Iron - mg/L	Dissolved Manganese -ug/L
MW-05	11/28/12	0.48	-137.2	-	-	2,900	< 0.021	< 0.032	< 0.039	0.36 Jq	< 0.074	< 0.044	7.9	11	9.5	< 0.0023	1.4 Jq
MW-08	11/26/12	0.31	-154.3	0.00	0.41	< 0.071	< 0.021	< 0.032	< 0.039	0.27 Jq	0.66	0.29	< 0.11	11	5.1	1.4	190
MW-13	11/26/12	2.61	-6.1	0.00	0.04	0.36	0.076	< 0.032	< 0.039	0.70	< 0.074	< 0.044	1.4	79	9.6	0.0048 Jq	< 0.24
MW-43	11/26/12	0.25	-119.5	0.01	0.11	54	< 0.021	< 0.032	< 0.039	0.29 Jq	2.2	210	0.77	8.7	5.8	0.096	4.5 Jq
MW-48	11/28/12	0.47	-122.3	0.01	1.15	< 0.071	< 0.021	< 0.032	< 0.039	1.1	< 0.074	1,700	< 0.11	6.3	7.9	1.9	250
MW-76B	11/28/12	0.40	-99.0	0.00	0.22	< 0.071	< 0.021	< 0.032	< 0.039	0.31 Jq	< 0.074	4.2	< 0.11	8.4	5.6	0.22	11
MW-101	11/28/12	0.31	-106.2	0.00	0.48	< 0.071	0.076	0.22	< 0.039	0.39 Jq	< 0.074	0.20	< 0.11	28	7.1	1.7	110
MW-102	11/28/12	0.65	-122.0	0.00	0.06	< 0.071	< 0.021	< 0.032	< 0.039	0.49 Jq	< 0.074	17	< 0.11	16	6.3	0.08	56
MW-103	11/27/12	2.81	20.4	0.00	0.01	31	< 0.021	< 0.032	< 0.039	1.1	1.0	7.0	< 0.11	6.9 Jd	7.5	2.1 Jcf	11
MW-104	11/27/12	0.70	-141.3	0.00	0.32	< 0.071	0.098	< 0.032	< 0.039	0.70	0.98	1.2	< 0.11	4.0	7.3	0.4	98
MW-105	11/27/12	0.47	-164.6	0.04	0.14	< 0.071	0.12	< 0.032	< 0.039	0.64 Jq	0.60	10	< 0.11	27	11	0.12	860
MW-106	11/27/12	0.79	-154.2	0.00	0.04	< 0.071	0.075	< 0.032	< 0.039	2.7	1.1	2.4	< 0.11	63	17	0.0051 Jq	120
MW-107	11/27/12	1.18	-177.5	0.01	0.02	53	< 0.021	< 0.032	< 0.039	1.7	1.5	1.2	< 0.11	26	14	0.012 Jq	62
MW-109	11/27/12	0.35	-153.3	0.00	0.03	310	< 0.021	< 0.032	0.44	0.48 Jq	1.4	4.4	1.3	23	9.3	0.0058 Jq	320
OW-02	11/28/12	2.97	-68.3	0.00	0.03	350	0.075	< 0.032	< 0.039	0.33 Jq	2.1	< 0.044	3.2	7.5	7.1	0.0066 Jq	0.26 Jq
P-02	11/26/12	0.40	-147.6	0.00	0.02	0.32	< 0.021	< 0.032	< 0.039	1.3	< 0.074	0.37	0.72	490	76	1.2	790
MI	DL	-	-	-	-	0.071	0.021	0.032	0.039	0.16	0.074	0.044	0.11	0.37	1.0	0.0023	0.24
MCL/E	OWNL	-	-	-	0.3	6	-	-	-	-	-	-	10	250	250	0.3	500 (2)

Only analytes positively detected are presented in this table. For a complete list, refer to the laboratory data package in Appendix H.

Notes:

(1) - Sulfide and ferrous iron sample analysis was performed in the field using a Hach DR 850 colorimeter.

(2) - DWNL

MCL - California Department of Public Health Services maximum contaminant level

DWNL - California Department of Public Health Services drinking water notification level

J - The analyte was positively identified, but the analyte concentration is an estimated value.

c - The Matrix Spike (MS) and/or Matrix Spike Duplicate (MSD) recoveries were outside control limits.

f - The duplicate Relative Percent Difference was outside the control limit.

q - The analyte detection was below the practical quantitation limit (PQL).

<# - Analyte not detected, method detection limit concentration is shown.</p>

MDL - Method detection limit

mg/L - milligrams per liter

 $\mu g/L$ - micrograms per liter

nM - NanoMoles

Bold - MCL or DWNL exceeded

"-" - Not available

Dissolved Oxygen and General Minerals

Software developed by the United States Geological Survey (USGS, 2009) was used to evaluate aquifer redox conditions based on dissolved oxygen, nitrate, manganese, iron, sulfate, and sulfide concentrations in the groundwater. The results of this analysis are provided in Appendix J. With the exception of monitoring wells MW-13 and OW-02, the aquifer redox category was anoxic. Wells for which redox category changed with respect to the Second Quarter 2012 conditions include

- MW-05 (changed from oxic to anoxic),
- MW-13 (changed from anoxic to oxic),
- MW-102 (changed from anoxic to mixed oxic/anoxic),
- MW-103 (changed from oxic to mixed oxic/anoxic),
- MW-104 (changed from anoxic to mixed oxic/anoxic),
- MW-105 (changed from anoxic to mixed anoxic),
- MW-107 (changed from anoxic to mixed oxic/anoxic), and
- P-02 (changed from anoxic to mixed anoxic).

It should be noted that in all of these cases, the redox classification changes are attributable to relatively small changes in the DO concentrations measured in the field.

Oxidation-Reduction Potential

Oxidation/reduction potential values are a general indicator of aquifer oxidation state. Table 3-16 shows that with the exception of monitoring well MW-103, ORP values are negative, consistent with the redox conditions necessary for perchlorate biodegradation (Lieberman and Borden, 2008).

Methane

Methane can be produced in an aquifer by a number of anaerobic biologic pathways, including fermentative breakdown of acetate to methane and carbon dioxide, and biological carbon dioxide reduction. In general, the presence of methane is indicative of highly-reduced aquifer conditions. Table 3-16 shows that methane was detected in all wells except for MW-05, MW-13, and OW-02,

which is consistent with the previously observed anoxic aquifer conditions in the riparian corridor located in the NPCA and MCEA.

Dissolved Hydrogen

Dissolved hydrogen is produced in anoxic environments by fermentative microorganisms, and is consumed by nitrate-, iron-, sulfate-, and carbon dioxide-reducing microorganisms. The efficiency of hydrogen utilization varies among these microorganisms, with nitrate-reducing bacteria being the most efficient, followed by iron-, sulfate-, and carbon dioxide-reducers. Thus, steady-state hydrogen concentrations in anoxic aquifers are lowest under nitrate-reducing conditions, and become progressively higher as conditions become more reducing. Table 3-16 shows that hydrogen was detected in nine of the monitoring wells at concentrations ranging from 0.60 to 2.1 nanoMoles (nM). These findings are consistent with anoxic aquifer conditions.

Dissolved Organic Carbon

Dissolved organic carbon was detected in all of the monitoring wells, at concentrations ranging from 0.27 to 2.7 milligrams per liter (mg/L) (Table 3-16). Volatile fatty acids are organic carbon molecules that are immediately available to the microorganisms involved in contaminant biodegradation. The presence of VFAs in groundwater, even at low concentrations, indicates that there is an excess of organic carbon available for biodegrading microorganisms. Detectable concentrations of VFAs were observed in seven of the monitoring wells (Table 3-16).

Perchlorate

The trend analysis conducted during Second Quarter 2012 indicates that only one of the monitoring wells sampled for contaminant attenuation parameters (MW-103) had an increasing or probably increasing perchlorate concentration trend. This is a relatively new well, which was installed in late 2010, and has been sampled six times. As a new well, the well was sampled quarterly and then, as specified in the groundwater monitoring sampling and analysis plan (Tetra Tech, 2003b), it was reclassified as a plume monitoring well and the sampling frequency was adjusted. Review of the analytical data used for the trend analysis suggests that perchlorate concentrations fluctuate seasonally, peaking in the fall and summer, and decreasing during the winter and spring. The perchlorate concentrations from the summer 2011 and 2012 events are similar, suggesting stability rather than an increasing trend. The DO and ORP appear to fluctuate

seasonally as well, most likely as a result of the influx of oxygenated water after storm events. Because the statistical analysis showed an increasing trend for perchlorate in this well, this well will be sampled semiannually. Future sampling events should provide more clarity on perchlorate trends in this well.

3.8 HABITAT CONSERVATION

Consistent with the United States Fish and Wildlife Service (USFWS)-approved Habitat Conservation Plan (HCP) (USFWS, 2005) and subsequent clarifications (Lockheed Martin Corporation, 2006a, 2006b, and 2006c) of the HCP describing activities for environmental remediation at the site, field activities were performed under the supervision of a USFWS-approved biologist. No impact to the Stephens' kangaroo rat occurred during the performance of field activities related to the Third Quarter 2012 and Fourth Quarter 2012 monitoring events.

SECTION 4 SUMMARY AND CONCLUSIONS

This section summarizes the results of the Third Quarter 2012 and Fourth Quarter 2012 groundwater monitoring events. A total of 182 groundwater level measurements were collected for the Third Quarter 2012 and Fourth Quarter 2012 water quality monitoring events. Four wells were observed to be dry during the Third Quarter and five wells were observed to be dry during the Fourth Quarter. For the Third Quarter 2012 monitoring event, 9 monitoring wells were proposed and sampled for water quality monitoring. For the Fourth Quarter 2012 monitoring event, 56 sampling locations (17 surface water, one alternate surface water location, and 38 monitoring wells) were proposed for water quality monitoring. Twelve surface water sample locations were not sampled because the locations were dry. The alternate surface water location SW-17 was also dry and was not sampled. Additionally one monitoring well, MW-111C, was unable to be sampled because this depth interval in the Water FLUTe™ system was clogged with sediment. Therefore, water quality data were collected from five surface water and 37 monitoring well locations.

4.1 GROUNDWATER ELEVATIONS

The Beaumont National Weather Service station reported approximately 0.30 inches of precipitation during Third Quarter 2012, and approximately 3.62 inches of precipitation during Fourth Quarter 2012. During this same period, groundwater elevations generally decreased approximately 2.54 feet across the site.

Groundwater elevations during the Third Quarter 2012 and Fourth Quarter 2012 monitoring events ranged from approximately 2,155 feet mean sea level (msl) upgradient of the former Burn Pit Area to approximately 1,789 feet msl in the Massacre Canyon Entrance Area.

Groundwater elevation differences in all wells from quarter to quarter appear to depend on the short- and long-term weather patterns. In general, the greatest differences in quarterly groundwater elevations occur during periods of seasonal precipitation. Wells in the Northern Potrero Creek Area and the Massacre Canyon Entrance Area appear to respond most quickly to precipitation compared to the former Burn Pit Area and Rocket Motor Production Area, which generally show a one-season lag before responding to seasonal precipitation. However, wells near Bedsprings Creek

just south of the Burn Pit Area also show rapid responses to precipitation due to surface water infiltration and mountain front recharge. The response also diminishes in each area with depth and distance from the Potrero and Bedsprings creeks. The site has experienced overall groundwater level declines since 2005.

4.2 SURFACE WATER FLOW

During the Third Quarter 2012 and Fourth Quarter 2012, Tetra Tech personnel walked the Potrero and Bedsprings creek riparian corridors to determine the presence, nature, and quantity of surface water in the creek beds. The locations where surface water was encountered were plotted and a determination was made whether the water was flowing or stagnant. At specific locations where flowing water was encountered, the flow rate was determined using a modified version of the *United States Environmental Protection Agency Volunteer Stream Monitoring: A Methods Manual* (USEPA, 1997).

Four fixed stream locations were previously chosen for stream flow measurements: SF-1, which is near Gilman Hot Springs at the southwest border of the site; SF-2, which is near MW-67; SF-3, which is near MW-15 and MW-18; and SF-4, which is near MW-101.

During Third Quarter 2012 and Fourth Quarter 2012 the four stream flow measurement locations, (SF-1 through SF-4), were dry at the time of measurement.

4.3 GROUNDWATER FLOW AND GRADIENTS

Groundwater flow directions from Third Quarter 2012 and Fourth Quarter 2012 were similar to previously observed patterns for a dry period. Generally, groundwater flows northwest from the southeastern limits of the valley (near the former Burn Pit Area) beneath the former Rocket Motor Production Area, toward Potrero Creek where groundwater flow then changes direction and begins heading southwest, parallel to the flow of Potrero Creek, into Massacre Canyon.

Between June 2012 (Second Quarter 2012) and September 2012 (Third Quarter 2012), the overall groundwater gradient (approximating a flowline from MW-36, upgradient of the Burn Pit Area, through the Rocket Motor Production Area and Northern Potrero Creek Area to MW-18, in the Massacre Canyon Entrance Area) remained constant at 0.013 feet/foot. Between September 2012 (Third Quarter 2012) and December 2012 (Fourth Quarter 2012), the overall groundwater gradient through the same flow path remained constant at 0.013 feet/foot. In general the horizontal gradient

was lowest between the Burn Pit Area and the Rocket Motor Production Area with a greatly increased flow through the Northern Potrero Creek Area and the Massacre Canyon Entrance Area. The flattening of the gradient in the Burn Pit Area and Rocket Motor Production Area appears to be attributable to the lithology, aquifer transmissivity, and aquifer thickness in these areas.

Vertical groundwater gradients between shallow and deeper monitoring well pairs are generally downward (negative) in the Burn Pit Area, Rocket Motor Production Area, and the Northern Potrero Creek Area, and upward (positive) in the Massacre Canyon Entrance Area. The response to seasonal changes in groundwater recharge, although dampened by depth, are consistent within the different vertical well pairs installed at the site. This suggests that there is vertical hydraulic communication within the aquifer.

4.4 WATER QUALITY

Both groundwater and surface water are collected and sampled as part of the Groundwater Monitoring Program. The program has a quarterly/semiannual/annual/biennial sampling frequency. The annual and biennial events are larger major monitoring events, and the quarterly and semiannual events are smaller minor events. All new wells are sampled quarterly for one year. The semiannual wells are sampled second and fourth quarters of each year, the annual wells are sampled second quarter of even-numbered years.

An evaluation of chemicals of potential concern is performed annually, and reported in the First and Second Quarter Semiannual Groundwater Monitoring Report. The chemicals of potential concern previously identified for the site during the 2011 evaluation, (Tetra Tech, 2011), are consistent with those identified in the 2012 evaluation (Tetra Tech, 2012c). The primary chemicals of potential concern identified for the site are perchlorate, 1,1-dichloroethene, trichloroethene, and 1,4-dioxane. The secondary chemicals of potential concern identified for the site are 1,1-dichloroethane, 1,2-dichloroethane, 1,1-trichloroethane, 1,1,2-trichloroethane, cis-1,2-dichloroethene, and vinyl chloride. The 2012 evaluation yielded no additions or deletions to the list of chemicals of potential concern. The results of surface and groundwater samples collected and tested during the Third and Fourth Quarter 2012 monitoring events are discussed below.

4.4.1 Surface Water

Surface water samples are collected semiannually during the second and fourth quarter sampling events, and during a storm event. Seventeen surface water sample locations and one alternate sample location have been identified for semiannual surface water sampling at the site. Sample locations have been chosen to include springs and manmade surface depressions fed by nearby springs, ephemeral ponds, and locations in the Bedsprings and Potrero creek drainages. Twelve locations in the active drainages and one ephemeral pond location have been identified for surface water sampling during a storm event. It is common for many of the locations to be dry at the time of sampling, due to the ongoing drought conditions and the ephemeral nature of the ponds and creeks.

During the Fourth Quarter 2012 sampling event, surface water samples were collected from five locations. The remaining 12 locations were dry at the time of sampling. Because surface water location SW-16 was dry, an attempt was made to collect a sample from the alternate location SW-17, but it was also dry and therefore not sampled. The sample results from the locations sampled are consistent with previous sample results obtained at the site.

The four primary chemicals of potential concern (1,4-dioxane, 1,1-dichloroethene, trichloroethene, and perchlorate) and four secondary chemicals of potential concern (1,1- dichloroethane, cis-1,2-dichloroethene, 1,1,1-trichloroethane, and vinyl chloride) were detected in surface water samples collected from locations SW-02, SW-03, and SW-04. These samples were collected from springs or surface depressions, in an area of discharging groundwater, located outside of the stream beds but near the intersection of Bedsprings and Potrero creeks, downgradient from the Rocket Motor Production Area.

Surface water sample locations SW-09 and SW-18 are in areas of flowing water in Potrero Creek, topographically downgradient of the springs and surface depressions discussed in the previous paragraph. 1,4-Dioxane was detected at concentrations exceeding the drinking water notification level (1 microgram per liter $[\mu g/L]$) in the surface water samples collected from these locations. No other chemicals of potential concern were detected in these locations during Fourth Quarter 2012.

The analyte 1,4-dioxane was detected above the drinking water notification level in all surface water samples collected during the Fourth Quarter 2012 sampling event. Additionally, perchlorate, 1,1-dichloroethene, and trichloroethene were detected above their respective maximum contaminant levels in the surface water sample collected from SW-02. Sample results for the surface water samples collected during Fourth Quarter 2012 are consistent with results from previous sampling events.

In general, the concentrations of chemicals of potential concern in surface water are highest in the area of the surface depressions, an area of discharging groundwater; the concentrations decrease rapidly to at or near the method detection limit, as one moves downgradient through the riparian zone toward the property boundary. The concentration gradient of 1,4-dioxane in surface water samples, however, is much smaller and appears to be less affected by movement through the riparian zone, since it is not biodegraded like the other chemicals of potential concern.

4.4.2 Groundwater

Groundwater monitoring wells were sampled during the third and fourth quarters. The third quarter event included the quarterly sampling of newly installed wells. The fourth quarter event included the quarterly sampling of newly installed wells and the semiannual sampling of increasing contaminant trend wells, guard wells, and contaminant attenuation wells (Tetra Tech, 2003b).

New Wells

Monitoring wells MW-110, MW-111A through MW-111E, and MW-112A through MW-112C were installed as part of the hydraulic testing in 2011 (Tetra Tech, 2012a) to help characterize the Mount Eden sandstone and contaminant concentrations with depth, and to provide additional hydraulic data to support the evaluation of remedial alternatives at the site. MW-110 was installed with a 2-inch diameter single completion pre-pack well across a zone of flowing sands. MW-111A through MW-111E, and MW-112A through MW-112C were installed with Water FLUTeTM multilevel monitoring systems. The results from MW-111A through MW-111E, and MW-112A through MW-112C are not considered representative at this time, due to (1) the extremely low permeability in the Burn Pit Area, and (2) the fact that wells with Water FLUTeTM system installed cannot be developed following traditional well development procedures.

Guard Wells

Four monitoring wells are designated as guard wells: MW-15, MW-18, MW-67, and MW-100. Wells MW-15 and MW-18 are a clustered well pair upstream of the Large Motor Washout Area (Feature F-33). All guard wells are located along Potrero Creek, downgradient of the source areas (Burn Pit Area and Rocket Motor Production Area). Well MW-18 is completed near the top of the alluvial aguifer and MW-15 is completed near the bottom of the alluvial aguifer. These wells are approximately three miles from the southern site boundary and are upgradient of the secondary sources identified at F-33, F-34, and F-39. Well MW-67, is the farthest downgradient site well and is approximately 0.9 miles upgradient of the southern site boundary. MW-100, an off-site well, is approximately 500 feet south of the southern site boundary near the mouth of Potrero Creek. Both MW-67 and MW-100 are located below the secondary sources identified at F-33, F-34, and F-39. The analyte 1,4-dioxane was detected in monitoring wells MW-15, MW-18, MW-67, and MW-100 at concentrations of 6.2 µg/L, 4.5 µg/L, 0.92 µg/L, and 0.23 µg/L, respectively. The analyte 1,4-dioxane was the only chemical of potential concern to be detected above the California Department of Public Health maximum contaminant level or drinking water notification level in these guard wells during the Fourth Quarter 2012 sampling event. The drinking water notification level for 1,4-dioxane is 1 µg/L. Sample results for the guard wells from Fourth Quarter 2012 are consistent with results from previous sampling events.

Increasing Trend Monitoring Wells

The number of increasing or probably increasing trend wells decreased from 24 wells in 2010 to 21 wells in the 2011 temporal trend analyses. The temporal trend analyses have been performed using data from Second Quarter 2002 to Second Quarter 2012. This period was chosen because operation of the Rocket Motor Production Area groundwater pump and treat system was discontinued in 2002. The 2011 temporal trend analysis updates the analysis performed following completion of the Second Quarter 2010 monitoring event (Tetra Tech, 2011). The temporal trends were analyzed using Mann-Kendall and linear regression methods. A summary of the trend analysis results for the 21 increasing or probably increasing trend locations is presented in Table 4-1. The percent change that these increases represent with respect to the mean of the data used to calculate each trend is also presented in Table 4-1. Twelve of the wells had trend magnitudes with less than a 20% change. Based on the magnitude of the trend (greater than a 20% change with respect to the mean) and the wells' location, nine increasing trend wells were included in Fourth Quarter 2012 semiannual sampling event. These locations are shaded in Table 4-1.

During this time period, the number of locations identified as having either a decreasing or probably decreasing trend has increased, and the number of wells identified as having no trend has decreased. Due to the new non-detect designation, several locations that were previously identified as having a stable trend have been reclassified, thus reducing the number of wells identified as having a stable trend.

In general, the plume morphology has remained stable with only slight modifications due to new wells better defining the lateral extent of the plume. The majority of the wells and the surface water locations display a stable trend, no trend, or are non-detect.

Possible reasons for the change in the number of increasing trend wells are: (1) the number of new wells that have been installed in the last several years (60 new wells since 2006); (2) longer monitoring histories at the individual locations, trends become more noticeable due to the ability to better define outliers; and (3) as time passes, changes created by operation of the former extraction system are dissipating.

In general, the chemicals of potential concern concentrations in the sampled increasing trend wells are consistent with previous results. These wells will continue to be sampled on a semiannual basis until the magnitude of the trend is less than a 20% change, or until the well is reclassified in the annual temporal trend analysis.

Table 4-1 Summary of Increasing Chemicals of Potential Concern Trends

Analyte:	Perchlorate			1,1-Dichloroethene			Trichloroethene			1,4-Dioxane		
Sample	T1	Magnitude	Magnitude	T 1	Magnitude	Magnitude		Magnitude	Magnitude	T	Magnitude	Magnitude
Location	Trend	(%/yr)	(µg/L/yr)	Trend	(%/yr)	(μg/L/yr)	Trend	(%/yr)	(µg/L/yr)	Trend	(%/yr)	(μg/L/yr)
Burn Pit Ar		1						1				
MW-07	No Trend			No Trend			No Trend			Increasing	6.6	0.04
		2.2	4 4 4	Probably		0.4.5.4					•	12.25
MW-26	Decreasing	-2.2	-166.44	Increasing	2.6	94.54	No Trend	_		Increasing	2.9	12.26
MW-60A	Increasing	23.7	1115.08	Increasing	4.9	17.74	Increasing	6.0	13.25	Increasing	4.9	5.42
							Probably					
MW-60B	Decreasing	-4.0	-56.21	Stable			Increasing	1.5	0.18	Increasing	25.6	1.33
MW-61B	Stable			No Trend			No Trend			Increasing	2.7	12.59
MW-61C	No Trend			Increasing	13.0	12.96	Increasing	11.3	2.49	Increasing	4.9	0.29
				Probably								
MW-71B	No Trend			Increasing	15.3	0.04	No Trend			Non-detect		
D. J. M.	D 1 di A											
	or Production Area		100.04	T	21.0	2.20	T	12.0	1.00	C4-1-1-		1
IW-04	Decreasing	-67.5	-108.04	Increasing	21.9	3.29	Increasing	12.8	1.66	Stable		
MW-05	ъ .		112.42	G. 11			Probably		5.70	NI TE 1		
	Decreasing	-5.1	-112.42	Stable			Increasing	6.6	5.78	No Trend		
MW-09	N. 1.			N7 1			N7 1			Probably	2.6	0.12
	Non-detect			Non-detect			Non-detect			Increasing	2.6	0.13
MW-19	0.11			Probably		0.00				a. 11		
	Stable			Increasing	1.2	0.33	No Trend			Stable		
				Probably		4.00						
MW-28	No Trend			Increasing	7.7	1.38	No Trend			No Trend		
MW-68	Increasing	21.9	1489.20	Increasing	29.2	2.83	No Trend			Increasing	34.7	3.22
	Probably											
MW-88	Increasing	56.6	735.48	Stable			No Trend			No Trend		
	Probably											
MW-91	Increasing	12.2	256.78	No Trend			No Trend			Increasing	15.3	0.25
MW-98B	No Trend			Increasing	20.1	2.61	Increasing	16.1	3.69	No Trend		
	trero Creek Area											
MW-76A	Non-detect			Non-detect			Non-detect			Increasing	18.1	0.38
MW-103	Increasing	346.8	221.92	Not Available			Not Available			Stable		
	anyon Entrance Are	ea		T			T			T		
F34-TW1	No Trend			No Trend			No Trend			Increasing	23.7	1.16
MW-70										Probably		
	No Trend	1		Decreasing	-27.4	-0.07	Decreasing	-13.3	-0.02	Increasing	5.3	0.15
	Probably			l						l		1
MW-93	Increasing	45.6	1.92	Stable		ļ	Stable		ļ	Stable		<u> </u>
SW-07	Stable			Non-detect			Non-detect			Increasing	4.6	0.04
				1		I			I	1		1
Notes:												

Shading indicates that based on the magnitude of the trend and the well location, the well was included in the Fourth Quarter 2012 sampling event. $\mu g/L/yr$ - Micrograms per liter per year %/yr - Percent change per year

4.5 CONTAMINANT ATTENUATION SAMPLING

A site-wide contaminant attenuation evaluation was completed in the Spring of 2012 (Tetra Tech, 2012b). The evaluation concluded that approximately 95% of the contaminant flux was being attenuated, and that contaminant attenuation was going to be an important component of any remedial strategy implemented at this site. The attenuation is primarily occurring in the riparian corridor located at the confluence of Bedsprings and Potrero creeks, with approximately 22% of the attenuation being due to evapotranspiration and 72% being due to biodegradation. The contaminant attenuation evaluation recommended the routine monitoring of degradation progress and aquifer redox conditions in up to 22 monitoring wells. With the exception of a few minor differences, the redox conditions observed in 2012 are consistent with the conditions observed in 2011 and documented in the contaminant attenuation evaluation finalized in 2012. The redox conditions and the contaminant trends observed indicate that conditions are essentially unchanged from the 2011 baseline, and that contaminant attenuation is continuing to occur.

4.6 PROPOSED CHANGES TO THE GROUNDWATER MONITORING PROGRAM

4.6.1 Groundwater Sampling Frequency

The sampling frequency of a monitoring well is based on the well's classification (i.e., its function) (Tetra Tech, 2003b). Groundwater monitoring well classifications are based on the evaluation of the temporal trends, spatial distribution, and other qualitative criteria. There are seven different well classifications. Currently no wells are designated as remedial monitoring wells, because remedial actions are not yet in place. A summary of the sampling frequency by well classification is presented in Table 4-2.

Table 4-2 Well Classification and Sampling Frequency

Classification	Sampling Frequency
Horizontal Extent (Plume) Wells	Annual
Vertical Distribution Wells	Biennial
Increasing Trend Wells	Semiannual
Remedial Monitoring Wells	Semiannual
Guard Wells	Semiannual
Redundant Wells	Suspend
New Wells	Quarterly

4.6.2 Proposed Changes

The analytical scheme is evaluated annually during the second quarter of each year, and changes may be proposed then to accommodate expanded site knowledge or changing site conditions. The classification of the wells in the network and their corresponding sampling frequency are also evaluated annually during the second quarter of each year, and are modified as needed. No unusual events or observations occurred during this reporting period that require consideration of modifying the monitoring program.

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