Semiannual Groundwater Monitoring Report Fourth Quarter 2009 and First Quarter 2010 Lockheed Martin Corporation, Beaumont Site 2 Beaumont, California



Prepared for:

LOCKHEED MARTIN

Prepared by:







Department of Toxic Substances Control

Maziar Movassaghi, Acting Director 5796 Corporate Avenue Cypress, California 90630



August 24, 2010

Ms. Denise Kato Remediation Analyst Senior Staff Lockheed Martin Corporation Energy, Environment, Safety & Health 2950 North Hollywood Way, Suite 125 Burbank, California 91505

SEMI-ANNUAL GROUNDWATER MONITORING REPORT, FOURTH QUARTER 2009 AND FIRST QUARTER 2010, LOCKHEED MARTIN CORPORATION, BEAUMONT SITE 2, BEAUMONT, CALIFORNIA (Site Code: 400261)

Dear Ms. Kato:

The Department of Toxic Substances Control (DTSC) has reviewed your responses to our comments regarding the subject groundwater monitoring report and found them to be acceptable. The subject groundwater monitoring report is approved.

Should you have any questions or comments, please contact me at (714) 484-5483.

Sincerely,

Daniel K Zogaib **Project Manager**

Brownfields and Environmental Restoration Program

CC: Mr. Gene Matsushita

Senior Manager

Environmental Remediation Lockheed Martin Corporation

Energy, Environment, Safety & Health 2950 North Hollywood Way, Suite 125

Burbank, California 91505





Department of Toxic Substances Control

Maziar Movassaghi, Acting Director 5796 Corporate Avenue Cypress, California 90630



August 17, 2010

Ms. Denise Kato Remediation Analyst Senior Staff Lockheed Martin Corporation Energy, Environment, Safety & Health 2950 North Hollywood Way, Suite 125 Burbank, California 91505

SEMI-ANNUAL GROUNDWATER MONITORING REPORT, FOURTH QUARTER 2009 AND FIRST QUARTER 2010, LOCKHEED MARTIN CORPORATION, BEAUMONT SITE 2, BEAUMONT, CALIFORNIA (Site Code: 400261)

Dear Ms. Kato:

The Department of Toxic Substances Control (DTSC) has reviewed the subject groundwater monitoring report. Enclosed are comments from DTSC's Geological Services Unit (GSU).

Please address the enclosed GSU comments by September 17, 2010.

Should you have any questions or comments, please contact me at (714) 484-5483.

Sincerely,

Daniel K. Zogaib

Project Manager

Brownfields and Environmental Restoration Program

See next page. CC:

Ms Denise Kato August 17, 2010 Page 2 of 2

Mr. Gene Matsushita CC:

Senior Manager

Environmental Remediation Lockheed Martin Corporation

Energy, Environment, Safety & Health 2950 North Hollywood Way, Suite 125 Burbank, California 91505





Department of Toxic Substances Control

Maziar Movassaghi, Acting Director 5796 Corporate Avenue Cypress, California 90630



TO:

Daniel Zogaib

Hazardous Substances Engineer

Brownfields and Environmental Restoration Program

FROM:

Dina Kourda, CEG CM___ **Engineering Geologist**

Geological Services Unit (GSU)

Cypress Office

DATE:

July 28, 2010 (August 11, 2010- FINALIZED)

SUBJECT: REVIEW OF SEMIANNUAL GROUNDWATER MONITORING REPORT 4Q.

2009 and 1Q 2010, LOCKHEED MARTIN CORPORATION, BEAUMONT

SITE 2. BEAUMONT, CALIFORNIA, DATED JUNE 9, 2010.

PCA 11050

SITE CODE 400261-00

Tracking # 1040017

At the request of the DTSC Project Manager, Mr. Daniel Zogaib, the Geological Services Unit (GSU) has reviewed the subject document received on July 13, 2010 for Lockheed Martin Corporation, Site 2 in Beaumont.

BACKGROUND

On behalf of Lockheed Martin Corporation (LMC), Tetra Tech, Inc. (Tetra Tech) prepared the subject document, for LMC's former Beaumont Site 2 (Jack Rabbit Trail) facility (the "Site") The Site is a 2,668-acre parcel located southwest of the city of Beaumont in San Bernardino County, California. Grand Central Rocket Company (GCR) purchased the site from the US Government in 1958. The Site was utilized for small rocket motor assembly, testing operations, propellant incineration, and minor disposal activities from 1958 to 1974, when Site closure took place under Lockheed.

According to Tetra Tech, chemicals of concern (COCs) include the following six: perchlorate, trichloroethene (TCE), methylene chloride, bis-(2-ethylhexyl) phthalate, Royal Demolition Explosives (RDX), and arsenic. Arsenic and bis-(2-ethylhexyl) phthalate were also identified, however, arsenic is a likely related to background and bis-(2-ethylhexyl) phthalate is likely a laboratory contaminant, according to Tetra Tech. Perchlorate has been identified as the primary COC. TCE, methylene chloride, and RDX are considered secondary COCs.

The subject document summarizes the results of the Fourth Quarter 2009 and First Quarter 2010 groundwater monitoring events conducted at the Site.

Mr. Daniel Zogaib

Semiannual Groundwater Monitoring Report, 4Q 2009 and 1Q 2010

Site 2, Lockheed Martin, Beaumont

August 11, 2010

Page 2 of 2

The following comments are based on a GSU critical flaw review.

SPECIFIC COMMENTS

- Section 3.3, Page 3-6 and Figure 3-3: Because of Site 2's unique configuration, it should be explained how horizontal flow directions are calculated. It appears that the gradient flow direction shifts slightly three times between TT-MW2-16 and TT-MW2-6S. Also, it should be specified if the groundwater gradient(s) is/are calculated site-wide or by area.
- 2 Section 3.6, Page 3-23: All new groundwater well construction detail/soil boring logs should be included as an appendix.

If you have any questions or comments, please do not hesitate to contact me at (714) 484-5408 or dkourda@dtsc.ca.gov

Peer reviewed by: Greg Neal, PG

cc: Fred Zanoria, CEG, CHg

RESPONSES TO DTSC COMMENTS ON THE SEMIANNUAL GROUNDWATER MONITORING REPORT FOURTH QUARTER 2009 AND FIRST QUARTER 2010, SITE 2 LOCKHEED MARTIN, BEAUMONT, CALIFORNIA TETRA TECH, INC AUGUST 2010

Specific Comments							
Comment	Response	Proposed Action					
1. Section 3.3, Page 3-6 and Figure 3-3: Because of Site 2's unique configuration it should be explained how horizontal flow directions are calculated. It appears that the gradient flow direction shifts slightly three times between TT-MW2-16 and TT-MW2-6S. Also, it should be specified if the groundwater gradient(s) is/are calculated site-wide or by area.	As alluded to in this comment, most of the monitoring wells at Site 2 lie essentially along a line, which precludes determination of groundwater flow direction by triangulation. The conceptual hydrogeologic model for the site suggests that shallow groundwater flow is focused within deeply weathered San Timoteo formation underlying the major canyons. This conceptual model is consistent with data showing very low to non-detectable perchlorate concentrations in monitoring wells TT-MW2-27 and TT-MW2-34A/B/C, which were installed in small side canyons off Laborde Canyon immediately south (generally downgradient) of areas with very high perchlorate concentration in groundwater. Assuming that the canyon walls approximate a no-flow boundary, the equipotential lines are drawn perpendicular to the sides of the canyons. Groundwater gradients are calculated using a segmented path approximating a flowline through the canyon. Appendix E of the report details the groundwater gradient calculation for both the entire site and individual segments along the flowline.	This will be explained in future reports.					
2. Section 3.6, Page 3-23: All new groundwater well construction detail/soil boring logs should be included as an appendix.	Groundwater monitoring well construction detail and soil boring logs are included in the well installation / site characterization report that details the well installation. The well construction table located in the appendix is updated with each groundwater monitoring report.	No further action is proposed.					

RESPONSES TO DTSC COMMENTS ON THE SEMIANNUAL GROUNDWATER MONITORING REPORT FOURTH QUARTER 2009 AND FIRST QUARTER 2010, SITE 2 LOCKHEED MARTIN, BEAUMONT, CALIFORNIA TETRA TECH, INC AUGUST 2010

REFERENCES

Tetra Tech, Inc. 2010. Dynamic Site Investigation and Summary Remedial Investigation Report, Lockheed Martin Corporation, Beaumont Site 2, Beaumont, California. April.





Department of Toxic Substances Control

Maziar Movassaghi, Acting Director 5796 Corporate Avenue Cypress, California 90630



August 24, 2010

Ms. Denise Kato Remediation Analyst Senior Staff Lockheed Martin Corporation Energy, Environment, Safety & Health 2950 North Hollywood Way, Suite 125 Burbank, California 91505

SEMI-ANNUAL GROUNDWATER MONITORING REPORT, FOURTH QUARTER 2009 AND FIRST QUARTER 2010, LOCKHEED MARTIN CORPORATION, BEAUMONT SITE 2, BEAUMONT, CALIFORNIA (Site Code: 400261)

Dear Ms. Kato:

The Department of Toxic Substances Control (DTSC) has reviewed your responses to our comments regarding the subject groundwater monitoring report and found them to be acceptable. The subject groundwater monitoring report is approved.

Should you have any questions or comments, please contact me at (714) 484-5483.

Sincerely,

Daniel K Zogaib **Project Manager**

Brownfields and Environmental Restoration Program

CC: Mr. Gene Matsushita

Senior Manager

Environmental Remediation Lockheed Martin Corporation

Energy, Environment, Safety & Health 2950 North Hollywood Way, Suite 125

Burbank, California 91505

Lockheed Martin Corporation, Shared Services Energy, Environment, Safety and Health 2950 North Hollywood Way, Suite 125 Burbank, CA 91505 Telephone: 818.847.0197 Facsimile: 818.847.0256



June 9, 2010

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, Fourth Quarter 2009 and First Quarter 2010, Lockheed Martin Corporation, Beaumont Site 2, Beaumont, California

Dear Mr. Zogaib:

Please find enclosed one hard copy of the body of the report and two compact disks with electronic copies of the report and appendices of the Semiannual Groundwater Monitoring Report, Fourth Quarter 2009 and First Quarter 2010, Lockheed Martin Corporation, Beaumont Site 2, Beaumont, California for your review and approval or comment.

If you have any questions regarding this submittal, please contact me at 408.756.9595 or denise.kato@Imco.com.

Sincerely,

Denise Kato

Remediation Analyst Senior Staff

Enclosure

Copy with Enc:

Gene Matsushita, LMC (CD and hard copy)
lan Lo, Camp, Dresser, McKee (CD)
Thomas J. Villeneuve, Tetra Tech, Inc. (CD and hard copy)

BUR121 Beau 2 Q409 Q110 GWMR

Semi-annual Groundwater Monitoring Report

Fourth Quarter 2009 and First Quarter 2010 Beaumont Site 2, Beaumont, California

Prepared for:

Lockheed Martin Corporation

Prepared by:

Tetra Tech, Inc.

June 2010

Christophel Patrick Environmental Scientist

,

Bill Muir, PG California (6762) Deputy Program Manager WILLIAM MUIR
No. 6762
Typ 7/31/11
FOR CALIFORNIA

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- Appendix A Recent Environmental Activities and Conceptual Site Model
- Appendix B Copies of the Field Data Sheets
- **Appendix C Well Construction Table**
- Appendix D Water Level Hydrographs and Precipitation Data
- **Appendix E Summary of Calculated Horizontal and Vertical Groundwater Gradients**
- Appendix F Validated Sample Results by Analytical Method
- Appendix G Laboratory Analytical Data Packages
- **Appendix H Consolidated Data Summary Table**
- **Appendix I COPC Time-Series Graphs**

Acronyms

ARCH air rotary casing hammer

bgs below ground surface

btoc below top of casing

BOS bottom of screen

CAM California Assessment Manual

CDHS California Department of Health Services

COPCs chemical(s) of potential concern

CSM Conceptual Site Model

DTSC Department of Toxic Substances Control

DWNL drinking water notification level

EC electrical conductivity

EPA United States Environmental Protection Agency

ft/ft feet per foot

ft/day feet per day

GCR Grand Central Rocket Company

GMP Groundwater Monitoring Program

HCP Habitat Conservation Plan

HSA hollow stem auger

HSUs hydrostratigraphic units

K hydraulic conductivity

LAC Lockheed Aircraft Corporation

LEBs Lockheed equipment blanks

LMC Lockheed Martin Corporation

LPC Lockheed Propulsion Company

LR Linear Regression

LTBs Lockheed trip blanks

MW monitoring well

MCLs maximum contaminant levels

MDLs method detection limits

mg/L milligrams per liter

MS matrix spike

MSD matrix spike duplicate

msl mean sea level

μg/L micrograms/liter

NA not applicable

NDMA N-nitrosodimethylamine

NWS National Weather Service

PW production well

PVC polyvinyl chloride

PZ piezometer

QAL Quaternary alluvium

QA/QC quality assurance/quality control

RDX Research Department composition X

SAP sampling and analysis plan

SKR Stephens' Kangaroo rat

SS stainless steel

STF San Timoteo formation

SVOCs semi-volatile organic compounds

TCE trichloroethene

1,2,3-TCP 1,2,3-trichloropropane

TOC top of casing

TOS top of screen

Unk. unknown

u-DMH unsymmetrical dimethyl hydrazine

U.S. United States

USFWS United States Fish and Wildlife Service

VOCs volatile organic compounds

WCA West Coast Analytical Services, Inc.

wSTF weathered San Timoteo formation

SECTION 1 INTRODUCTION

This Semi-annual Groundwater Monitoring Report (Report) has been prepared by Tetra Tech, Inc. (Tetra Tech), on behalf of Lockheed Martin Corporation (LMC), and presents the results of the Fourth Quarter 2009 and First Quarter 2010 groundwater quality monitoring activities for the Beaumont Site 2 (Site) Groundwater Monitoring Program (GMP). The Site is located southwest of the City of Beaumont, Riverside County, California (Figure 1-1). Currently, the Site is inactive with the exception of ongoing investigative activities performed under Consent Order (88/89 034) with the Department of Toxic Substances Control (DTSC).

The objectives of this Report are to:

- Briefly summarize the Site history;
- Document the water quality monitoring procedures and results;
- Analyze and evaluate the water quality monitoring data generated.

This Report is organized into the following sections: 1) Introduction, 2) Summary of Monitoring Activities, 3) Groundwater Monitoring results, and 4) Summary and Conclusions. A brief description of the previous site environmental investigations and the current conceptual site model (CSM) can be found in Appendix A.

1.1 SITE BACKGROUND

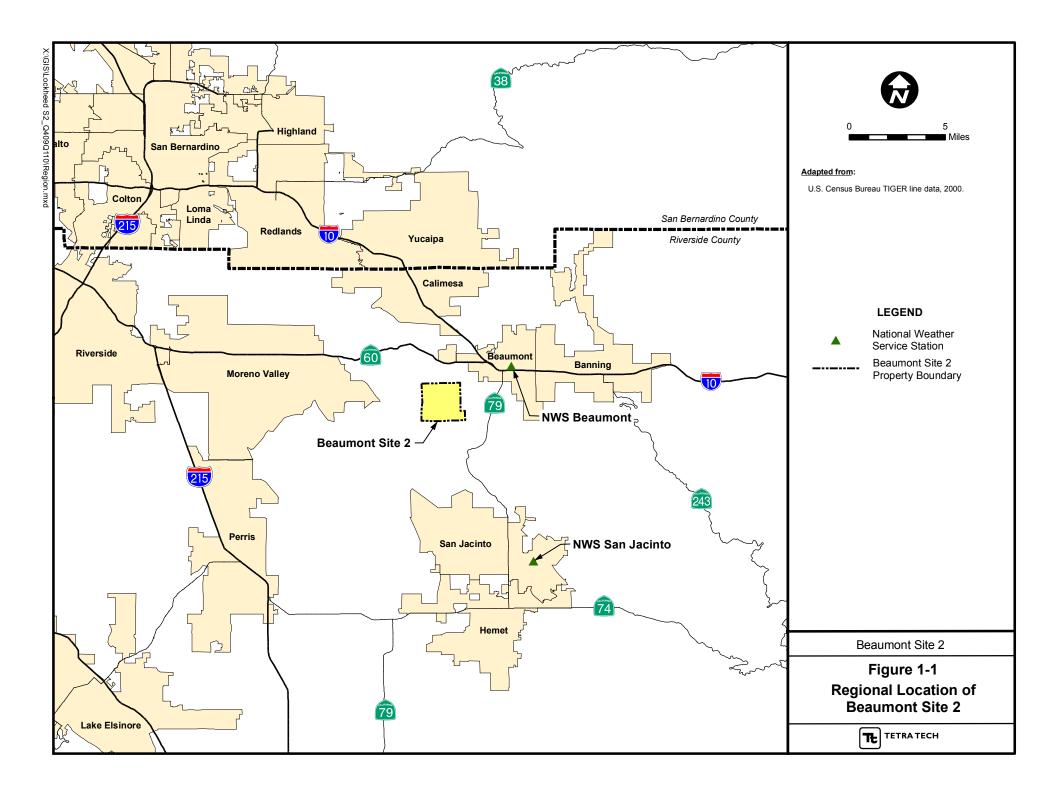
The Site is a 2,668 acre parcel located southwest of Beaumont, California. The parcels that comprise the Site were owned by individuals and the United States (U.S.) government prior to 1958. Between 1958 and 1960, portions of the Site were purchased by the Grand Central Rocket Company (GCR) and utilized as a remote test facility for early space and defense program efforts. In 1960, Lockheed Aircraft Corporation (LAC) purchased one-half interest in GCR. GCR became a wholly-owned subsidiary of LAC in 1961. The remaining parcels of land that comprise the Site were purchased from the U.S. government between 1961 and 1964. In 1963, Lockheed Propulsion Company (LPC) became an operating division of LAC and was responsible for the operation of the Site until its closure in 1974. The Site was utilized by GCR and LPC from 1958 to 1974 for

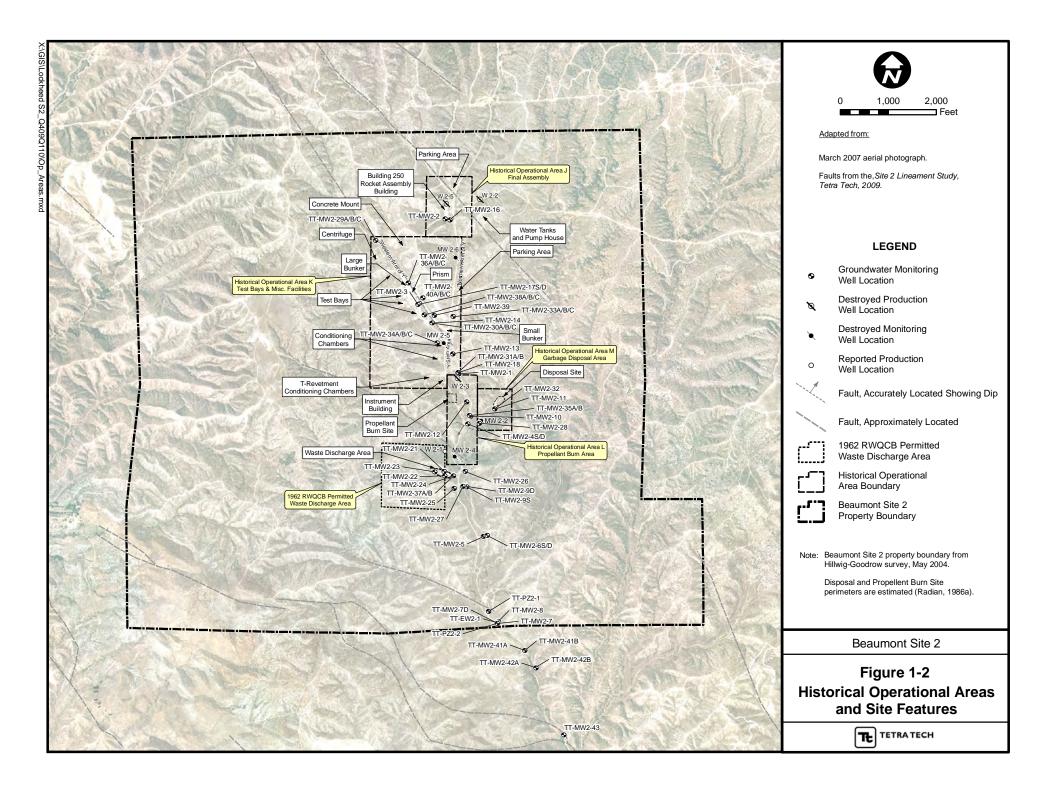
small rocket motor assembly, testing operations, propellant incineration, and minor disposal activities. Ogden Labs is known to have leased portions of the Site in the 1970s (Radian, 1986a).

In 1989, the DTSC issued a consent order requiring LMC to cleanup contamination at the Site related to past testing activities (CDHS, 1989). Based on investigative and cleanup activities performed at the Site, the DTSC issued a no further remedial action letter to LMC in 1993.

Based on regulatory interest in perchlorate and 1,4-dioxane, a groundwater sample was collected from an inactive groundwater production well (identified as W2-3) at the Site in January 2003. The sample was analyzed for volatile organic compounds (VOCs), perchlorate, and 1,4-dioxane to determine the potential presence and concentration of those chemicals in groundwater. The analytical results indicated that VOCs and 1,4-dioxane were not present at or above their respective method detection limits (MDLs). However, perchlorate was reported at a concentration of 4,080 micrograms per liter (μ g/L), which exceeded the California Department of Public Health drinking water notification level (DWNL) which existed at that time of 6 μ g/L. In October 2007 the DWNL was replaced by the California Department of Public Health Maximum Contaminant Level (MCL). Based on the detection of perchlorate in the groundwater sample collected, the DTSC reopened the Site for further assessment.

Four primary historical operational areas have been identified at the Site (Figure 1-2). Each operational area was responsible for various activities associated with rocket motor assembly, testing, and propellant incineration. A brief description of each operational area follows:





Historical Operational Area J (Area J) –Final Assembly

Rocket motor casings with solid propellant were transported to Building 250 where final assembly of the rocket hardware was conducted. The building was used from 1970 to 1974 for final assembly and shipment of short range attack missile rocket motors. Rocket motor assembly operations included installation of the nozzle and headcap, pressure check of the motor, installation of electrical systems, and preparations for shipment. During plant closure in 1974, all usable parts of this facility were dismantled, taken off the Site, and sold (Radian, 1986a).

<u>Historical Operational Area K (Area K) – Test Bays and Miscellaneous Facilities</u>

The primary features included a large earthen structure known as the "Prism," conditioning chambers, a centrifuge, and four test bays and two associated bunkers.

The Prism was reportedly built between 1984 and 1990 and was used to test radar by General Dynamics (Tetra Tech, 2007a). Details concerning construction of the Prism are not available, but it appears to have been constructed with soils from near the test bays.

The conditioning chambers were used to examine the effects of extreme temperatures on rocket motors and to meet specification requirements (Radian, 1986a). A centrifuge was located in the northwestern portion of Area K, where rocket motors were tested in order to determine if the solid propellant would separate from its casing under increased gravitational forces.

Previously, only three test bays were known; however, a former employee reported during a recent interview that a fourth test bay, located north of the other three bays, was also previously used in Area K (Tetra Tech, 2009). The initial testing activities had a history of explosions that destroyed complete test areas, especially during the period when GCR operated at the Site (Radian, 1986a). While vestiges from three test bays are currently visible at the Site, the fourth was reportedly destroyed by such an explosion during testing. Also reportedly, after motor failure, the area was checked to recover unburned propellant.

Historical Operational Area L (Area L) – Propellant Burn Area

Solid propellant was reportedly transported to the burn area and set directly on the ground surface for burning (Radian, 1986a). No pits or trenches were dug as part of the burning process. The solid propellant was saturated with diesel fuel to initiate combustion. Reportedly, the solid propellant would burn rapidly. There is no evidence or physical features that identify the precise location of

burning activities, and previous site investigations (Tetra Tech, 2005; Tetra Tech, 2010b) found no evidence of significant contamination in Area L.

Two production wells were located in this area (W2-1 and W2-3). W2-1 was reported to have been part of the agricultural homestead. The origin of W2-3 is unknown. The use of the wells is unknown.

Historical Operational Area M (Area M) – Garbage Disposal Site

A garbage disposal area was located adjacent to a small creek at the Site (Radian, 1986a). Scrap metal, paper, wood, and concrete materials were disposed of at the disposal site by LPC. Hazardous materials, including explosives and propellants, were never disposed of at the disposal site by LPC according to employee interviews. Ogden Labs, a company that tested valves and explosive items, also used this disposal site. Reportedly, Ogden Labs disposed hazardous waste at the disposal site. In 1972, a Lockheed Safety Technician was exposed to toxic vapors of unsymmetrical dimethyl hydrazine (u-DMH) from a pressurized gas container located within the disposal site. Based on potential exposure risks to occupants, LPC's safety group required Ogden Labs to take measures to remove any potentially hazardous materials at the disposal site. Shortly thereafter, a disposal company was contracted by Ogden Labs to clean up the disposal site (Radian, 1986a).

Waste Discharge Area

In 2007, LMC discovered the existence of Santa Ana River Basin Regional Water Pollution Control Board Resolution 62-24, dated September 14, 1962. Resolution 62-24 prescribed requirements for the "discharge of industrial wastes (rocket fuel residuum) to excavated pits." The discharge area was described as two shallow basins protected by two-foot berms, located in a small canyon on the western side of Laborde Canyon, in the SW ¼ of the NW ¼ of Section 19, Township 3 South, Range 1 West, San Bernardino Baseline and Meridian. Resolution 62-24 further describes the wastes to be discharged as "residue remaining after the manufacturing refuse is burned," and indicates that amount of material to be discharged was "approximately 5,000 gallons per year."

The exact nature of the waste proposed for discharge is not clear from the Resolution 62-24. The description of the waste material suggests that the area may have been used for burning propellant; but the description of the quantity of material to be discharged suggests that the waste may have

been liquid rather than solid. A 1961 aerial photograph shows the waste discharge area (WDA) as a large cleared area with roads leading to two circular structures, suggesting that the WDA was in use by 1961 (Tetra Tech, 2009a). Investigation of this area (Tetra Tech, 2007c; 2008a) found evidence for perchlorate impacts in both soil and groundwater.

Features remaining at the WDA include two roughly circular depressions surrounded by earthen berms, at the location of the circular structures identified in the 1961 aerial photograph.

SECTION 2 SUMMARY OF MONITORING ACTIVITIES

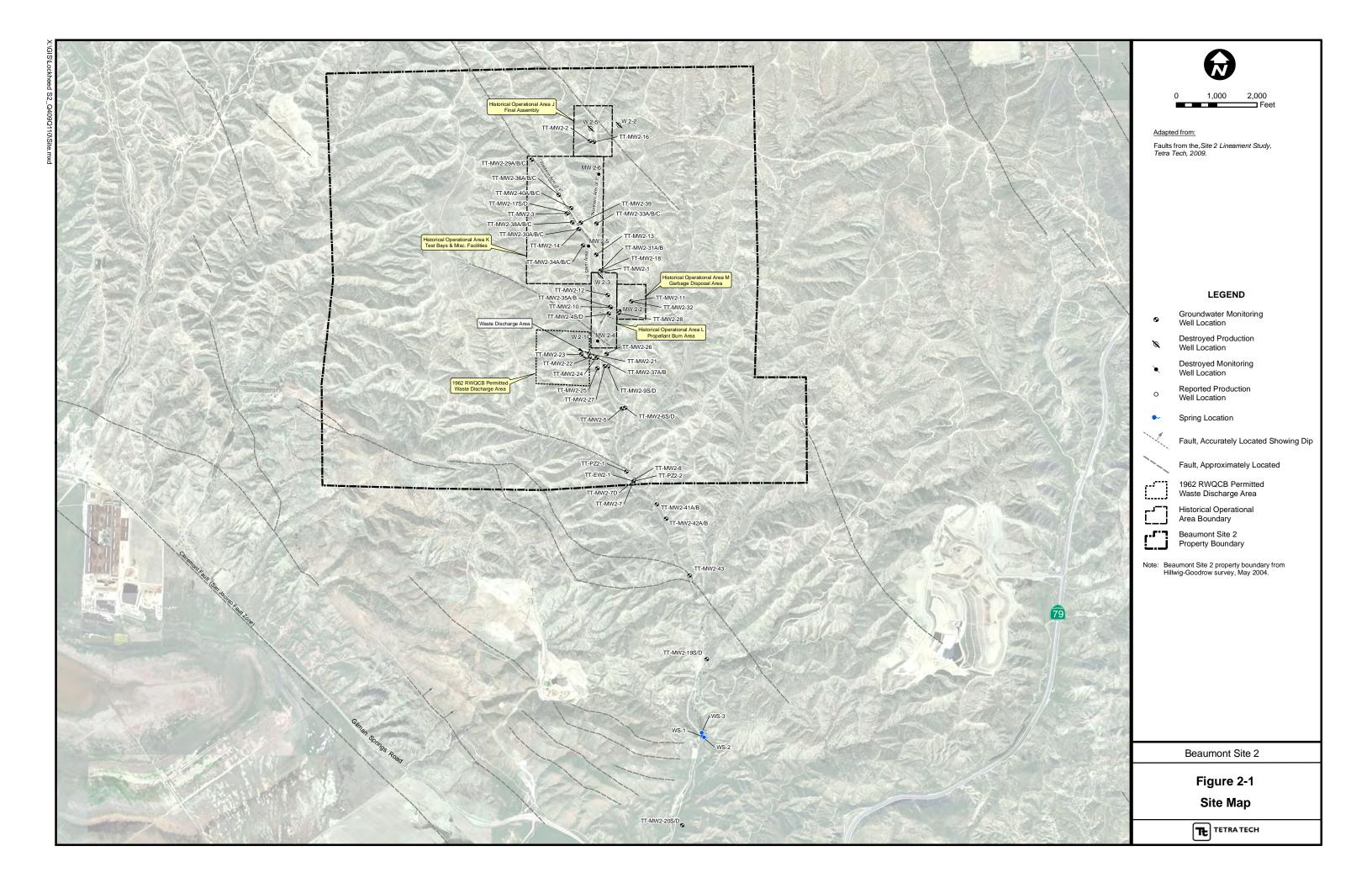
Section 2 summarizes the Fourth Quarter 2009 and First Quarter 2010 groundwater monitoring events conducted at the Site. The results from these monitoring events are discussed in Section 3.

2.1 GROUNDWATER LEVEL MEASUREMENTS

The Fourth Quarter 2009 groundwater level measurements were collected from 69 monitoring wells and two piezometers on 12 December 2009. The First Quarter 2010 groundwater level measurements were collected from 69 monitoring wells and two piezometers on 15 February 2010. Two monitoring wells, TT-MW2-29A and TT-MW2-43, were found to be dry during both quarters. Figure 2-1 presents a site map showing the well locations. Copies of the field data sheets from the water quality monitoring events are presented in Appendix B. A summary of well construction details is presented in Appendix C.

2.2 GROUNDWATER SAMPLING

The GMP has a quarterly, semi-annual, and annual frequency. Both groundwater and surface water are sampled as part of the GMP. The annual event is the major monitoring event and the quarterly and semi-annual events are smaller, minor events. All new wells are sampled quarterly for one year after which they are evaluated and reclassified. The semi-annual event includes, horizontal extent, vertical distribution, increasing contaminant, and guard wells, and is sampled during the second and fourth quarter of each year. In addition to the quarterly and semi-annual wells, the annual event includes background wells and takes place during the second quarter of each year. The groundwater monitoring schedule is reviewed and modified as necessary annually during the Second Quarter groundwater monitoring event. Modifications are done in accordance with the approved Groundwater Sampling and Analysis Plan (SAP) (Tetra Tech, 2007b). Fourth Quarter 2009 and First Quarter 2010 follow the schedule proposed in the Second and Third Quarter 2008 monitoring report (Tetra Tech, 2009b) which was presented to the DTSC in May 2009 and approved with no comments to the proposed schedule.



During the Fourth Quarter 2009 monitoring event 62 groundwater samples and three surface water samples were collected between 2 December and 17 December 2009; two additional locations, TT-MW2-29A and TT-MW2-43, were dry and were not sampled. During the First Quarter 2010 monitoring event six groundwater samples and two surface water samples were collected between 16 February and 17 February 2010. Additionally, storm water samples were collected during storm events on 27 January and 6 February 2010 from 5 locations. Tables 2-1 and 2-2 list the locations monitored for the Fourth Quarter 2009 and First Quarter 2010 monitoring events, analytical methods, sampling dates, and Quality Assurance/Quality Control (QA/QC) samples collected. Figures 2-2 and 2-3 illustrate the well locations sampled for the Fourth Quarter 2009 and First Quarter 2010 monitoring events, respectively. Groundwater sampling, analytical, and QA/QC procedures for the monitoring event were described in the Groundwater Monitoring Well Installation Work Plan (Tetra Tech, 2004a) and the SAP.

The following water quality field parameters were measured and recorded on field data sheets (Appendix B) during well purging activities: water level, temperature, pH, electrical conductivity (EC), turbidity, dissolved oxygen (DO) and oxidation reduction potential (ORP). Measurement of water quality parameters was initiated 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, and EC \pm 3%, turbidity < 10 nephelometric turbidity units (NTUs) (or \pm 10% if turbidity stabilizes at > 10 NTUs), DO \pm 0.3 mg/L and ORP \pm 10 mV. Sampling instruments and equipment were maintained, calibrated, and operated in accordance with the manufacturer's specifications, guidelines, and recommendations. Groundwater monitoring wells were purged and sampled using low-flow purging and sampling techniques with dedicated double valve sampling pumps or a portable bladder pump.

For the Fourth Quarter 2009 and First Quarter 2010 monitoring events, every effort was made to collect groundwater samples in order of increasing perchlorate and TCE concentration. Samples were placed in appropriate EPA method specified containers. A sample identification label was affixed to each sample container, and sample custody was maintained by a chain-of-custody record. Groundwater samples collected for the monitoring events were chilled and transported to

E. S. Babcock & Sons, Inc., a state-accredited analytical laboratory, via courier, thus maintaining proper temperatures and sample integrity. Trip blanks (LTBs) were collected on each day of the monitoring events to assess potential cross-contamination of water samples while in transit. Equipment blanks (LEBs) were collected when sampling with non-dedicated equipment to assess cross-contamination potential of water samples via sampling equipment.

2.3 SURFACE WATER SAMPLING

Storm water locations SW-01 through SW-07 are located in the ephemeral creek bed that runs through Laborde Canyon. Storm water runoff collects in the creek during periods of heavy precipitation and runs south through the Site and the former Wolfskill property, eventually crossing under Gilman Hot Springs Road. Water is present in the creek bed only during periods of heavy, prolonged precipitation. Surface water locations WS-1, WS-2 and WS-3 are spring locations on the former Wolfskill property. Water is generally present at one or more of these locations throughout the year.

During Fourth Quarter 2009, surface water samples were collected from three locations, WS-1, WS-2, and WS-3, and were analyzed for perchlorate. During First Quarter 2010, surface water samples were collected from two locations, WS-1 and WS-2, and were analyzed for perchlorate and general minerals. Additionally, storm water samples were collected from five locations, SW-02, SW-03, SW-05, SW-06, and SW-07, during storm events on 27 January and 6 February 2010. Storm water samples were analyzed for VOCs, and perchlorate. No other surface or storm water samples were collected during this reporting period. Figure 2-4 presents the surface and storm water sampling locations.

Table 2-1 Sampling Schedule and Analysis Method - Fourth Quarter 2009

Monitoring Well Location	Sample Date	VOCs (8260B)	Perchlorate (332.0)	1,4-Dioxane (8270C SIM)	Natural Attenuation Parameters	NDMA (E521)	Comments and QA /QC Samples
WS-1	12/02/09	-	X	-	-	-	Spring Sample
WS-2	12/02/09	-	X	-	-	-	Spring Sample
WS-3	12/02/09	-	X	-	-	-	Spring Sample
TT-MW2-1	12/09/09	-	X	-	X	-	Sample with Dedicated Pump, MS/MSD
TT-MW2-4S	12/04/09	-	X	-	-	-	Sample with Dedicated Pump
TT-MW2-5	12/3/2009 - 12/5/2009	-	X	X	X	-	Sample with Dedicated Pump
TT-MW2-6S	12/03/09	-	X	-	-	-	Sample with Dedicated Pump
TT-MW2-6D	12/03/09	-	X	-	-	-	Sample with Dedicated Pump
TT-MW2-7	12/03/09	-	X	-	X	-	Sample with Dedicated Pump
TT-MW2-7D	12/03/09	X	X	-	-	-	Sample with Dedicated Pump
TT-MW2-8	12/03/09	-	X	-	-	-	Sample with Dedicated Pump, Duplicate TT-MW2-8-Dup
TT-MW2-9S	12/08/09	-	X	X	X	-	Sample with Dedicated Pump
TT-MW2-9D	12/04/09	-	X	-	-	-	Sample with Dedicated Pump
TT-MW2-10	12/09/09	-	X	-	X	-	Sample with Dedicated Pump
TT-MW2-11	12/04/09	X	X	-		-	Sample with Dedicated Pump, Duplicate TT-MW2-11-Dup
TT-MW2-12	12/14/09	-	X	-	X	-	Sample with Dedicated Pump
TT-MW2-13	12/14/09	-	X	-	-	-	Sample with Dedicated Pump, Duplicate TT-MW2-13-Dup
TT-MW2-14	12/14/09	-	X	-	X	-	Sample with Dedicated Pump
TT-MW2-17S	12/17/09	-	X	-	-	-	Sample with Dedicated Pump
TT-MW2-17D	12/10/09	-	X	-	X	-	Sample with Dedicated Pump
TT-MW2-18	12/14/09	-	X	-	-	-	Sample with Dedicated Pump
TT-MW2-19S	12/11/09	-	X	X	-	X	Sample with Dedicated Pump, MS/MSD
TT-MW2-19D	12/02/09	-	X	-	-	X	Sample with Dedicated Pump
TT-MW2-20S	12/02/09	-	X	-	-	X	Sample with Dedicated Pump
TT-MW2-20D	12/02/09	-	X	-	-	X	Sample with Dedicated Pump
TT-MW2-21	12/04/09	X	X	X	-	X	Sample with Dedicated Pump
TT-MW2-22	12/04/09	X	X	X	-	-	Sample with Dedicated Pump, Duplicate TT-MW2-22-Dup
TT-MW2-23	12/11/09	X	X	-	X	_	Sample with Dedicated Pump
TT-MW2-24	12/11/09	X	X	X	X	X	Sample with Dedicated Pump, Duplicate TT-MW2-24-Dup
TT-MW2-25	12/04/09	X	X	-	-	X	Sample with Dedicated Pump
TT-MW2-26	12/04/09	X	X	X	-	X	Sample with Dedicated Pump
TT-MW2-27	12/03/09	X	X	-	-	-	Sample with Dedicated Pump
TT-MW2-28	12/04/09	X	X	-	-	X	Sample with Dedicated Pump
TT-MW2-29A	NA	-	-	-	-	-	Dry Well
	ple Locations:				67		
			ples Collected:				65
Notes: EPA -	United States Environme					NDMA -	

Notes: EPA - United States Environmental Protection Agency.

QA/QC - Quality assurance / quality control VOCs - Volatile Organic Compounds

NA - Not available

NDMA - N-Nitrosodimethylamine

MS / MSD- Matrix Spike / Matrix Spike Duplicate.

"-" Not analyzed

Table 2 1 Sampling Schedule and Analysis Method - Fourth Quarter 2009 (continued)

Monitoring Well Location	Sample Date	VOCs (8260B)	Per chlorate (332.0)	1,4-Dioxane (8270C SIM)	Natural Attenuation NDMA Parameters (E521)		Comments and QA /QC Samples		
TT-MW2-29B	12/07/09	X	X	-	-	X	Sample with Dedicated Pump		
TT-MW2-29C	12/07/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-30A	12/14/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-30B	12/11/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-30C	12/11/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-31A	12/09/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-31B	12/09/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-32	12/09/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-33A	12/11/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-33B	12/15/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-33C	12/15/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-34A	12/15/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-34B	12/15/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-34C	12/15/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-35A	12/09/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-35B	12/09/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-36A	12/10/09	X	X	-	X	X	Sample with Dedicated Pump, Duplicate TT-MW2-36A-Dup		
TT-MW2-36B	12/10/09	X	X	-	-	-	Sample with Dedicated Pump, MS/MSD		
TT-MW2-36C	12/10/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-37A	12/08/09	X	X	X	-	-	Sample with Dedicated Pump		
TT-MW2-37B	12/08/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-38A	12/14/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-38B	12/17/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-38C	12/14/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-39	12/15/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-40A	12/10/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-40B	12/10/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-40C	12/10/09	X	X	-	-	-	Sample with Dedicated Pump		
TT-MW2-41A	12/16/09	X	X	X	-	X	Sample with Dedicated Pump		
TT-MW2-41B	12/16/09	X	X	X	-	X	Sample with Portable Bladder Pump		
TT-MW2-42A	12/16/09	X	X	X	-	X	Sample with Portable Bladder Pump		
TT-MW2-42B	12/16/09	X	X	X	-	X	Sample with Portable Bladder Pump		
TT-MW2-43	NA	-	-	-	-	-	Dry Well		
	Fourth Qua	arter 2009: Sa	mple Locations:				67		
		Total Sar	nples Collected:				65		
Notes: EPA -	Notes: EPA - United States Environmental Protection Agency. NDMA - N-Nitrosodimethylamine								

Notes: EPA - United States Environmental Protection Agency.

QA/QC - Quality assurance / quality control VOCs - Volatile Organic Compounds

NA - Not available.

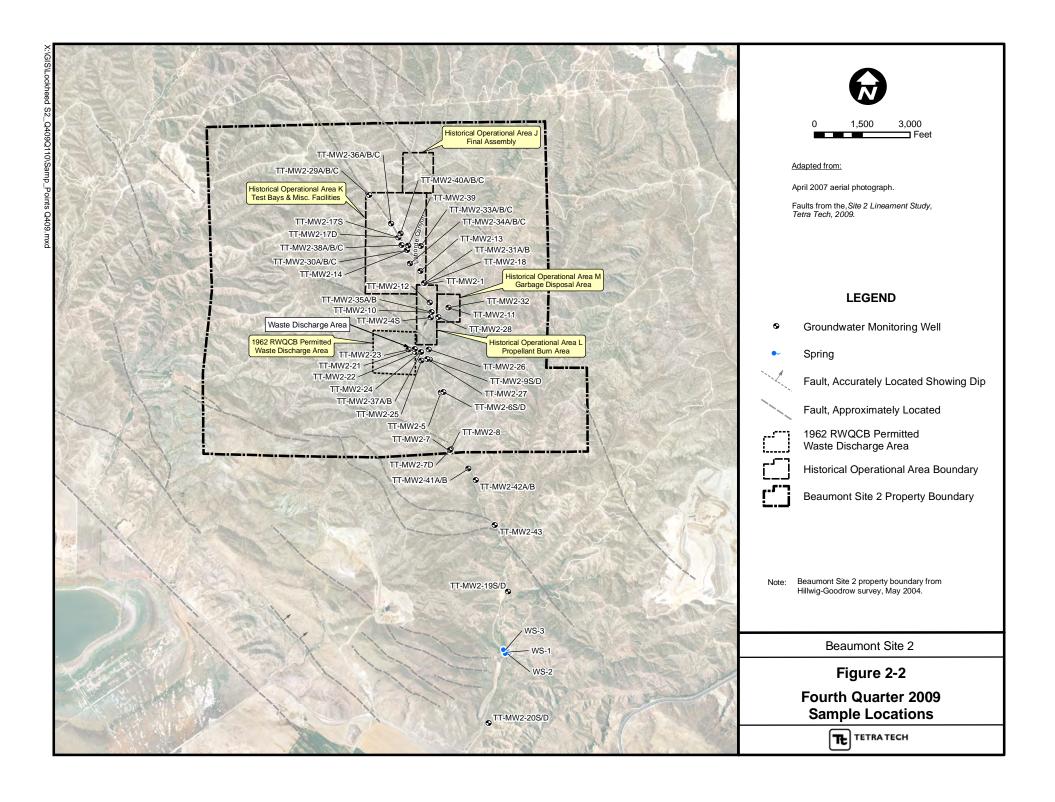
NDMA - N-Nitrosodimethylamine

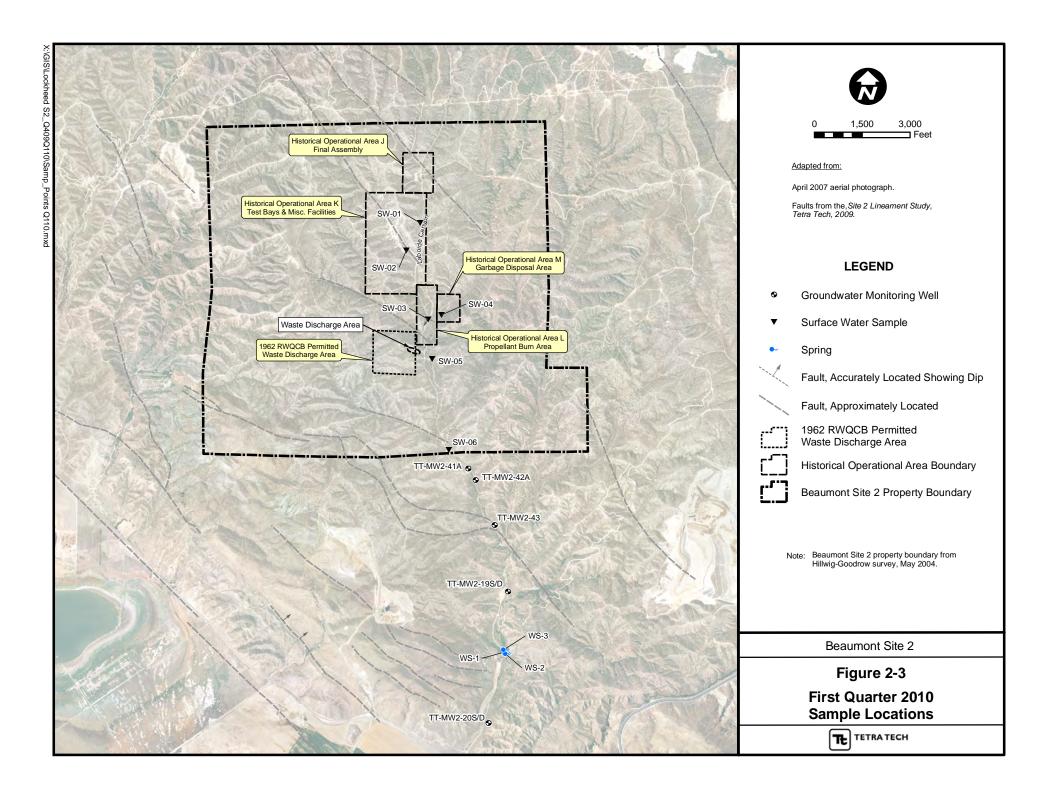
MS / MSD- Matrix Spike / Matrix Spike Duplicate.

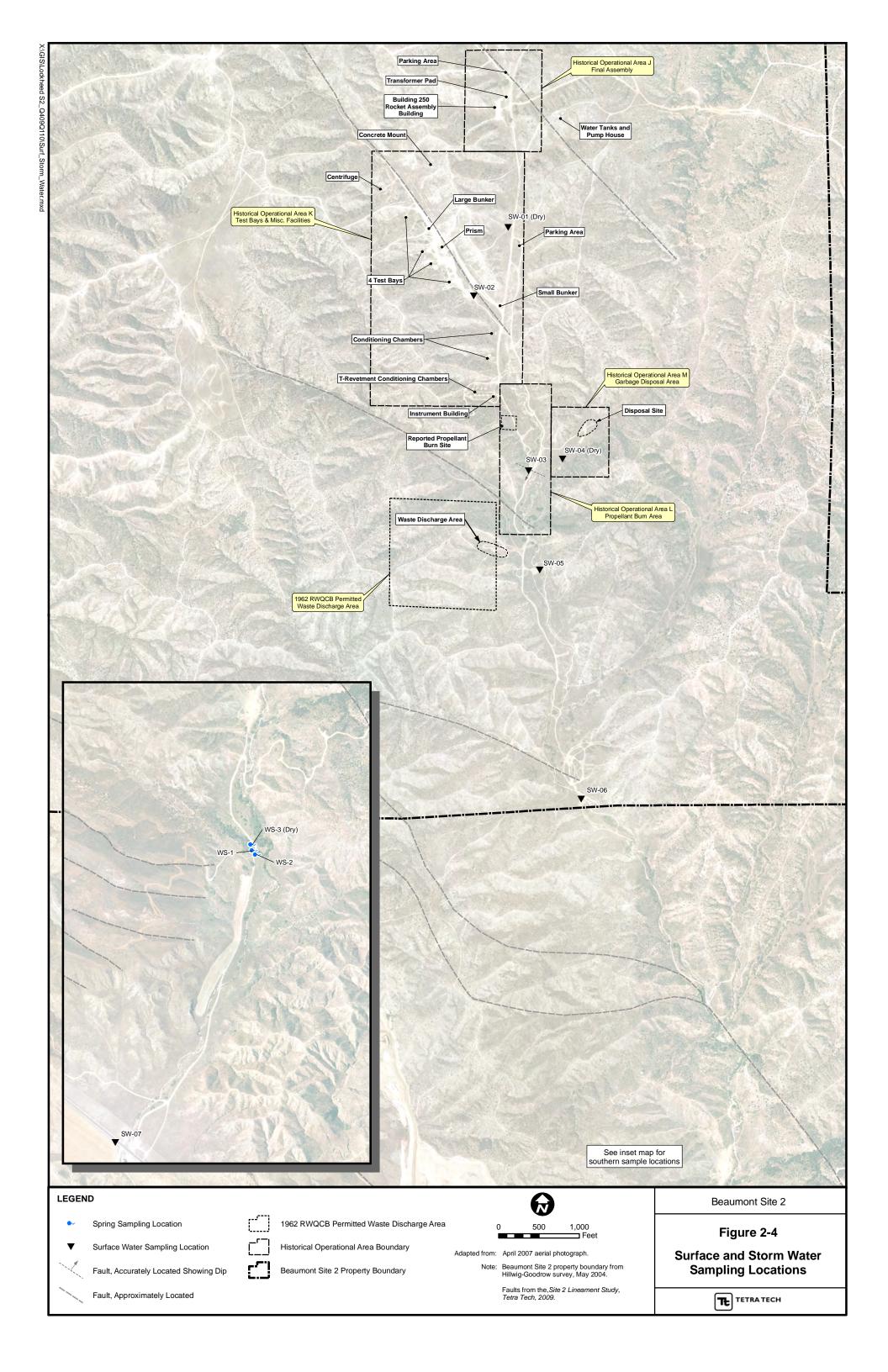
"-" Not analyzed

Table 2-2 Sampling Schedule and Analysis Method - First Quarter 2010

Monitoring Well Location	Sample Date	VOCs (EPA 8260B)	Perchlorate (EPA 331.0)	Ca, Mg, K, Na (SW6010B)	Total Dissolved Solids (E 160.1)	Chloride, Nitrate, Sulfate (E 300.0)	Carbonate, Bicarbonate (E310.1)	Comments and QA /QC Samples
WS-1	2/16/2010	-	X	X	X	X	X	Spring Sample, MSMSD, Duplicate WS-1-Dup
WS-2	2/16/2010	-	X	X	X	X	X	Spring Sample
WS-3	NA	-	-	-	-	-	-	Mud
SW-01	NA	-	-	-	-	-	-	Dry
SW-02	2/6/2010	X	X	-	-	-	-	Stom Water Sample
SW-03	2/6/2010	X	X	-	-	-	-	Stom Water Sample
SW-04	NA	-	-	-	-	-	-	Dry
SW-05	02/06/10	X	X	-	-	-	-	Stom Water Sample
SW-06	02/06/10	X	X	-	-	-	-	Stom Water Sample
SW-07	01/27/10	X	X	-	-	-	-	Stom Water Sample
TT-MW2-19S	02/16/10	-	X	-	-	-	-	Sample with Portable Bladder Pump, Duplicate TT-MW2-19S-Dup
TT-MW2-19D	02/16/10	-	X	-	-	-	-	Sample with Dedicated Pump
TT-MW2-20S	02/16/10	-	X	-	-	-	-	Sample with Dedicated Pump
TT-MW2-20D	02/16/10	-	X	-	-	-	-	Sample with Dedicated Pump
TT-MW2-41A	02/17/10	-	X	-	-	-	-	Sample with Dedicated Pump
TT-MW2-42A	02/17/10	-	X	-	-	-	-	Sample with Portable Bladder Pump
TT-MW2-43	NA	-	-	-	-	-	-	Dry
Firs	st Quarter 201	10: Total Sa	mple Locations:					17
		Total San	nples Collected:					13
Notes: EPA -	United States	s Environmer	ntal Protection Ag	ency.		Ca -	Calcium	
QA/QC -	Quality assurance / quality control Mg - Magnesium							
MS / MSD-	Matrix Spike / Matrix Spike Duplicate. K - Potassium							
"_"	Not analyzed	i				Na -	Sodium	
VOCs -	Volatile Org	anic Compou	nds					
NA -	Not available	e.						







2.4 ANALYTICAL DATA QA/QC

The groundwater samples collected were analyzed using approved EPA methods. Since the analytical data were obtained by following EPA-approved method criteria, the data were validated using the EPA-approved evaluation methods described in the National Functional Guidelines (EPA, 2004 and EPA, 2008).

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 U.S. Fish and Wildlife Service approved Habitat Conservation Plan (HCP) [USFWS, 2005] and subsequent clarifications (LMC, 2006a and 2006b) of the HCP. Groundwater sampling activities were conducted with light duty vehicles, and as specified in the Low Affect HCP, do not require biological monitoring.

SECTION 3 GROUNDWATER MONITORING RESULTS

The results of the Fourth Quarter 2009 and First Quarter 2010 groundwater monitoring events are presented in the following subsections. These subsections include tabulated summaries of the groundwater elevation and water quality data, groundwater elevation maps, and analyte results figures.

3.1 GROUNDWATER ELEVATION

Based on the groundwater levels measured during the Fourth Quarter 2009 and First Quarter 2010 monitoring events, groundwater elevations at the Site range from about 2,075 feet msl at TT-MW2-16, located in the northern portion of the Site, to about 1,819 feet msl at TT-MW2-8, located in the southern portion of the Site. Depth to groundwater ranged from about 118 feet bgs at TT-MW2-29B to about 15 feet bgs at TT-MW2-8. A tabulated summary of groundwater depths and elevations is presented in Table 3-1. Changes in groundwater elevations from the previous monitoring event for wells monitored for the Fourth Quarter 2009 and First Quarter 2010 monitoring events are shown on Figures 3-1 and 3-2, respectively, and hydrographs for individual wells are presented in Appendix D.

When compared to Third Quarter 2009, groundwater elevation during Fourth Quarter 2009 decreased an average of 0.16 feet. When compared to Fourth Quarter 2009, groundwater elevation during First Quarter 2010 increased an average of 0.30 feet. Changes in groundwater elevations by area for wells monitored for the Fourth Quarter 2009 and First Quarter 2010 monitoring events are shown in Table 3-2.

 Table 3-1 Groundwater Elevation Data - Fourth Quarter 2009 and First Quarter 2010

			Four	th Quarter 2009			Firs	st Quarter 2010	
	Measuring Point Elevation (feet	Date	Depth to Water (from Measuring	Groundwater Elevation	Groundwater Elevation Change from Third Quarter	Date	Depth to Water (from Measuring	Groundwater Elevation	Groundwater Elevation Change from Fourth Quarter
Well ID	msl)	Measured	Point, feet)	(feet msl)	2009 (feet)	Measured	Point, feet)	(feet msl)	2009 (feet)
TT-EW2-1	1840.24	12/01/09	22.27	1817.97	NA	02/15/10	20.90	1819.34	1.37
TT-MW2-1	2035.21	12/01/09	57.97	1977.24	-0.33	02/15/10	58.21	1977.00	-0.24
TT-MW2-2	2137.75	12/01/09	70.25	2067.50	-0.04	02/15/10	70.29	2067.46	-0.04
TT-MW2-3	2094.66	12/01/09	70.55	2024.11	-0.04	02/15/10	70.65	2024.01	-0.10
TT-MW2-4S	1986.94	12/01/09	50.86	1936.08	-0.12	02/15/10	50.80	1936.14	0.06
TT-MW2-4D	1987.17	12/01/09	58.05	1929.12	-0.17	02/15/10	57.80	1929.37	0.25
TT-MW2-5	1911.31	12/01/09	40.16	1871.15	-0.13	02/15/10	39.65	1871.66	0.51
TT-MW2-6S	1908.00	12/01/09	36.71	1871.29	-0.16	02/15/10	35.63	1872.37	1.08
TT-MW2-6D	1908.07	12/01/09	37.67	1870.40	-0.16	02/15/10	36.78	1871.29	0.89
TT-MW2-7	1839.25	12/01/09	21.52	1817.73	-0.80	02/15/10	20.22	1819.03	1.30
TT-MW2-7D	1838.96	12/01/09	19.00	1819.96	-0.74	02/15/10	17.95	1821.01	1.05
TT-MW2-8	1836.32	12/01/09	18.21	1818.11	-0.05	02/15/10	16.90	1819.42	1.31
TT-MW2-9S	1938.38	12/01/09	39.35	1899.03	-0.85	02/15/10	39.30	1899.08	0.05
TT-MW2-9D	1938.78	12/01/09	43.24	1895.54	-0.34	02/15/10	42.91	1895.87	0.33
TT-MW2-10	2001.57	12/01/09	57.83	1943.74	-0.26	02/15/10	57.78	1943.79	0.05
TT-MW2-11	2004.51	12/01/09	49.88	1954.63	-0.13	02/15/10	49.84	1954.67	0.04
TT-MW2-12	2016.26	12/01/09	50.85	1965.41	-0.10	02/15/10	50.91	1965.35	-0.06
TT-MW2-13	2049.39	12/01/09	66.59	1982.80	-0.10	02/15/10	66.69	1982.70	-0.10
TT-MW2-14	2074.78	12/01/09	66.31	2008.47	-0.24	02/15/10	66.55	2008.23	-0.24
TT-MW2-16	2137.20	12/01/09	62.20	2075.00	-0.40	02/15/10	62.34	2074.86	-0.14
TT-MW2-17S	2095.55	12/01/09	71.25	2024.30	0.06	02/15/10	71.39	2024.16	-0.14
TT-MW2-17D	2095.33	12/01/09	71.45	2023.88	-0.04	02/15/10	71.38	2023.95	0.07
TT-MW2-18	2035.32	12/01/09	57.88	1977.44	-0.31	02/15/10	58.11	1977.21	-0.23
TT-MW2-19S	1698.34	12/01/09	45.42	1652.92	0.43	02/15/10	45.58	1652.76	-0.16
TT-MW2-19D	1698.37	12/01/09	26.36	1672.01	-1.45	02/15/10	24.60	1673.77	1.76
TT-MW2-20S	1587.77	12/01/09	34.54	1553.23	-0.44	02/15/10	34.88	1552.89	-0.34
TT-MW2-20D	1587.48	12/01/09	33.79	1553.69	-0.48	02/15/10	34.09	1553.39	-0.30
TT-MW2-21	1978.45	12/01/09	66.32	1912.13	0.02	02/15/10	66.36	1912.09	-0.04
TT-MW2-22	1975.86	12/01/09	65.14	1910.72	0.08	02/15/10	65.19	1910.67	-0.05
TT-MW2-23	1995.17	12/01/09	82.94	1912.23	-0.09	02/15/10	82.96	1912.21	-0.02
TT-MW2-24	1964.26	12/01/09	53.78	1910.48	0.09	02/15/10	53.74	1910.52	0.04
TT-MW2-25	1966.96	12/01/09	63.96	1903.00	0.05	02/15/10	64.02	1902.94	-0.06
TT-MW2-26	1944.43	12/01/09	38.19	1906.24	-0.68	02/15/10	37.51	1906.92	0.68
TT-MW2-27	1948.27	12/01/09	50.25	1898.02	-0.84	02/15/10	50.15	1898.12	0.10
TT-MW2-28	1995.65	12/01/09	62.02	1933.63	-0.84	02/15/10	61.79	1933.86	0.23
TT-MW2-29A	2147.77	12/01/09	Dry	Dry	NA	02/15/10	Dry	Dry	NA

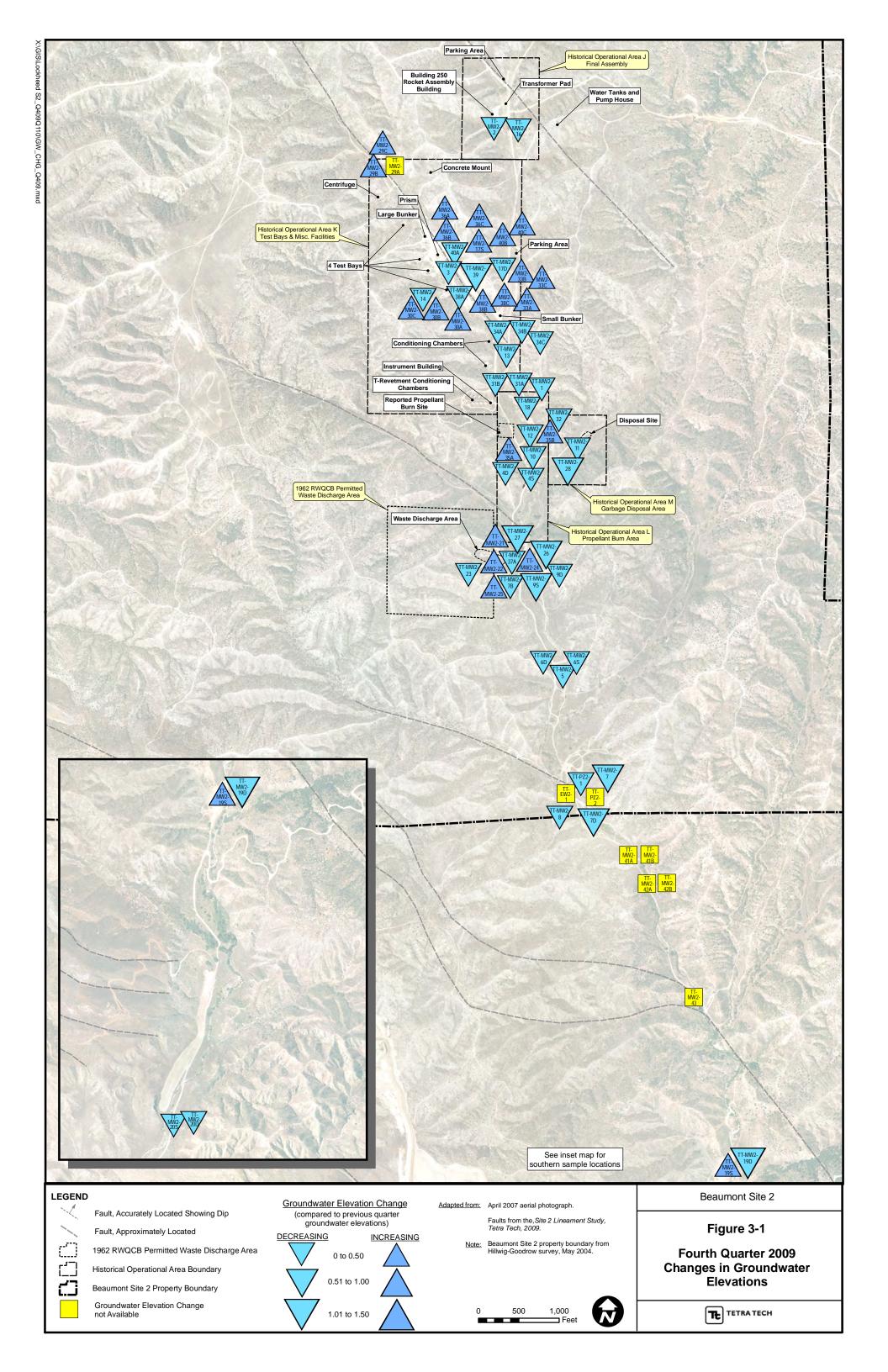
NA - Not applicable msl - Mean sea level #.## - Denotes an increase in groundwater elevation - #.## - Denotes a decrease in groundwater elevation

Table 3-1 Groundwater Elevation Data - Fourth Quarter 2009 and First Quarter 2010 (Continued)

			Four	th Quarter 2009			Firs	st Quarter 2010	
	Measuring Point Elevation (feet	Date	Depth to Water (from Measuring	Groundwater Elevation	Groundwater Elevation Change from Third Quarter	Date	Depth to Water (from Measuring	Groundwater Elevation	Groundwater Elevation Change from Fourth Quarter
Well ID	msl)	Measured	Point, feet)	(feet msl)	2009 (feet)	Measured	Point, feet)	(feet msl)	2009 (feet)
TT-MW2-29B	2147.90	12/01/09	121.60	2026.30	0.35	02/15/10	121.82	2026.08	-0.22
TT-MW2-29C	2147.83	12/01/09	127.71	2020.12	0.10	02/15/10	127.60	2020.23	0.11
TT-MW2-30A	2074.37	12/01/09	72.95	2001.42	0.13	02/15/10	72.93	2001.44	0.02
TT-MW2-30B	2074.41	12/01/09	75.43	1998.98	0.04	02/15/10	75.28	1999.13	0.15
TT-MW2-30C	2074.35	12/01/09	77.68	1996.67	0.08	02/15/10	77.63	1996.72	0.05
TT-MW2-31A	2036.11	12/01/09	58.86	1977.25	-0.22	02/15/10	59.04	1977.07	-0.18
TT-MW2-31B	2036.15	12/01/09	66.61	1969.54	-0.07	02/15/10	66.25	1969.90	0.36
TT-MW2-32	2004.87	12/01/09	53.69	1951.18	-0.12	02/15/10	53.58	1951.29	0.11
TT-MW2-33A	2070.54	12/01/09	61.22	2009.32	0.04	02/15/10	61.21	2009.33	0.01
TT-MW2-33B	2070.54	12/01/09	65.95	2004.59	0.08	02/15/10	65.95	2004.59	0.00
TT-MW2-33C	2070.54	12/01/09	64.05	2006.49	0.12	02/15/10	64.02	2006.52	0.03
TT-MW2-34A	2066.84	12/01/09	65.97	2000.87	-0.06	02/15/10	66.04	2000.80	-0.07
TT-MW2-34B	2066.85	12/01/09	73.15	1993.70	-0.05	02/15/10	73.11	1993.74	0.04
TT-MW2-34C	2066.84	12/01/09	74.77	1992.07	-0.01	02/15/10	74.70	1992.14	0.07
TT-MW2-35A	2003.20	12/01/09	49.59	1953.61	0.12	02/15/10	49.08	1954.12	0.51
TT-MW2-35B	2003.20	12/01/09	55.05	1948.15	0.02	02/15/10	54.75	1948.45	0.30
TT-MW2-36A	2100.99	12/01/09	79.01	2021.98	0.09	02/15/10	78.97	2022.02	0.04
TT-MW2-36B	2101.04	12/01/09	79.78	2021.26	0.02	02/15/10	79.75	2021.29	0.03
TT-MW2-36C	2100.88	12/01/09	79.76	2021.12	0.04	02/15/10	79.75	2021.13	0.01
TT-MW2-37A	1963.62	12/01/09	63.25	1900.37	-0.24	02/15/10	62.98	1900.64	0.27
TT-MW2-37B	1963.67	12/01/09	71.26	1892.41	-0.15	02/15/10	70.94	1892.73	0.32
TT-MW2-38A	2084.56	12/01/09	59.48	2025.08	-0.29	02/15/10	59.63	2024.93	-0.15
TT-MW2-38B	2084.42	12/01/09	81.45	2002.97	0.05	02/15/10	81.35	2003.07	0.10
TT-MW2-38C	2084.63	12/01/09	88.90	1995.73	0.05	02/15/10	88.76	1995.87	0.14
TT-MW2-39	2079.53	12/01/09	61.95	2017.58	-0.74	02/15/10	61.81	2017.72	0.14
TT-MW2-40A	2096.28	12/01/09	72.48	2023.80	-0.03	02/15/10	72.56	2023.72	-0.08
TT-MW2-40B	2096.24	12/01/09	83.82	2012.42	0.03	02/15/10	83.77	2012.47	0.05
TT-MW2-40C	2096.28	12/01/09	88.88	2007.40	0.12	02/15/10	88.71	2007.57	0.17
Tt-MW2-41A	1812.47	12/01/09	23.90	1788.57	NA	02/15/10	21.92	1790.55	1.98
Tt-MW2-41B	1812.22	12/01/09	21.06	1791.16	NA	02/15/10	18.56	1793.66	2.50
Tt-MW2-42A	1799.06	12/01/09	28.02	1771.04	NA	02/15/10	26.24	1772.82	1.78
Tt-MW2-42B	1799.07	12/01/09	25.60	1773.47	NA	02/15/10	24.32	1774.75	1.28
Tt-MW2-43	1771.44	12/01/09	Dry	Dry	NA	02/15/10	Dry	Dry	NA
TT-PZ2-1	1847.06	12/01/09	19.91	1827.15	-0.27	02/15/10	19.15	1827.91	0.76
TT-PZ2-2	1840.76	12/01/09	22.40	1818.36	NA	02/15/10	20.97	1819.79	1.43

NA - Not applicable msl - Mean sea level

#.## - Denotes an increase in groundwater elevation - #.## - Denotes a decrease in groundwater elevation



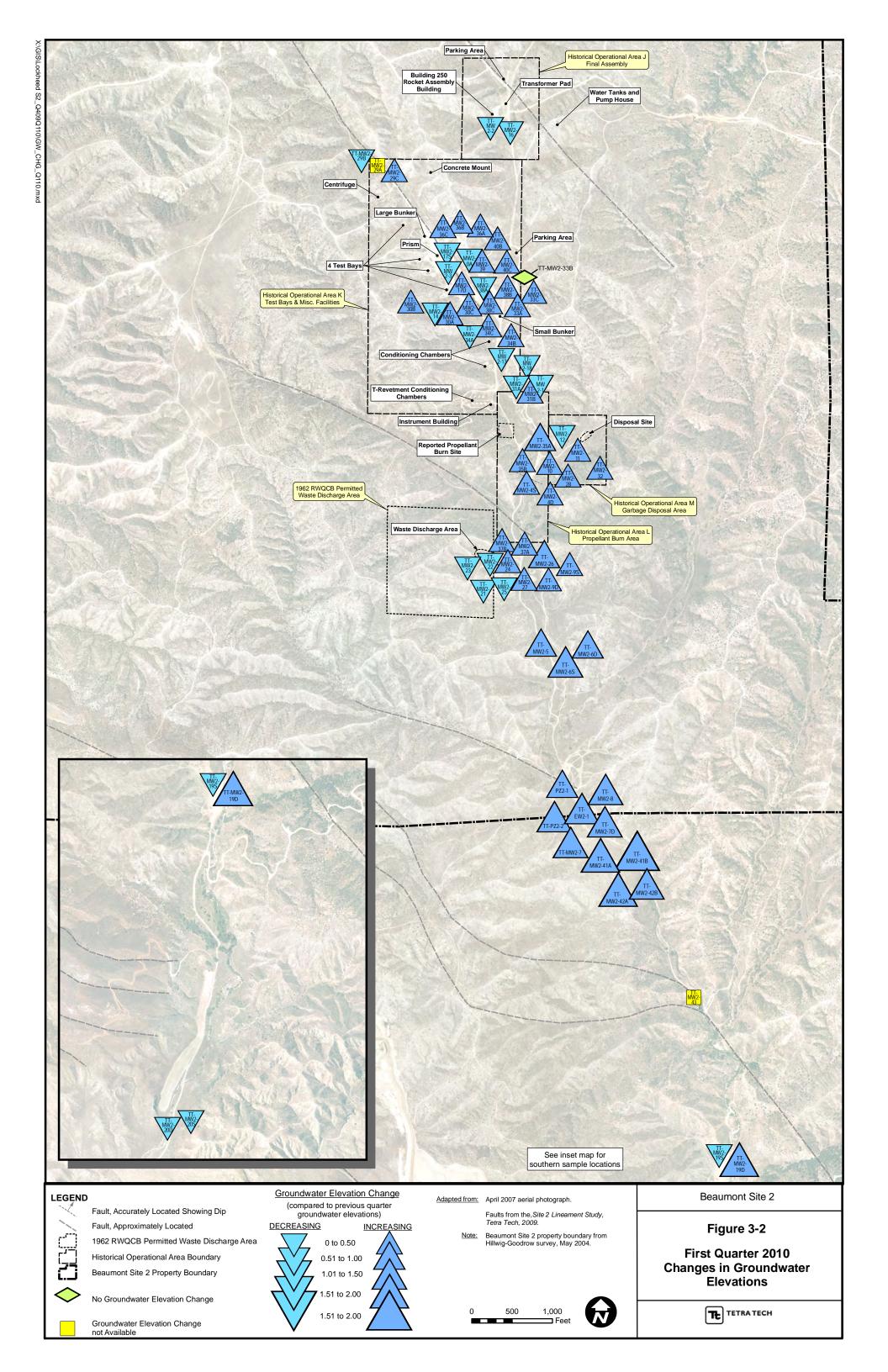


Table 3-2 Groundwater Elevation Change - Fourth Quarter 2009 and First Quarter 2010

Site Area	0	dwater Elevation h Quarter 2009	Average Change By Area	Range of Groundwate Change - First Qua		Average Change By Area
J	-0.40	-0.04	-0.22	-0.14	-0.04	-0.09
K	-0.74	0.35	-0.04	-0.24	0.36	-0.01
L	-0.26	0.12	-0.08	-0.06	0.51	0.19
M	-0.84	-0.12	-0.36	0.04	0.23	0.13
FWDA	-0.24	0.09	-0.05	-0.05	0.32	0.09
LC	-0.85	0.05	-0.41	-0.06	2.50	1.02
WS	-1.45	0.43	-0.48	-0.34	1.76	0.24

J - Final Assembly Area

K - Former Test Bay Area

L - Former Burn Area

M - Garbage disposal Area

FWDA - Former waste discharge area

LC - Lower Canyon

WS - Former Wolfskill property

3.2 GROUNDWATER FLOW

Groundwater contour maps for first groundwater screened wells from Fourth Quarter 2009 and First Quarter 2010 groundwater levels are presented in Figure 3-3 and Figures 3-4 respectively. Hydrographs for individual wells are presented in Appendix D.

3.3 GROUNDWATER GRADIENTS

The average horizontal groundwater gradients calculated between TT-MW2-16 and TT-MW2-6S from the Fourth Quarter 2009 and First Quarter 2010 groundwater monitoring events for the shallow Quaternary Alluvium and weathered San Timoteo Formation (QAL/wSTF) screened wells were 0.030 ft/ft. The horizontal groundwater gradients calculated between TT-MW2-2 and TT-MW2-6D for the Fourth Quarter 2009 and First Quarter 2010 groundwater monitoring events for the deeper San Timoteo Formation (STF) screened wells were 0.029 ft/ft.

Vertical groundwater gradients are calculated from individual clusters of wells. Well clusters are used to measure the differences in static water level at different depths within the aquifer. The vertical gradient is a comparison of static water level between wells at different depths within the aquifer and is an indication of the vertical flow, (downward - negative gradient, upward - positive gradient), of groundwater. Vertical groundwater gradients at the Site are generally downward. The vertical gradients range from -0.28 at well cluster TT-MW2-4S and 4D located in Area L to +0.19 at well cluster TT-MW2-19S and 19D located on the former Wolfskill property. A summary of calculated horizontal and vertical groundwater gradients is presented in Table 3-3. A complete listing of historical horizontal and vertical groundwater gradients and associated calculations is presented in Appendix E.

Table 3-3 Summary of Horizontal and Vertical Groundwater Gradient

Horizontal Groundwater Gradients (fee	et / foot), approximating a flow	v line perpendicular to	groundwater contours
<u> </u>	Overall	Overall	
	STF	QAL/WSTF	
<u> </u>	TT-MW2-2	TT-MW2-16	
	to	to	
	TT-MW2-6D	TT-MW2-6S	
Fourth Quarter (December) 2009	0.029	0.030	
FIrst Quarter (February) 2010	0.029	0.030	

Vertical Groundwater Gradients (feet / foot)

-					Southern portion of Site	Southern portion of Site	Southern portion of	Former Wolfskill	Former Wolfskill
_	Area J	Area K	Area K	Area L	2	2	Site 2	Property	Property
				TT- MW2-					
	TT-MW2-2	TT-MW2-17D	TT-MW2-18	4D	TT-MW2-9D	TT-MW2-6D	TT-MW2-7D	TT-MW2-19D	TT-MW2-20D
deep screen	(STF)	(QAL/WSTF)	(STF)	(STF)	(STF)	(STF)	(STF)	(MEF)	(MEF)
				TT-					
			TT-MW2-1	MW2-					
	TT-MW2-16	TT-MW2-17S	(QAL /	4S	TT-MW2-9S	TT-MW2-6S	TT-MW2-7	TT-MW2-19S	TT-MW2-20S
shallow screen	(QAL/WSTF)	(QAL/WSTF)	WSTF)	(STF)	(QAL/WSTF)	(QAL/WSTF)	(QAL/WSTF)	(QAL/MEF)	(QAL)
Fourth Quarter (December) 2009	-0.16	-0.02	0.01	-0.28	-0.13	-0.05	0.05	0.17	0.01
First Quarter (February) 2010	-0.16	-0.01	0.01	-0.27	-0.12	-0.06	0.05	0.19	0.03

Notes:

STF - San Timoteo Formation
MEF - Mt. Eden Formation
QAL - Quaternary Alluvium

QAL/WSTF - Quaternary Alluvium and weathered San Timoteo Formation

QAL/MEF - Quaternary Alluvium and Mt. Eden Formation

3.4 ANALYTICAL DATA SUMMARY

Groundwater samples collected during the Fourth Quarter 2009 monitoring event were analyzed for perchlorate. Select wells were also sampled for VOCs, NDMA, 1,4-dioxane, and natural attenuation parameters,. VOCs and perchlorate are contaminants of potential concern at the Site. During the First Quarter 2010 monitoring event, groundwater samples were analyzed for perchlorate, spring samples were analyzed for perchlorate and general minerals parameters, and storm water samples were tested for perchlorate and VOCs.

Summaries of validated laboratory analytical results for analytes detected above their respective MDLs during the Fourth Quarter 2009 monitoring event are presented in Table 3-4. Summaries of validated laboratory analytical results for analytes detected above their respective MDLs during the First Quarter 2010 monitoring event are presented in Tables 3-5. A complete list of the analytes tested along with validated sample results by analytical method are provided in Appendix F. VOC and perchlorate sample results above the published MCL (federal or state, whichever is lower) or DWNL are bolded in Tables 3-4 and 3-5. Tables 3-6 and 3-7 present summary statistics for validated organic and inorganic analytes detected during the monitoring events. Laboratory analytical data packages, which include all environmental, field QC, and laboratory QC results, are provided in Appendix G. A consolidated laboratory data summary table is presented in Appendix H.

3.4.1 Data Quality Review

The quality control samples were reviewed as described in the Revised Groundwater Sampling and Analysis Plan (Tetra Tech, 2003b). The data for the groundwater sampling activities were contained in analytical data packages generated by E.S. Babcock & Sons, Inc. These data packages were reviewed using the latest versions of the National Functional Guidelines for Organic and Inorganic Data Review documents from the EPA (EPA, 2008 and 2005).

Preservation criteria, holding times, field blanks, laboratory control samples (LCS), 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 include comparing statistically calculated control limits to percent recoveries of all spiked analytes and duplicate spiked analytes. Relative Percent Difference (RPD) control limits are compared to actual spiked (MS/MSD) 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 A209B for total dissolved solids, Method AM23G for volatile fatty acids, Method AM20GAX for hydrogen, Method E300.0 for nitrate, sulfate, and chloride, Method E521 for low level NDMA, Method E332.0 for perchlorate, Method A5310 for total and dissolved organic carbon, Method RSK-175 for methane, ethane, ethene, Method SW8270C SIM for 1,4-dioxane, Method SW6010B for metals, and Method 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.

Method E521 for low level NDMA had surrogate recovery errors and field duplicate RPD errors that qualified as estimated in 33.4 percent of the total E521 data. The data qualified as estimated is usable for the intended purpose. Field blank contamination caused 13.3 percent of the total E521 data to be qualified for blank contamination. Data qualified for blank contamination is generally not usable. Additional sample volumes will be collected for future E521 analyses so the laboratory can perform corrective action to mitigate surrogate recovery errors.

Method SW8260B for VOCs had a surrogate recovery error that qualified as estimated in 0.05 percent of the total SW8260B data. Trip blank contamination caused 0.5 percent of the total SW8260B data to be qualified for blank contamination. The data qualified as estimated is usable for the intended purpose. Blank qualified data is generally not usable.

Method RSK-175 for methane, ethane, and ethene had holding time errors that qualified as estimated 7.7 percent of the total RSK-175 data. The data qualified as estimated is usable for the intended purpose. Method blank contamination caused 12.8 of the total RSK-175 data to be qualified for blank contamination. Additional sample volumes will be collected for future RSK-175 analyses so the laboratory can perform corrective action to mitigate method blank contamination.

Method E300.0 for anions had errors that qualified as estimated in the data listed below

• Two (66%) chloride samples were qualified for LCS recovery errors and two (66%) were qualified for matrix spike recovery errors.

- Five out of 16 nitrate samples (31.3%) had holding time errors. The holding time errors were due to instrument failure. The instrument has since been repaired.
- Two out of 16 sulfate samples (12.5%) were qualified for LCS recovery.

The E300 data qualified as estimated are usable for the intended purpose. Additional sample volumes will be collected for future E300 analyses so the laboratory can perform corrective action to mitigate laboratory LCS recovery errors.

Method AM23G for volatile fatty acids had field duplicate errors that qualified as estimated 1.7 percent of the total AM23G data. The data qualified as estimated is usable for the intended purpose. Method blank contamination qualified 14.5% of the total AM23G data. Data qualified for blank contamination is generally considered not usable. Additional sample volumes will be collected for future AMG23G analyses so the laboratory can perform corrective action to mitigate method blank contamination.

Method A5310 for dissolved organic carbon had field duplicate errors that qualified 7.7 percent of the total A5310 data. The estimated data is usable for the intended purpose.

Table 3-4 Summary of Validated Detected Organic and Inorganic Analytes - Fourth Quarter 2009

Sample Location	Sample Date	Per chlorate	1,4- Dioxane	NDMA	Acetone	2- Butanone	Benzene	Carbon Disulfide	Chloro form	Chloro methane	1,1- Dichloro ethane	1,2- Dichloro ethane	1,1- Dichloro ethene	c-1,2- Dichloro ethene	t-1,2- Dichloro ethene	Ethyl benzene	2- Hexanone	4-Methyl- 2- Pentanone	Methylene Chloride	Toluene	1,1,2- Trichloro ethane	Trichloro ethene	Tetrachloro ethene	m,p- Xylenes	o- Xylene
*****				<u> </u>	<u> </u>	1	1						g/L unless otl			1			1		1	1	I		
WS-1	12/2/2009	< 0.35	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
WS-2	12/2/2009	<0.35	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
WS-3	12/2/2009	<0.35	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-1 TT-MW2-4S	12/9/2009	7,700	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-4S	12/4/2009	0.54	NA 1.0	NA NA	NA NA	NA NA	NA	NA	NA NA	NA NA	NA	NA	NA	NA	NA NA	NA	NA NA	NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA	NA NA
TT-MW2-6S	12/3/2009	930	1.0	NA NA	NA NA	NA NA	NA NA	NA	NA	NA	NA	NA	NA	NA	NA NA	NA	NA NA	NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA	NA
TT-MW2-6D	12/3/2009	1,400	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA
TT-MW2-7	12/3/2009 12/3/2009	<0.71 410	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA	NA NA
TT-MW2-7D	12/3/2009	0.91	NA NA	NA NA	<5.0	<1.2	<0.14	2.1	<0.17	<0.36	<0.098	<0.21	<0.12	<0.18	<0.10	<0.26	<1.2	<0.95	<0.15	<0.22	<0.31	<0.17	<0.17	<0.36	<0.41
TT-MW2-8	12/3/2009	350	NA NA	NA NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-9S	12/8/2009	4,700	4.3	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-9D	12/4/2009	0.22	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-10	12/9/2009	< 0.35	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-11	12/4/2009	280	NA	NA	<5.0	<1.2	< 0.14	< 0.36	< 0.17	< 0.36	< 0.098	< 0.21	<0.12	0.39 Jq	< 0.10	< 0.26	<1.2	< 0.95	0.20 Jq	<0.22	<0.31	9.2	< 0.17	< 0.36	<0.41
TT-MW2-12	12/14/2009	< 0.71	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-13	12/14/2009	2,700	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-14	12/14/2009	47,000	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-17S	12/17/2009	1,500	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-17D	12/10/2009	53,000	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-18	12/14/2009	14,000	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-19S	12/11/2009	5.7	< 0.10	0.0026	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-19D	12/2/2009	< 0.071	NA	0.0044	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-20S	12/2/2009	< 0.35	NA	0.0012 Jq	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-20D	12/2/2009	< 0.35	NA	0.0100	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-21	12/4/2009	2.1	< 0.10	0.0039 Jb	< 5.0	<1.2	< 0.14	< 0.36	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	4.9	< 0.22	< 0.31	2.8	< 0.17	< 0.36	< 0.41
TT-MW2-22	12/4/2009	< 0.071	42	NA	< 5.0	<1.2	0.87	0.37 Jq	0.29 Jq	< 0.36	3.1	1.7	29	1.4	0.35 Jq	< 0.26	<1.2	< 0.95	9.3	< 0.22	< 0.31	460	< 0.17	< 0.36	< 0.41
TT-MW2-23	12/11/2009	< 0.071	NA	NA	< 5.0	<1.2	< 0.14	< 0.36	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	< 0.15	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-24	12/11/2009	140,000	320	0.0040 Jf	< 5.0	<1.2	0.21 Jq	< 0.36	3.8	< 0.36	0.89	0.78	2.7	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	0.78 BJkq	< 0.22	0.60	100	< 0.17	< 0.36	< 0.41
TT-MW2-25	12/4/2009	< 0.071	NA	0.0031	<5.0	<1.2	< 0.14	< 0.36	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	< 0.15	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-26	12/4/2009	90	< 0.10	0.0032	<5.0	<1.2	<0.14	<0.36	<0.17	<0.36	<0.098	<0.21	<0.12	<0.18	<0.10	<0.26	<1.2	<0.95	<0.15	<0.22	<0.31	<0.17	<0.17	<0.36	<0.41
TT-MW2-27	12/3/2009	200	NA	NA	<5.0	<1.2	< 0.14	1.2	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	< 0.15	<0.22	< 0.31	< 0.17	< 0.17	< 0.36	<0.41
TT-MW2-28	12/4/2009	29	NA	0.0100	<5.0	<1.2	< 0.14	< 0.36	< 0.17	< 0.36	< 0.098	< 0.21	<0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	0.15 Jq	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
Method De	tection Limit	0.071	0.10	0.0007	5	1.2	0.14	0.36	0.17	0.36	0.098	0.21	0.12	0.18	0.10	0.26	1.2	0.95	0.15	0.22	0.31	0.17	0.17	0.36	0.41
MCL (unless no	ted) / DWNL	6	3 (1)	0.01(1)	-	-	1	160(1)	-	-	5	0.5	6	6	10	300	-	-	5	150	5	5	5	1750	1750

Notes: Only analytes positively detected in samples are presented in this table. For a complete list of constituents analyzed, refer to the laboratory data package.

μg/L - Micrograms per liter

NDMA - N-Nitrosodimethylamine

MCL - California Department of Health Services Maximum Contaminant Level.

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

(1) - DWNL

" - " MCL/DWNL not established.

< # - Method detection limit concentration is shown.

Bold - Maximum Contaminant Level exceeded.

- VA Not analyze
- B The sample result was less than 5 times blank contamination. Cross contamination is suspected.
- J The analyte was positively identified, but the concentration is an estimated value.
- b The surrogate spike recovery was outside control limits.
- f The duplicate Relative Percent Difference (RPD) was outside the control limit
- k The analyte was found in a field blank.
- q $\;\;$ The analyte detected was below the Practical Quantitation Limit (PQL).

Table 3 4 Summary of Validated Detected Organic and Inorganic Analytes - Fourth Quarter 2009 (continued)

Sample	Sample	Per	1,4-	NDMA	Acetone	2-	Benzene	Carbon	Chloro	Chloro	1,1-	1,2-	1,1-	c-1,2-	t-1,2-	Ethyl	2-	4-Methyl-	Methylene	Toluene	1,1,2-	Trichloro	Tetrachloro	m,p-	0-
Location	Date	chlorate	Dioxane			Butanone		Disulfide	form	methane	Dichloro	Dichloro	Dichloro	Dichloro	Dichloro	benzene	Hexanone	2-	Chloride		Trichloro	ethene	ethene	Xylenes	Xylene
											ethane	ethane	ethene	ethene	ethene			Pentanone			ethane			<u>'</u>	
				•		•					All results re	eported in µg	z/L unless oth	erwise state	d		•				•	•			
TT-MW2-29B	12/7/2009	0.25	NA	0.0110	< 5.0	<1.2	< 0.14	1.3	< 0.17	< 0.36	< 0.098	<0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	< 0.15	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-29C	12/7/2009	0.22	NA	NA	< 5.0	<1.2	< 0.14	0.96	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	< 0.15	0.26 Jq	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-30A	12/14/2009	5,300	NA	NA	< 5.0	<1.2	< 0.14	< 0.36	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	< 0.15	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-30B	12/11/2009	1,100	NA	NA	< 5.0	<1.2	< 0.14	0.70	0.20 Jq	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	< 0.15	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-30C	12/11/2009	< 0.071	NA	NA	15	24	< 0.14	1.7	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	0.20 BJkq	1.8	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-31A	12/9/2009	< 0.35	NA	NA	<5.0	<1.2	< 0.14	1.0	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	< 0.15	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-31B	12/9/2009	< 0.35	NA	NA	< 5.0	<1.2	0.25 Jq	2.0	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	< 0.15	1.4	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-32	12/9/2009	< 0.071	NA	NA	< 5.0	<1.2	< 0.14	0.98	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	< 0.15	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-33A	12/11/2009	< 0.071	NA	NA	< 5.0	<1.2	< 0.14	< 0.36	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	< 0.15	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-33B	12/15/2009	< 0.071	NA	NA	<5.0	<1.2	< 0.14	1.4	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	0.17 Jq	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-33C	12/15/2009	0.37	NA	NA	< 5.0	<1.2	0.22 Jq	0.93	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	< 0.15	7.4	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-34A	12/15/2009	0.39	NA	NA	< 5.0	<1.2	< 0.14	2.1	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	0.19 Jq	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-34B	12/15/2009	0.43	NA	NA	<5.0	<1.2	< 0.14	0.67	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	0.16 Jq	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-34C	12/15/2009	< 0.071	NA	NA	< 5.0	<1.2	< 0.14	1.4	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	0.16 Jq	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-35A	12/9/2009	< 0.071	NA	NA	< 5.0	<1.2	< 0.14	1.3	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	< 0.15	0.49 Jq	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-35B	12/9/2009	< 0.071	NA	NA	< 5.0	<1.2	0.24 Jq	0.93	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	0.15 BJkq	3.0	< 0.31	< 0.17	0.22 Jq	< 0.36	< 0.41
TT-MW2-36A	12/10/2009	< 0.071	NA	0.0067 Jf	< 5.0	<1.2	< 0.14	0.71	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	0.23 BJkq	0.60	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-36B	12/10/2009	< 0.071	NA	NA	< 5.0	<1.2	< 0.14	< 0.36	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	0.22 BJkq	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-36C	12/10/2009	< 0.071	NA	NA	37	4.8	0.45 Jq	57	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	0.25 BJkq	1.0	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-37A	12/8/2009	4100	7.5	NA	100	16	0.52	< 0.36	0.19 Jq	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	2.5 Jq	1.4 Jq	< 0.15	0.53	< 0.31	0.86	< 0.17	< 0.36	< 0.41
TT-MW2-37B	12/8/2009	0.49	NA	NA	< 5.0	<1.2	< 0.14	13	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	< 0.15	0.66	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-38A	12/14/2009	170,000	NA	NA	< 5.0	<1.2	< 0.14	< 0.36	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	< 0.15	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-38B	12/17/2009	15,000	NA	NA	< 5.0	1.4 Jq	< 0.14	10	< 0.17	0.73	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	< 0.15	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-38C	12/14/2009	0.092 Jq	NA	NA	< 5.0	<1.2	< 0.14	0.59	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	< 0.15	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-39	12/15/2009	81,000	NA	NA	< 5.0	<1.2	< 0.14	< 0.36	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	0.15 Jq	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-40A	12/10/2009	< 0.071	NA	NA	< 5.0	<1.2	< 0.14	< 0.36	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	< 0.15	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-40B	12/10/2009	4	NA	NA	< 5.0	<1.2	21	< 0.36	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	1.6	<1.2	< 0.95	0.21 BJkq	13	< 0.31	< 0.17	< 0.17	2.4	1.6
TT-MW2-40C	12/10/2009	< 0.071	NA	NA	< 5.0	1.9 Jq	0.57	2.0	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	0.25 BJkq	0.51	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-41A	12/16/2009	< 0.071	< 0.10	0.0042 Bk	< 5.0	<1.2	< 0.14	< 0.36	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	< 0.15	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-41B	12/16/2009	0.11	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-42A	12/16/2009	< 0.071	< 0.10	0.0020 Bk	< 5.0	<1.2	< 0.14	< 0.36	< 0.17	< 0.36	< 0.098	< 0.21	< 0.12	< 0.18	< 0.10	< 0.26	<1.2	< 0.95	< 0.15	< 0.22	< 0.31	< 0.17	< 0.17	< 0.36	< 0.41
TT-MW2-42B	12/16/2009	0.099 Jq	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
Method De	tection Limit	0.071	0.10	0.0007	9.1	6.9	0.28	1.9	0.33	0.36	0.37	0.31	0.40	0.49	0.10	0.22	1.2	1.0	2.6	0.33	0.54	0.30	0.17	0.45	0.24
MCL (unless not		6	3 (1)	0.01(1)	-	-	1	160 (1)	-	-	5	0.5	6	6	10	300	-	-	5	150	5	5	5	1750	1750
(,. =	-	- (-)		1	l	_	(-)									l		-				_		

Notes: Only analytes positively detected in samples are presented in this table. For a complete list of constituents analyzed, refer to the laboratory data package.

μg/L - Micrograms per liter

NDMA - N-Nitrosodimethylamine

MCL - California Department of Health Services Maximum Contaminant Level.

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

" - " MCL/DWNL not established.

< # - Method detection limit concentration is shown.

Bold - Maximum Contaminant Level exceeded

B - The sample result was less than 5 times blank contamination. Cross contamination is suspected.

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

b - The surrogate spike recovery was outside control limits.

f - The duplicate Relative Percent Difference (RPD) was outside the control limit

The analyte was found in a field blank.

q - The analyte detected was below the Practical Quantitation Limit (PQL).

Table 3-5 Summary of Validated Detected Organic and Inorganic Analytes - First Quarter 2010

Sample Name	Sample Date	Methylene Chloride - ug/L	Perchlorate - ug/L	Calcium - mg/L	Magnesium -mg/L	Potassium -mg/L	Sodium - mg/L	Alkalinity, Total (as CaCO3) - mg/L	Bicarbonate (as CaCO3) -mg/L	Total Dissolved Solids - mg/L	Chloride - mg/L	Sulfate - mg/L
WS-1	2/16/2010	NA	< 0.071	38	6.9	4.8	100	100	120	410	45 Jcf	160
WS-2	2/16/2010	NA	< 0.071	42	8.2	6.4	110	110	130	420	46 Jd	160
SW-02	2/6/2010	< 0.15	11	NA	NA	NA	NA	NA	NA	NA	NA	NA
SW-03	2/6/2010	< 0.15	0.59	NA	NA	NA	NA	NA	NA	NA	NA	NA
SW-05	2/6/2010	< 0.15	0.66	NA	NA	NA	NA	NA	NA	NA	NA	NA
SW-06	2/6/2010	< 0.15	1.1	NA	NA	NA	NA	NA	NA	NA	NA	NA
SW-07	1/27/2010	0.20 Jbq	0.91	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-19S	2/16/2010	NA	5.4	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-19D	2/16/2010	NA	< 0.071	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-20S	2/16/2010	NA	< 0.071	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-20D	2/16/2010	NA	< 0.071	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-41A	2/17/2010	NA	0.18	NA	NA	NA	NA	NA	NA	NA	NA	NA
TT-MW2-42A	2/17/2010	NA	0.17	NA	NA	NA	NA	NA	NA	NA	NA	NA
Method De	tection Limit	0.15	0.071	0.50	0.50	0.50	0.50	1.7	1.7	11	0.50	0.37
MCL (unless not	ed) / DWNL	5	6	-	-	-	-	-	-	500	250	250

Notes:	Only analytes positively detected in samples are presented in this table. For a complex	ete fist of	constituents analyzed, refer to the laboratory data package.
μg/L -	Micrograms per liter	<#-	Method detection limit concentration is shown.
mg/L -	Milligrams per liter	J -	The analyte was positively identified, but the concentration is an estimated value.
MCL -	California Department of Health Services Maximum Contaminant Level.	b -	the surrogate spike recovery was outside control limits.
DWNL -	California Department of Health Services drinking water notification level.	c -	The Matrix Spike (MS) and/or Matrix Spike Duplicate (MSD) recoveries were
(1) -	DWNL		outside control limits.
" - "	MCL/DWNL not established.	d -	The Laboratory Control Sample (LCS) recovery was outside control limits.
NA -	Not analyzed	f -	The duplicate Relative Percent Difference (RPD) was outside the control limit
Bold -	Maximum Contaminant Level exceeded.	a -	The analyte detection was below the Practical Quantitation Limit (PQL).

Table 3-6 Summary Statistics of Validated Detected Organic and Inorganic Analytes - Fourth Quarter 2009

Compounds Detected	Total Number of Samples Analyzed (1)	Total Number of Detections (1)	Number of Detections Exceeding MCL or DWNL (1)	Correspo MCL (u	ınless	Minin Concent Detec	ration	Maxin Concent Detec	ration
Organic Analytes:									
1,4-Dioxane	10	5	4	3 (2)	μg/L	1.0	μg/L	320	μg/L
NDMA	13	11	3	0.01(2)	μg/L	0.0012	μg/L	0.011	μg/L
Acetone	40	3	0	-	μg/L	15	μg/L	100	μg/L
2-Butanone	40	5	0	-	μg/L	1.4	μg/L	24	μg/L
Benzene	40	9	1	1	μg/L	0.21	μg/L	21	μg/L
Carbon Disulfide	40	23	0	160 (2)	μg/L	0.37	μg/L	57	μg/L
Chloroform	40	4	0	-	μg/L	0.19	μg/L	3.8	μg/L
Chloromethane	40	1	0	300	μg/L	0.73	μg/L	0.73	μg/L
1, 1-Dichloroethane	40	2	0	5	μg/L	0.89	μg/L	3.1	μg/L
1, 2-Dichloroethane	40	2	2	0.5	μg/L	0.78	μg/L	1.7	μg/L
1, 1-Dichloroethene	40	2	1	6	μg/L	2.7	μg/L	29	μg/L
cis-1, 2-Dichloroethene	40	2	0	6	μg/L	0.39	μg/L	1.4	μg/L
trans-1,2-Dichloroethene	40	1	0	10	μg/L	0.35	μg/L	0.35	μg/L
Ethylbenzene	40	1	0	300	μg/L	1.6	μg/L	1.6	μg/L
2-Hexanone	40	1	0	-	μg/L	2.5	μg/L	2.5	μg/L
4-Methyl-2-Pentanone	40	1	0	-	μg/L	1.4	μg/L	1.4	μg/L
Methylene Chloride	40	9	1	5	μg/L	0.15	μg/L	9.3	μg/L
Toluene	40	12	0	150	μg/L	0.26	μg/L	13	μg/L
1, 1, 2-Trichloroethane	40	1	0	5	μg/L	0.6	μg/L	0.6	μg/L
Trichloroethene	40	5	3	5	μg/L	0.86	μg/L	460	μg/L
Tetrachloroethene	40	1	0	5	μg/L	0.22	μg/L	0.22	μg/L
m,p-Xylenes	40	1	0	1750	μg/L	2.4	μg/L	2.4	μg/L
o-Xylene	40	1	0	1750	μg/L	1.6	μg/L	1.6	μg/L
Inorganic Analytes:									
Perchlorate	65	37	22	6	μg/L	0.092	μg/L	170,000	μg/L

Only analytes positively detected in groundwater or surface water samples are presented in this table.

For a complete list of constituents analyzed, refer to the laboratory data package.

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

(2) - California Department of Health Services state drinking water notification level.

MCL - California Department of Health Services Maximum Contaminant Level.

DWNL - California Department of Health Services state drinking water notification level.

" - " MCL/DWNL not established.

 $\mu g/L$ - $\;$ Micrograms per liter.

NDMA - N-Nitrosodimethylamine

Table 3-7 Summary Statistics of Validated Detected Organic and Inorganic Analytes - First Ouarter 2010

Compounds Detected	Total Number of Samples Analyzed (1)	Total Number of Detections (1)	Number of Detections Exceeding MCL or DWNL (1)	MCL	ponding (unless / DWNL	Minim Concent Detec	ration	Maxin Concent Detec	ration
Organic Analytes:									
Methylene Chloride	5	1	0	5	μg/L	0.20	μg/L	0.20	μg/L
Inorganic Analytes:									
Perchlorate	13	8	1	6	μg/L	0.17	μg/L	11	μg/L
Total Alkalinity (as CaCO3)	2	2	0	-	mg/L	100	mg/L	110	mg/L
Bicarbonate (as CaCO3)	2	2	0	-	mg/L	120	mg/L	130	mg/L
Total Dissolved Solids	2	2	0	500	mg/L	410	mg/L	420	mg/L
Chloride	2	2	0	250	mg/L	46	mg/L	46	mg/L
Sulfate	2	2	0	250	mg/L	160.0	mg/L	160	mg/L
Calcium	2	2	0	-	mg/L	39.0	mg/L	42	mg/L
Magnesium	2	2	0	-	mg/L	7.1	mg/L	8	mg/L
Potassium	2	2	0	-	mg/L	4.9	mg/L	6	mg/L
Sodium	2	2	0	-	mg/L	100	mg/L	110	mg/L
Notes: (1) -	For a complete	list of constituents	in groundwater or surfa analyzed, refer to the l aple duplicates, trip bla	aboratory	data packa	ge.	n this tab	le.	

California Department of Health Services state drinking water notification level.

California Department of Health Services Maximum Contaminant Level.

California Department of Health Services state drinking water notification level.

MCL/DWNL not established.

 $\mu g/L$ -Micrograms per liter.

Milligrams per liter mg/L -

3.5 CHEMICALS OF POTENTIAL CONCERN

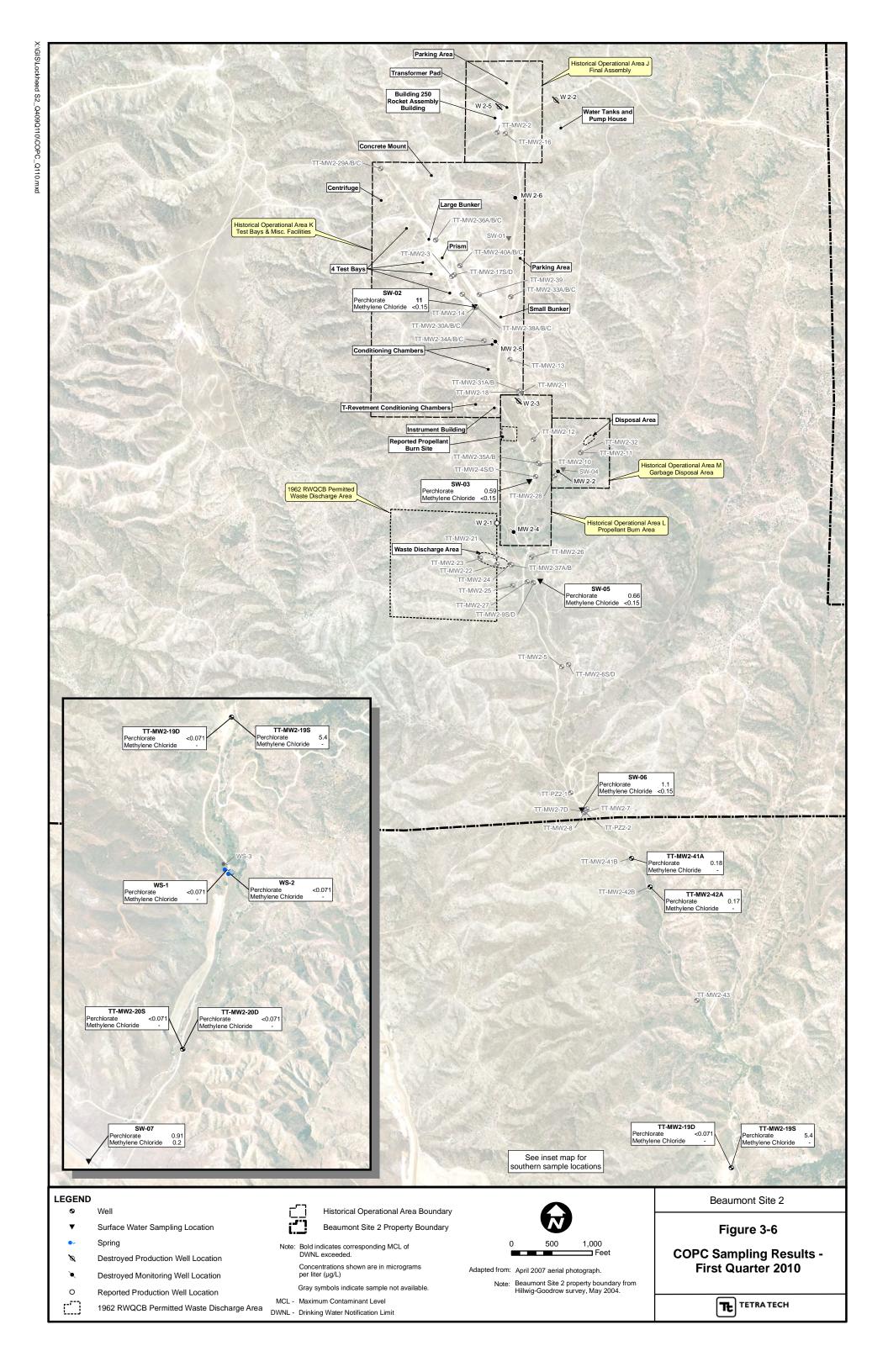
COPC evaluation and trend analysis are performed annually as part of the Second Quarter monitoring event. The analytes detected were screened against the MCLs or DWNLs (if an MCL is not established). The analytes were organized and evaluated in two groups, organic and inorganic analytes, and divided into primary and secondary COPCs. Table 3-8 presents a summary of the Site 2 COPCs. Laboratory analytical results from the Fourth Quarter 2009 and First Quarter 2010 monitoring events are presented in the following two subsections. Data which are B qualified because of association with either laboratory or field contamination is not included in the COPC evaluation. Figures 3-5 and 3-6 present summaries of COPC laboratory results for groundwater samples collected for the Fourth Quarter 2009 and First Quarter 2010.

Table 3-8 Groundwater Chemicals of Potential Concern

Analyte	Classification					
Perchlorate	Primary					
Trichloroethene	Primary					
Methylene Chloride	Primary					
1,4-Dioxane	Primary					
RDX	Secondary					
Notes:						
RDX - Hexahydro-1,3,5-trinitro-1,3,5-triazine						

3.5.1 Organic Analytes

Seven organic analytes (1,4-dioxane, 1,2-DCA, 1,1-DCE, benzene, methylene chloride, NDMA, and TCE) were detected above a published MCL or DWNL during the Fourth Quarter 2009 and First Quarter 2010 monitoring events. Tables 3-4 and 3-5 presents summaries of validated organic analyte concentrations reported in groundwater samples collected during the Fourth Quarter 2009 and First Quarter 2010 groundwater monitoring events.



1,4-Dioxane was reported in groundwater samples collected from five monitoring wells: TT-MW2-5, TT-MW2-9S, TT-MW2-22, TT-MW2-24, and TT-MW2-37A, during the Fourth Quarter 2009 monitoring event at concentrations ranging from 1.0 μ g/L to 320 μ g/L. All wells are located within or just downgradient of the former WDA. The DWNL for 1,4-dioxane is 3 μ g/L.

1,2-DCA was reported in groundwater samples collected from two monitoring wells, TT-MW2-22 and TT-MW2-24, located in the former WDA, during the Fourth Quarter 2009 monitoring event at concentrations of 1.7 μ g/L and 0.78 μ g/L respectively. The MCL for 1,2-DCA is 0.5 μ g/L.

1,1-DCE was reported in groundwater samples collected from two monitoring wells, TT-MW2-22 and TT-MW2-24, located in the former WDA, during the Fourth Quarter 2009 monitoring event at concentrations of 29 μ g/L and 2.7 μ g/L respectively. The MCL for 1,1-DCE is 6 μ g/L.

Benzene was reported in groundwater samples collected from nine monitoring wells, TT-MW2-22, TT-MW2-31B, TT-MW2-33C, TT-MW2-35B, TT-MW2-36C, TT-MW2-37A, TT-MW2-40B, and TT-MW2-40C during the Fourth Quarter 2009 monitoring event, at concentrations ranging from $0.21 \,\mu\text{g/L}$ to $21 \,\mu\text{g/L}$. The MCL for benzene is $1 \,\mu\text{g/L}$.

During Fourth Quarter 2009, methylene chloride was reported in groundwater samples collected from nine monitoring wells, TT-MW2-11, TT-MW2-21, TT-MW2-22, TT-MW2-28, TT-MW2-33B, TT-MW-34A, TT-MW2-34B, TT-MW2-34C, and TT-MW2-39. With the exception of monitoring wells TT-MW2-21 and TT-MW2-22, all detections were at or near the method detection limit of 0.15 μ g/L. Methylene chloride was detected in monitoring wells TT-MW2-21 and TT-MW2-22, at concentrations of 4.9 μ g/L and 9.3 μ g/L respectively. During First Quarter 2010, methylene chloride was detected in surface water sample SW-07 at a concentration of 0.20 μ g/L. Methylene chloride has previously been reported in groundwater samples collected from monitoring well TT-MW2-22 at concentration up to 220 μ g/L. The MCL for methylene chloride is 5 μ g/L.

NDMA was reported in groundwater samples collected from 11 monitoring wells during the Fourth Quarter 2009 monitoring event at concentrations ranging from 0.0012 μ g/L to 0.011 μ g/L. The DWNL for NDMA is 0.01 μ g/L.

TCE was reported in groundwater samples collected from four monitoring wells, TT-MW2-21, TT-MW2-22, TT-MW2-24, and TT-MW2-37A, located in the former WDA during the Fourth Quarter 2009 monitoring event at concentrations of 2.8 μ g/L, 460 μ g/L, 100 μ g/L, and 0.86 μ g/L respectively. TCE was also reported in monitoring well TT-MW2-11, located in Area M at concentrations of 9.2 μ g/L. The MCL for TCE is 5 μ g/L. Time-series graphs of TCE are provided in Appendix I.

Other organic analytes detected at low levels and below their respective MCLs or DWNLs during the Fourth Quarter 2009 and First Quarter 2010 groundwater monitoring events were acetone, 2-butanone, carbon disulfide, chloromethane, chloroform, 1,1-DCA, cis-1,2-DCE, trans-1,2-DCE, ethylbenzene, 2-hexanone, 4-methyl-2-pentanone, toluene, 1,1,2-trichloroethane, tetrachloroethene (PCE), m,p-xylenes, and o-xylenes. None of these compounds exceeded their MCL or DWNL, and generally they are not detected consistently from monitoring event to event.

3.5.2 Organic COPCs

Based on the analysis above and the concentrations detected during previous groundwater monitoring events, TCE, methylene chloride, and 1,4-dioxane have been identified as primary COPCs at the Site. Based on the limited and relatively low RDX concentrations reported in groundwater samples collected from the Site, RDX is regarded as a secondary COPC. NDMA will be further evaluated to determine whether the source is groundwater contamination or if it is being introduced as cross contamination from an outside source (i.e., dedicated sampling pumps). The remaining 16 organic analytes were either detected below their respective MCL or DWNL or at relatively low concentrations. Their distribution and concentration in groundwater will continue to be monitored and the results evaluated.

3.5.3 Inorganic Analytes

One inorganic analyte, perchlorate, was detected in groundwater above a published MCL or DWNL. Tables 3-4 and 3-5 present a summary of validated inorganic analyte concentrations reported in groundwater samples collected during the Fourth Quarter 2009 and First Quarter 2010 groundwater monitoring events.

Perchlorate was reported in groundwater samples collected from 37 of 65 locations sampled during the Fourth Quarter 2009 groundwater monitoring event and eight of 13 locations sampled during the First Quarter 2010 monitoring event, at concentrations up to 170,000 μ g/L. The

California MCL for perchlorate is 6 μ g/L. Time-series graphs of perchlorate are provided in Appendix I.

3.5.4 Inorganic COPCs

Based on the analysis above and the concentrations detected during previous groundwater monitoring events, perchlorate is the only inorganic primary COPC identified at the Site. No secondary COPCs were identified. Metals will continue to be evaluated on an annual basis and as additional monitoring points are added to the network.

3.6 NEW MONITORING WELLS

Three groundwater monitoring wells, TT-MW-41A, TT-MW2-42A, and TT-MW2-43, three piezometers, TT-MW2-41B, TT-MW2-42B, and TT-PZ2-2, and one extraction well, TT-EW2-1, were installed between September 2009 and October 2009 as part of the Site 2 Additional Offsite Well Installation and Aquifer Testing study (Tetra Tech, 2010c). Following completion of the well construction and development activities, as part of the installation process groundwater samples were collected from the newly installed monitoring wells and analyzed for perchlorate, and select wells were analyzed for VOCs, 1,4-dioxane, NDMA, and natural attenuation parameters. A complete description of the monitoring well installation activities and results is presented in the Site 2 Additional Offsite Well Installation and Aquifer Testing Report (Tetra Tech, 2010c).

Table 3-9 presents a summary of validated organic and inorganic analyte concentrations reported in new well samples collected as part of the installation process and during the Fourth Quarter 2009 and First Quarter 2010 groundwater monitoring events.

Table 3-9 Summary of Validated Detected COPCs in New Wells

Sample Location	Sample Date	Perchlorate 1,4-Dioxane ND		NDMA	Methylene Chloride	Trichloroethene					
All results reported in µg/L unless otherwise stated											
TT-EW2-1	10/23/2009	2.6	0.2	< 0.0007	< 0.15	< 0.17					
TT-EW2-1	10/30/2009	5.4	NA	NA	NA	NA					
TT-MW2-41A	10/21/2009	0.66	< 0.10	0.0030	< 0.15	< 0.17					
TT-MW2-41A	11/23/2009	0.36	< 0.10	0.0030	< 0.15	< 0.17					
TT-MW2-41A	12/16/2009	< 0.071	< 0.10	0.0042 Bk	< 0.15	< 0.17					
TT-MW2-41A	2/17/2010	0.18	NA	NA	NA	NA					
TT-MW2-41B	12/16/2009	0.11	NA	NA	NA	NA					
TT-MW2-42A	10/22/2009	< 0.071	< 0.10	0.002 Je	0.62 Jq	< 0.17					
TT-MW2-42A	11/23/2009	< 0.071	< 0.10	< 0.0007	< 0.15	< 0.17					
TT-MW2-42A	12/16/2009	< 0.071	< 0.10	0.0020 Bk	< 0.15	< 0.17					
TT-MW2-42A	2/17/2010	0.17	NA	NA	NA	NA					
TT-MW2-42B	12/16/2009	0.099 Jq	NA	NA	NA	NA					
Method Detection Limit		0.071	0.10	0.0007	0.15	0.17					
MCL (unless noted) / DWNL		6	3 (1)	0.01 (1)	5.0	5.0					

Notes: Only analytes positively detected in samples are presented in this table.

For a complete list of constituents analyzed, refer to the laboratory data package.

NDMA - N-Nitrosodimethylamine

μg/L - Micrograms per liter

MCL - California Department of Health Services Maximum Contaminant Level.

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

(1) - DWNL

" - " MCL/DWNL not established.

Bold - Maximum Contaminant Level exceeded.

< # - Method detection limit concentration is shown.

NA - Not analyzed

B - The sample result was less than 5 times blank contamination. Cross contamination is suspected.

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

e - A holding time violation occurred.

k - The analyte was found in the field blank.

q - The analyte deteced was below the Practical Quantitation Limit (PQL).

3.7 SURFACE WATER SAMPLING RESULTS

Surface water samples were collected for perchlorate at three locations, WS-1, WS-2 and WS-3, from a spring on the former Wolfskill property during the Fourth Quarter 2009 and for perchlorate and general minerals parameters at two locations, WS-1 and WS-2, during First Quarter 2010 (Figure 2-4). Perchlorate was not detected above the MCL in WS-1, WS-2, or WS-3 during Fourth Quarter 2009 or First Quarter 2010. The MCL for perchlorate is 6 µg/L.

Storm water samples were collected from five locations during storm events on 27 January and 6 February 2010. Storm water samples were tested for VOCs, and perchlorate. Methylene chloride was detected in one sample, SW-07, at a concentration of 0.20 μ g/L. No other VOCs were detected. Perchlorate was detected in all five storm water samples at concentrations ranging from 0.59 μ g/L to 11 μ g/L. No other surface water samples were collected during this reporting period.

Table 3-10 presents a summary of validated organic and inorganic analyte concentrations reported in surface water and storm water samples collected during the Fourth Quarter 2009 and First Quarter 2010 groundwater monitoring events.

Table 3-10 Summary of Validated COPCs in Surface Water

Sample Name	Sample Date	Methylene Chloride - ug/L	Perchlorate µg/L						
All results reported in µg/L unless otherwise stated									
WS-1	12/2/2009	NA	< 0.35						
WS-1	2/16/2010	NA	< 0.071						
WS-2	12/2/2009	NA	< 0.35						
WS-2	2/16/2010	NA	< 0.071						
WS-3	12/2/2009	NA	< 0.35						
SW-02	2/6/2010	< 0.15	11						
SW-03	2/6/2010	< 0.15	0.59						
SW-05	2/6/2010	< 0.15	0.66						
SW-06	2/6/2010	< 0.15	1.1						
SW-07	1/27/2010	0.20 Jbq	0.91						
Method l	Detection Limit	0.15	0.071						
MCL (unless	noted) / DWNL	5	6						

Notes:

μg/L - Micrograms per liter

MCL - California Department of Health Services Maximum Contaminant Level.

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

< # - Method detection limit concentration is shown.

3.8 MONITORED NATURAL ATTENUATION SAMPLING

Eleven monitoring wells, six associated with the groundwater perchlorate plume originating from the former Test Bay Area (TT-MW2-36A, TT-MW2-17D, TT-MW2-14, TT-MW2-1, TT-MW2-12 and TT-MW2-10), and five associated with the groundwater perchlorate plume originating from the former WDA (TT-MW2-23, TT-MW2-24, TT-MW2-9S, TT-MW2-5, and TT-MW2-7) were sampled and analyzed for monitored natural attenuation parameters (MNA) during the Fourth Quarter 2009 monitoring event. Samples for laboratory analysis were analyzed for total organic carbon (TOC), dissolved organic carbon (DOC), total iron, sulfate, methane, hydrogen, and volatile fatty acids (VFAs). Due to the short laboratory holding times associated with ferrous iron and sulfide, ferrous iron and sulfide were analyzed in the field just prior to sampling using a field instrument. Additionally, DO and ORP were monitored with field instruments during purging and sampling. Figure 3-7 presents monitoring well locations sampled for MNA during the Fourth Quarter 2009 monitoring event. Table 3-11 presents a summary of detected analytes and field measurements.

Former Test Bay Area

The wells selected for natural attenuation parameter sampling in the Test Bay perchlorate plume form a longitudinal traverse extending from TT-MW2-36A, which lies upgradient of the perchlorate plume, to wells TT-MW2-10 and TT-MW2-12, which lie downgradient of the perchlorate plume. Aquifer conditions at the upgradient well (TT-MW2-36A) are anaerobic, based on the low DO (<1 mg/L) and ORP (<100 mV). The hydrogen concentration (2 nmol/L) is suggestive of anaerobic conditions, and is within the range of sulfate reduction, consistent with the relatively low sulfate concentration (22 mg/L) and relatively high sulfide concentration (0.34 mg/L). Methane concentrations are somewhat elevated (68 μ g/L), and are also consistent with anaerobic conditions. Nitrate, which competes with perchlorate for organic carbon, was not detected. Upgradient aquifer conditions therefore appear to be generally conducive to perchlorate biodegradation.

Aquifer conditions within the perchlorate plume range from mildly anaerobic in TT-MW2-17D (DO = 0.65 mg/L and ORP = 53 mV) to mildly aerobic in TT-MW2-14 (DO = 4.4 m/L and ORP = 81 mV) and TT-MW2-1 (DO = 4.5 mg/L and ORP = 103 mV). In contrast, hydrogen concentrations are relatively high (3.5 to 6.5 nmol/L), suggestive of sulfate-reducing to methanogenic conditions. The hydrogen data do not appear to be consistent with the DO and ORP measurements, the presence of sulfate concentrations greater than upgradient well TT-MW2-36A, very low sulfide concentrations, and very low methane concentrations. The hydrogen results may be an artifact caused by minor corrosion of the stainless steel well screens. Organic carbon concentrations are relatively low, and nitrate concentrations in all three wells are elevated, ranging from 7.2 to 15 mg/L. The apparent slightly anaerobic to aerobic aquifer conditions and elevated nitrate concentrations suggest that conditions are unfavorable for perchlorate biodegradation.

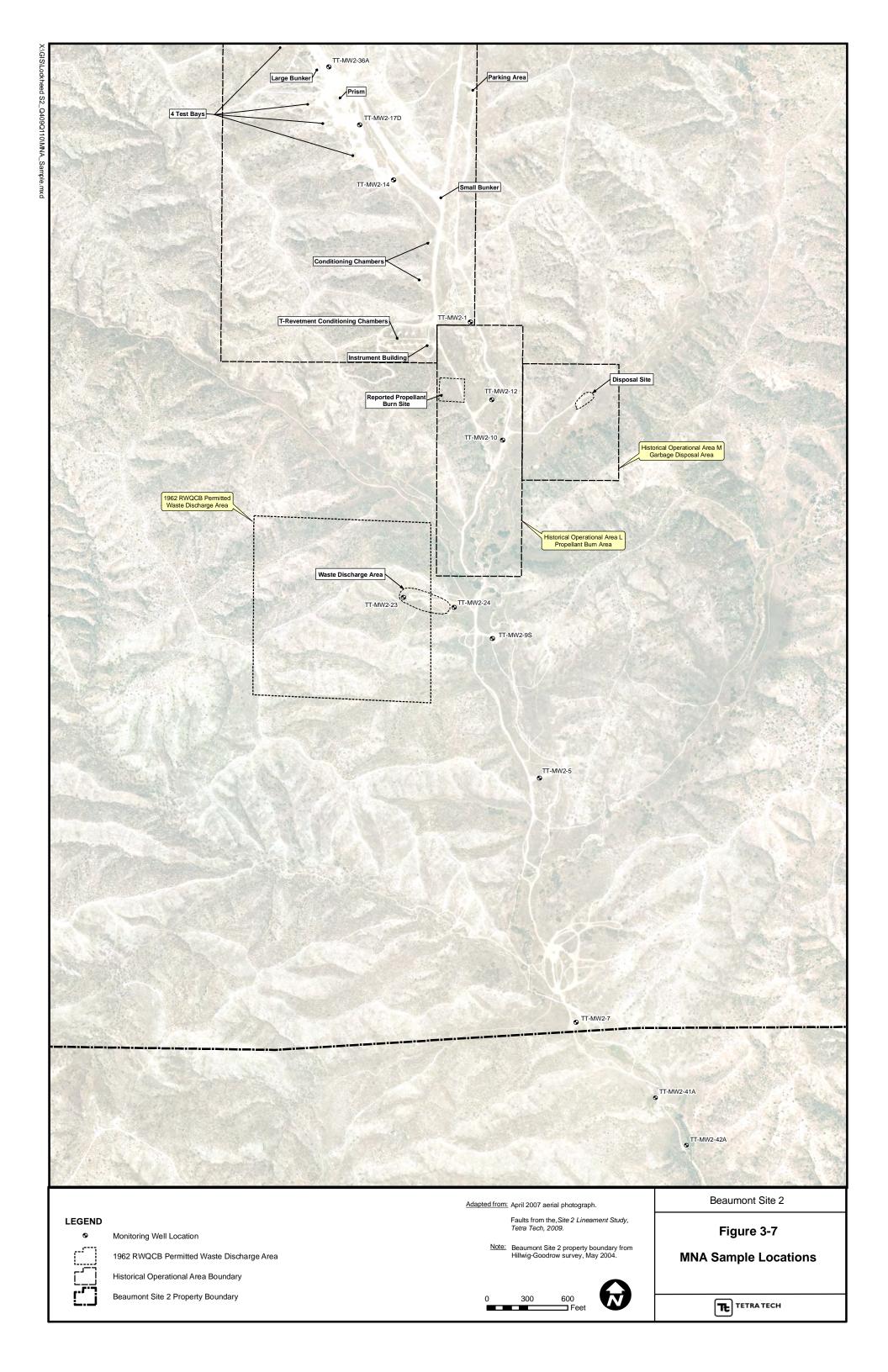


Table 3-11 Summary of Validated Detected Natural Attenuation Analytes and Field Measurements – Fourth Quarter 2009

Field Parameters				Analytes													
Sample Location	Sample Date	DO - mg/L	ORP - mVs	Sulfide -mg/L	Ferrous Iron - mg/L	Per chlorate -ug/L	Acetic Acid	Hexanoic Acid - mg/L	Lactic Acid and HIBA - mg/L	Propionic Acid - mg/L	Dissolved Organic Carbon - mg/L	Total Organic Carbon -mg/L	Hydrogen -nM	Methane - ug/L	Nitrate (as N) - mg/L	Sulfate -mg/L	Iron - mg/L
TT-MW2-1	12/9/09	4.45	102.9	0.00	0.00	7,700	0.1 Ba	< 0.007	0.27 Ba	0.035 Jq	0.98	0.86	6.5	0.06 BJaq	7.2	31 Jd	< 0.012
TT-MW2-5	12/3/09	4.66	159.1	0.01	0.11	930	0.11	0.01 Jq	< 0.042	< 0.002	0.93	0.80	9.9	0.08 Jq	10	140	0.047
TT-MW2-7	12/3/09	0.37	196.0	0.00	0.04	410	0.085	< 0.007	< 0.042	0.064 BJaq	1.6	1.2	3.4	0.04 Jq	5.4	190	< 0.012
TT-MW2-9S	12/8/09	1.11	92.8	0.00	0.06	4,700	0.061 BJaq	< 0.007	0.19 Ba	< 0.002	3.6	1.4	5	0.08 BJaeq	11	130	< 0.012
TT-MW2-10	12/9/09	3.11	110.7	0.00	0.03	< 0.35	0.06 BJaq	< 0.007	0.16 Ba	0.034 Jq	3.1	1.0	11	0.08 BJaq	< 0.11	61 Jd	0.16
TT-MW2-12	12/14/09	2.82	57.5	0.01	0.03	< 0.71	0.073 Ba	0.3	0.15 Ba	< 0.002	5.2	1.2	5.4	0.76	< 0.11	50	0.14
TT-MW2-14	12/14/09	4.36	80.8	0.00	0.04	47,000	0.046 BJaq	< 0.007	0.14 Ba	< 0.002	2.5	1.9	3.5	0.17 Ba	15	180	0.11
TT-MW2-17D	12/10/09	0.65	52.7	0.02	0.00	53,000	0.042 BJaq	0.11	0.097 BJaq	< 0.002	1.2	0.94	4.1	0.17 Ba	8.8 Je	55	< 0.012
TT-MW2-23	12/11/09	0.68	91.9	0.07	0.05	< 0.071	0.23	< 0.007	< 0.042	< 0.002	0.52 Jq	0.17 Jq	1.4	19	< 0.11	25	1.2
TT-MW2-24	12/11/09	0.80	109.6	0.00	0.02	140,000	0.16 BJaf	< 0.007	< 0.042	< 0.002	7.5	7.2	0.91	0.2	61 Je	95	0.013 Jq
TT-MW2-36A	12/10/09	0.78	-124.9	0.34	0.00	< 0.071	0.11	< 0.007	0.26 Ba	< 0.002	1.3 Jf	1.1	2	68	<0.11 UJe	22	0.027
TT-MW2-41A	11/23/2009	0.65	-35.3	0.41	0.33	0	0.052 BJkq	< 0.007	0.23 Bk	< 0.002	1.6	11	-	1.2	0.34	130	11.0
TT-MW2-42A	11/23/2009	0.93	-6.1	0.03	0.03	< 0.071	0.11	< 0.007	0.4	< 0.002	0.74 Ba	8.2	-	3.6	< 0.11	46	0.28
Method De	tection Limit	-	-	-	-	0.071	0.07	0.07	0.10	0.07	0.021	0.021	0.6	0.00784	0.11	0.37	0.012
MCL/D	WNL (µg/L)	-	-	-	-	6	-	-	-	-	-	-	-	-	10	250	0.3

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

mg/L - milligrams per liter

μg/L - micrograms per liter.

nM - nanomoles

mV - millivolts

MCL - Maximum Contaminant Level.

DWNL - California Department of Health Services state drinking water notification level.

- "-" Not available.
- <# Analyte not detected, method detection limit concentration is shown.

Bold - Maximum Contaminant Level exceeded.

- B The sample result was less than 5 times blank contamination. Cross contamination is suspected.
- J The analyte was positively identified, but the concentration is an estimated value.
- U The analyte was analyzed for , but was not detected above the method detection limit.
- a The analyte was found in the method blank.
- d The Laboratory Control Sample (LCS) recovery was outside control limits.
- e A holding time violation occurred.
- f The duplicate Relative Percent Differnce was outside the control limit.
- q -The analyte detection was below the Practical Quantitation Limit (PQL).

Conditions in downgradient wells TT-MW2-12 and TT-MW2-10 are mildly aerobic based on DO concentrations (2.8 to 3.2 mg/L) and ORP values (58 to 111 mV). As noted for the wells within the plume, hydrogen concentrations (5.4 to 11 nmol/L) are suggestive of highly reducing conditions, which is not consistent with the relatively high sulfate, low sulfide, and low methane concentrations. As previously noted, the hydrogen concentrations may be an artifact caused by minor corrosion of the stainless steel well screens. Organic carbon concentrations are relatively low, but still higher than those observed in the plume wells, and nitrate was not detected in either well. The apparently aerobic aquifer conditions are unfavorable for perchlorate biodegradation.

Waste Discharge Area

The wells selected for natural attenuation parameter sampling in the WDA perchlorate plume form a longitudinal traverse extending from TT-MW2-23, which lies within the WDA upgradient of the perchlorate plume, to wells TT-MW2-42A, which lies downgradient of the perchlorate plume and to the south of the site. Aquifer conditions at upgradient well TT-MW2-23 are mildly anaerobic, as indicated by the low DO (0.7 mg/L) and ORP (92 mV). The hydrogen concentration (1.4 nmol/L) also suggests mildly anaerobic redox conditions, which are compatible with the slightly elevated dissolved iron (1.2 mg/L), relatively low sulfate concentration (25 mg/L), slightly elevated sulfide (0.07 mg/L), and slightly elevate methane (19 µg/L). Organic carbon concentrations are relatively low, and nitrate was not detected. Upgradient aquifer conditions appear to be generally conducive to perchlorate biodegradation.

Aquifer conditions within the plume range from mildly anaerobic in wells TT-MW2-24, TT-MW2-9S (DO 0.8 and 1.1 mg/L, ORP 110 to 93 mV, respectively) to mildly aerobic in TT-MW2-5 and TT-MW2-7 (DO 0.37 to 4.7, ORP 159 to 196 mV, respectively). The hydrogen concentration in TT-MW2-24 (which has a PVC well screen) is 0.91 nmol/L, consistent with the mildly anaerobic conditions suggested by the DO and ORP data. Hydrogen concentration in TT-MW2-9S, TT-MW2-5, and TT-MW2-7 range from 3.4 to 9.9 nmol/L, which are suggestive of highly reducing conditions and inconsistent with the DO and ORP data. As previously noted, the hydrogen concentrations may be an artifact caused by minor corrosion of the stainless steel well screens. Sulfate concentrations are high and sulfide concentrations are low, which supports mildly anaerobic to mildly aerobic aquifer conditions. Total organic carbon concentrations are relatively high (7.5 mg/L) in TT-MW2-24, and are lower (3.6 to 0.93 mg/L) in the downgradient wells.

Nitrate concentrations are elevated in all wells, ranging from 5.4 to 61 mg/L. Aquifer conditions within the plume do not appear to be conducive to perchlorate biodegradation.

Downgradient wells TT-MW2-41A and TT-MW2-42B are located in a riparian corridor south of the site property boundary. Both wells have low DO (0.65 to 0.9 mg/L) and low ORP (-6 to -35 mV), which are suggestive of anaerobic conditions. Dissolved iron and sulfide concentrations (11 and 0.41 mg/L, respectively) are elevated in TT-MW2-41A; methane is slightly elevated in both wells (1.2 µg/L in TT-MW2-41A, 3.6 µg/L in TT-MW2-42A). These observations are consistent with the presence of reducing conditions. Total organic carbon concentration are low, but dissolved organic carbon concentrations are relatively high (8.2 to 11 mg/L). Nitrate concentrations are low to non-detectable. Aquifer conditions in the downgradient area appear to be conducive to perchlorate biodegradation.

3.9 HABITAT CONSERVATION

Consistent with the U.S. Fish and Wildlife Service (USFWS) approved HCP (USFWS, 2005) and subsequent clarifications (LMC, 2006a and 2006b) of the HCP describing environmental activities proposed at the Site, all field activities were performed under the supervision of a USFWS approved biologist who monitored each work location. Groundwater sampling activities were conducted with light duty vehicles and, as specified in the Low Affect HCP, do not require biological monitoring. As a result, no impact to SKR occurred during the performance of the field activities related to the Fourth Quarter 2009 and First Quarter 2010 monitoring events.

SECTION 4 SUMMARY AND CONCLUSIONS

This section summarizes the results of the Fourth Quarter 2009 and First Quarter 2010 groundwater monitoring events. During the Fourth Quarter 2009 monitoring event 69 monitoring well locations and two piezometers were measured for groundwater levels and 62 monitoring wells and three surface water locations were sampled for groundwater quality. Two monitoring wells, TT-MW2-29A and TT-MW2-43, were found to be dry during the Fourth Quarter 2009 monitoring event. During the First Quarter 2010 monitoring event 69 monitoring wells, two surface water locations, and five storm water locations were sampled for groundwater quality. Two monitoring wells, TT-MW2-29A and TT-MW2-43, were found to be dry during the First Quarter 2010 monitoring event.

4.1 GROUNDWATER ELEVATION AND FLOW

During the Fourth Quarter 2009 and First Quarter 2010 monitoring events, groundwater elevations at the Site ranges from about 2,075 feet msl in the northern portion of the Site, to about 1,819 feet msl in the southern portion of the Site. Depth to groundwater ranged from about 118 feet bgs to about 15 feet bgs.

Based on the measured groundwater elevations, the current CSM, and the southward sloping topography at the Site, groundwater flow in the QAL/wSTF and STF screened wells appears to be southerly and generally follows the topography of Laborde Canyon. Groundwater flow will be refined as additional data are acquired.

Generally, groundwater elevations at the Site are relatively stable and demonstrated a limited seasonal rise and fall. The exception is the shallow wells near the property boundary that appear to show stronger seasonal fluctuations. Although the data is limited, the overall long term trend appears to correspond to the long term precipitation pattern.

4.1.1 Groundwater Gradients

Horizontal groundwater gradients across the Site are relatively constant. The horizontal groundwater gradients calculated between TT-MW2-16 and TT-MW2-6S from the Fourth Quarter

2009 and First Quarter 2010 groundwater monitoring events for the QAL/wSTF screened wells averaged 0.030 ft/ft. The horizontal groundwater gradients calculated between TT-MW2-2 and TT-MW2-6D for the Fourth Quarter 2009 and First Quarter 2010 groundwater monitoring events for the deeper STF screened wells averaged 0.029 ft/ft.

Generally the vertical gradients are downward on site and upward from the Site boundary south. The vertical gradients range from negative 0.28 to positive 0.19. A summary of calculated horizontal and vertical groundwater gradients is presented in Table 3-3 and in Appendix E.

4.2 WATER QUALITY MONITORING

Groundwater samples collected during the Fourth Quarter 2009 and First Quarter 2010 monitoring events were analyzed for perchlorate. Select locations were analyzed for VOCs, natural attenuation parameters, general minerals, 1,4 dioxane, and NDMA. Based on the historical operations at the Site and groundwater monitoring results, perchlorate, TCE, methylene chloride, and 1,4-dioxane were identified as a primary COPCs. RDX was identified as a secondary COPC. NDMA will be further evaluated to determine whether the source is groundwater contamination or if it is being introduced as cross contamination from an outside source.

Perchlorate has not been detected in the groundwater above the MCL (6.0 μg/L) in Area J. In Area K, perchlorate in the groundwater was detected at concentrations ranging from below the MDL to 170,000 μg/L during Fourth Quarter 2009 and First Quarter 2010. Previously, perchlorate was detected as high as 190,000 μg/L. A source of perchlorate was identified in Area K. In Area L, downgradient of both Areas J and K, perchlorate in the groundwater was detected at concentrations ranging from below the MDL to 0.54 μg/L during Fourth Quarter 2009 and First Quarter 2010. Previously, perchlorate was detected as high as 2.1 μg/L. There are currently no indications that a source of perchlorate is present in Area L; the perchlorate detected in Area L groundwater appears to have originated in Area K. In Area M, the Garbage Disposal area, perchlorate in the groundwater was detected at concentrations ranging from below the MDL to 280 μg/L during Fourth Quarter 2009 and First Quarter 2010. Previously, perchlorate was detected as high as 469 μg/L. Area M has been identified as a source of perchlorate in groundwater. In the former WDA, downgradient of the operational areas J, K, L, and M, perchlorate in the groundwater was detected at concentrations ranging from below the MDL to 140,000 μg/L during Fourth Quarter 2009 and First Quarter 2010. Previously, perchlorate was detected as high as

190,000 μ g/L. The former WDA has been identified as a source of perchlorate in groundwater. In the lower section of Laborde Canyon, downgradient of the operational areas and the former WDA, perchlorate in the groundwater was detected at concentrations ranging from below the MDL to 4,700 μ g/L during Fourth Quarter 2009 and First Quarter 2010. Previously, perchlorate was detected as high as 4,100 μ g/L in the lower section of Laborde Canyon and as high as 519 μ g/L at the southern site boundary. No source of perchlorate has been identified at the southern site boundary. On the former Wolfskill property, south of the southern site boundary, perchlorate was detected in TT-MW1-19S during Fourth Quarter 2009 and First Quarter 2010 at concentrations of 5.7 μ g/L and 5.4 μ g/L respectively.

TCE was reported in groundwater samples collected from monitoring well TT-MW2-11 located in Area M at a concentration of 9.2 μ g/L during the Fourth Quarter 2009. In the former WDA, TCE was detected in monitoring wells TT-MW2-21, TT-MW2-22, TT-MW2-24, and TT-MW2-37A at concentrations of 2.8 μ g/L, 460 μ g/L, 100 μ g/L, and 0.86 μ g/L, respectively during the Fourth Quarter 2009. The TCE MCL is 5 μ g/L. Based on the data available at this time, the extent of the TCE plumes in groundwater appear to be isolated to small areas, and it does not extend offsite.

Methylene chloride was reported in groundwater samples collected from monitoring wells TT-MW2-21 and TT-MW2-22 located in the former WDA at concentrations of 4.9 μ g/L and 9.2 μ g/L respectively during the Fourth Quarter 2009. Additionally, methylene chloride was detected at or near the method detection limit of 0.15 μ g/L during Fourth Quarter 2009 in monitoring wells TT-MW2-11, TT-MW2-28, TT-MW2-33B, TT-MW2-34A, TT-MW2-34B, TT-MW2-34C, and TT-MW2-30 and in storm water sample SW-07 during First Quarter 2010. The methylene chloride MCL is 5 μ g/L. Based on the data available at this time, the extent of the methylene chloride plumes in groundwater appear to be isolated, and do not extend offsite.

1,4-dioxane was reported in groundwater samples collected from monitoring wells TT-MW2-5, TT-MW2-9S, TT-MW2-22, TT-MW2-24, TT-MW2-37A, which are located in or downgradient of the former WDA, at concentrations ranging from 1.0 μ g/L to 320 μ g/L during Fourth Quarter 2009. The 1,4-dioxane DWNL is 3 μ g/L.

RDX samples are collected and analyzed only during the Second Quarter monitoring event so no samples were collected during this monitoring period. RDX was initially analyzed for as part of a

screening evaluation for emerging contaminants. The origin of the RDX is unknown. Based on the data available at this time, the extent of the RDX plume in groundwater appears to be isolated to Area K and the former WDA and does not extend offsite.

NDMA was detected in groundwater samples collected from 11 monitoring wells during the Fourth Quarter 2009 monitoring event at concentrations ranging from $0.0012 \,\mu\text{g/L}$ to $0.011 \,\mu\text{g/L}$. The DWNL for NDMA is $0.01 \,\mu\text{g/L}$ The origin of the NDMA is unknown. However, based on the data available at this time, the source of the NDMA contamination may be the site approved dedicated purging and sampling pumps.

Natural Attenuation Sampling

The objective of the MNA sampling and analyses effort is to understand the geochemical characteristics upgradient, within and downgradient from the Test Bay Area and WDA perchlorate plumes. A preliminary evaluation of MNA suggests that there is limited potential for natural perchlorate biodegradation, except within the riparian area downgradient of the WDA perchlorate plume.

4.3 GROUNDWATER MONITORING PROGRAM AND THE GROUNDWATER QUALITY MONITORING NETWORK

Twenty-two quarters of water quality monitoring have been conducted at the Site since the September 2004 well installation activities. Groundwater samples have been routinely analyzed for VOCs and perchlorate. Selected testing for CAM 17 metals, general minerals, 1,4-dioxane, RDX, NDMA, 1,2,3-TCP and hexavalent chromium has also been performed. A groundwater monitoring sampling and analysis plan (SAP) was prepared to optimize and better define the GMP at the Site (Tetra Tech, 2007b). In concurrence with the DTSC, groundwater monitoring will be performed in accordance with the SAP.

Perchlorate, TCE, methylene chloride, and 1,4-dioxane have been identified as primary COPCs. All monitored wells will be tested for perchlorate semi-annually, and select wells will be sampled for VOCs and 1,4-dioxane semi-annually or annually. Because of the previous detections of RDX in TT-MW2-1 and TT-MW2-13, and the recent detection of RDX in TT-MW2-24, sampling for RDX will continue to be conducted annually in monitoring wells TT-MW2-1, TT-MW2-13, and TT-MW2-24.

Due to the uncertainty of the source of the contamination, select wells will continue to be sampled for NDMA semi-annually until the source of the contamination can be determined.

Because of previous detections of arsenic above the MCL, unfiltered metals will be analyzed for all monitored wells sampled annually during the Second Quarter sampling event until background levels for metals can be determined. Filtered metals will be collected for ecological risk assessment in water table monitoring wells where the water level is less than 25 feet below ground surface, and where the well screen is less than 25 feet below ground surface.

The analytical scheme is evaluated annually during the Second Quarter of each year and changes may be proposed to accommodate expanded site knowledge or changing site conditions. The classifications of the wells in the network and the corresponding sampling frequency is also evaluated annually during the Second Quarter of each year and modified to accommodate expanded Site knowledge or changing Site conditions.

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