

**GROUNDWATER INFORMATION SHEET**

**Hexavalent Chromium**

*The purpose of this groundwater information sheet is to provide general information regarding a specific constituent of concern (COC). The information provided herein relates to groundwater sources used for public drinking water, not water served at the tap.*

<b>GENERAL INFORMATION</b>	
<b>Constituent of Concern</b>	Hexavalent Chromium
<b>Synonyms</b>	Chromium VI, Chromium Six, Chrome 6, Cr <sup>6</sup>
<b>Chemical Formula</b>	Cr <sup>6</sup>
<b>CAS No.</b>	18540-29-9
<b>Storet No.</b>	01032
<b>Summary</b>	In 2014, California State Water Resources Control Board (SWRCB) established a Maximum Contaminant Level (MCL) for hexavalent chromium at 10 micrograms per liter (µg/L) that was withdrawn by SWRCB in August 2017. Until a revised MCL is adopted by SWRCB, the total chromium MCL will be used as the drinking water standard. In order to show a spatial distribution of hexavalent chromium in public wells the Detection Limit for Purposes of Reporting (DLR) of 1 µg/L is used for this fact sheet. Based on SWRCB data from 2007 to 2017, 3,798 active and standby public water wells had concentrations of hexavalent chromium above the DLR. Most detections have occurred in Los Angeles (516), San Bernardino (430) and Fresno (324) counties.

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<b>REGULATORY AND WATER QUALITY LEVELS<sup>1</sup></b> <b>HEXAVALENT CHROMIUM</b>		
<b>Type</b>	<b>Agency</b>	<b>Concentration</b>
Federal MCL <sup>2</sup>	US EPA <sup>3</sup>	None established
State MCL	SWRCB <sup>4</sup>	In progress
Detection Limit for Purposes of Reporting (DLR)	SWRCB	1 µg/L for Cr6
Public Health Goal (PHG)	OEHHA <sup>5</sup>	0.02 µg/L
Others: HBSL <sup>6</sup> (non-cancer health effect) Cal/EPA Cancer Potency Factor as a drinking water level	USGS <sup>7</sup> Cal/EPA	20 µg/L 0.07 µg/L

<sup>1</sup> These levels generally relate to drinking water. Other water quality levels may exist. For further information see A Compilation of Water Quality Goals, 17<sup>th</sup> Edition (SWRCB, 2016).

<sup>2</sup> MCL - Maximum Contaminant Level

<sup>3</sup> US EPA - United States Environmental Protection Agency

<sup>4</sup> SWRCB—State Water Resources Control Board

<sup>5</sup> OEHHA = Office of Environmental Health and Human Hazard Assessment

<sup>6</sup> HBSL - Health Based Screening Level

<sup>7</sup> USGS – United State Geological Survey

<b>HEXAVALENT CHROMIUM DETECTIONS IN PUBLIC WATER WELLS<sup>8</sup></b>	
<b>Detection Type</b>	<b>Number of Groundwater Sources</b>
Number of active and standby public wells with hexavalent chromium detections <sup>9</sup>	Hexavalent chromium was detected in 4,475 wells (8,800 wells tested)
Number of active and standby public water wells with Cr <sup>6</sup> concentrations above 1 µg/L.	3,798 wells
Top counties having public water wells with Cr <sup>6</sup> concentrations above 1 µg/L.	Los Angeles (516), San Bernardino (430), Fresno (324)

<sup>8</sup>Based on SWRCB 2007-2017 public and standby well data.

<sup>9</sup>Data from private domestic wells and wells with less than 15 service connections are not available.

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<b>ANALYTICAL INFORMATION</b>		
<b>Method</b>	<b>Detection Limit</b>	<b>Note</b>
<b>US EPA 218.7</b>	0.01 µg/L	By ion chromatography with post-column derivatization and UV-visible spectroscopic detection (SWRCB approved for drinking water)
<b>US EPA 218.6</b>	0.3 µg/L	By ion chromatography (SWRCB approved for drinking water)
<b>USGS by GFAAS</b>	0.05 µg/L	Cr <sup>6</sup> separation in the field, not time sensitive
<b>Known Limitations to Analytical Methods</b>	Water sample pH must be adjusted to 9.0-9.5, stored at 4° C and analyzed within 24 hours.	
<b>Public Drinking Water Testing Requirements</b>	In January 2001, hexavalent chromium was identified as an unregulated chemical requiring monitoring. As a result, public water systems began to test for hexavalent chromium in their drinking water supplies to the DLR of 1µg/L. In 2014, MCL for hexavalent chromium was established at 10 µg/L. On August 1, 2017 the MCL was repealed. Compliance with the MCL for total chromium continues to be required.	

<b>OCCURRENCE</b>	
<b>Anthropogenic Sources</b>	Chromium is a metallic chemical that originates as a contaminant in the environment from the discharges of dye and paint pigments, wood preservatives, chrome-plating liquid wastes, and leaching from hazardous waste sites. The greatest use of chromium is in metal alloys such as stainless steel; protective coatings on metal; magnetic tapes; and pigments for paints, cement, paper, rubber, composition floor covering, etc. The two largest sources of chromium emission in the atmosphere are from the chemical manufacturing and combustion of natural gas, oil and coal.

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<b>Natural Sources</b>	Chromium is a metal found in natural deposits of ores containing other elements, mostly as chrome-iron ore. It is also widely present in soil and plants. Under most conditions, natural chromium in the environment occurs as Cr <sup>3+</sup> . Under oxidizing conditions, alkaline pH range, presence of MnO <sub>2</sub> , and minerals containing chromium, part of it may occur as hexavalent chromium dissolved in groundwater. Recent sampling of drinking water sources throughout California suggests that hexavalent chromium may occur naturally in groundwater at many locations. Naturally occurring hexavalent chromium may be associated with serpentinite-containing rock or chromium containing geologic formations.
<b>History of Occurrence</b>	Hexavalent chromium has been detected in groundwater at several industrial sites where wood treatment or metal plating solutions were used. Between 1952 and 1966, Pacific Gas & Electric (PG&E) used hexavalent chromium to reduce corrosion in its natural gas compressor plant in Hinkley, near Barstow. Hexavalent chromium contaminated groundwater was suspected of causing cancer and tumors in local residents beginning in the mid 1980's. Since then, elevated levels of hexavalent chromium has been detected in groundwater at several other locations including: Glendale, Topock, and Kettleman City. Hexavalent chromium also occurs naturally in groundwater at the Presidio of San Francisco and Lawrence Livermore National Laboratory.
<b>Transport Characteristics</b>	Hexavalent chromium is readily soluble in water. Under high Eh (oxidizing) and alkaline (pH above 7) conditions, hexavalent chromium can be predominant in groundwater. However, in the presence of organic matter, ferrous iron (Fe II) and sulfide, hexavalent chromium can be readily reduced to Cr <sup>3+</sup> and immobilized. Adsorption of hexavalent chromium by clayey soil and natural aquifer materials is low to moderate under near-neutral pH ranges commonly encountered in groundwater.

## **REMEDICATION & TREATMENT TECHNOLOGIES**

### **In-situ Treatment:**

In several laboratory and field pilot tests, and full-scale remediation systems, hexavalent chromium has been removed using a permeable reactive barrier filled with zero-valent iron granules or surfactant-modified zeolite. Engineered chemical reduction technologies involve the addition or in-situ injection of an electron donor such as hydrogen sulfite, sodium dithionite, sodium metabisulfite, calcium metabisulfite calcium polysulfite or tin(II) chloride. Other methods include geochemical fixation, soil flushing and extraction, bioremediation and electrokinetics.

### **Above-Ground Treatment**

Drinking water can be treated by different pump and treat remediation systems.  $\text{Cr}^3$  and  $\text{Cr}^6$  can be removed by reverse osmosis or ion exchange resin. The ion exchange method should be used with caution, as presence of other metals may interact with the process and decrease system effectiveness. Removal of  $\text{Cr}^6$  by seaweed biosorbent and bacteria (*Bacillus* sp.) within packed bed reactors has also been used.

### **Natural Attenuation**

Natural attenuation of hexavalent chromium may occur in the subsurface environment through reduction by organic matter, iron hydroxides or sulfides. Prior to selection of natural attenuation as an option for remediation, the following conditions need to be demonstrated: 1) there are natural reducers present within the aquifer, 2) the amount of hexavalent chromium and other reactive constituents do not exceed the capacity of the aquifer to reduce them, 3) the rate of hexavalent chromium reduction is greater than the rate of transport of the aqueous hexavalent chromium off the impacted site, 4) the hexavalent chromium remains immobile, and 5) there is no net oxidation of  $\text{Cr}^3$  to  $\text{Cr}^6$ .

**HEALTH EFFECT INFORMATION**

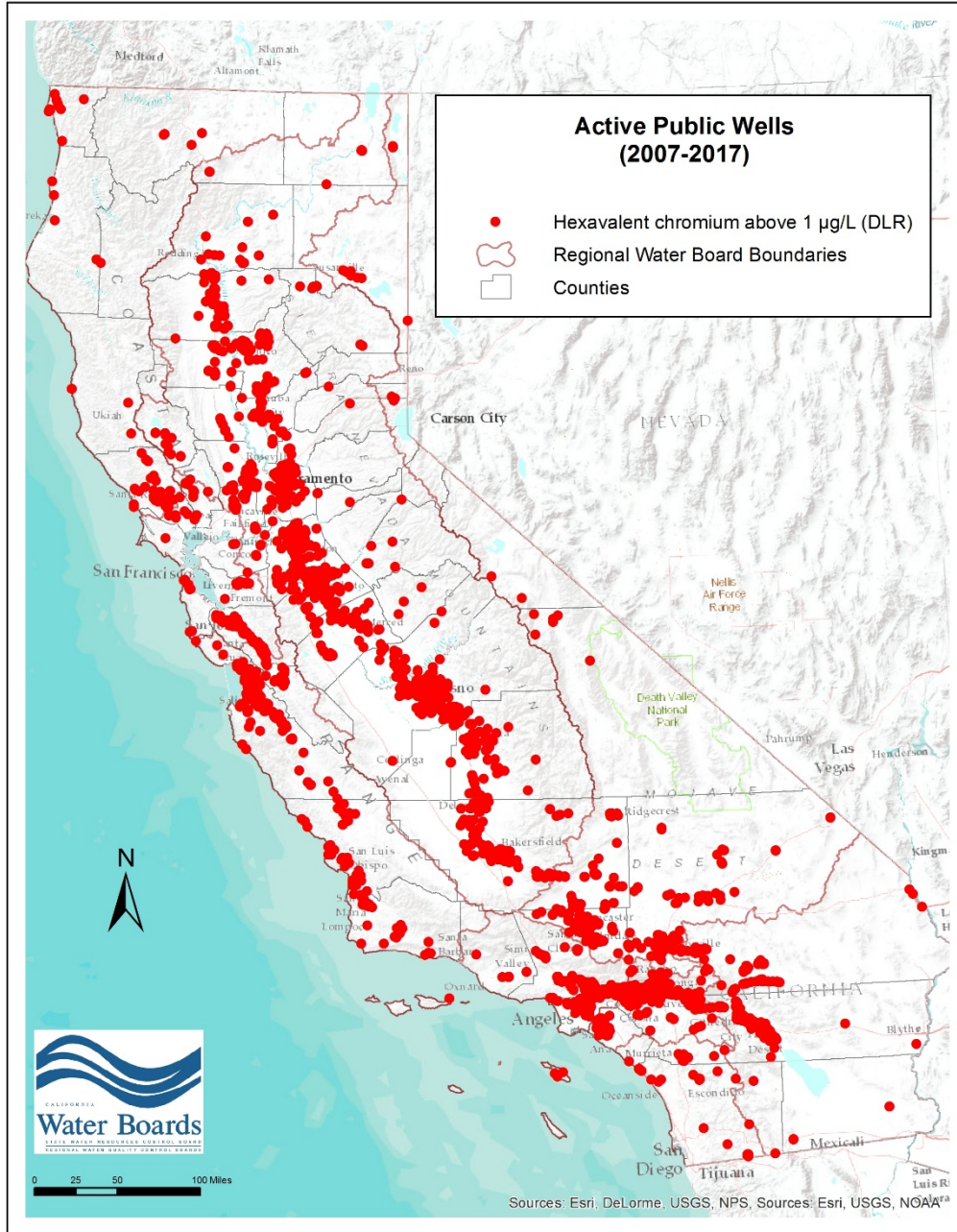
Hexavalent chromium is known to cause cancer in humans when inhaled. It can also damage the lining of the nose and throat and irritate the lungs. A number of scientific studies have found elevated rates of lung cancer in workers with occupational exposure to hexavalent chromium by inhalation. A few studies of workers exposed by inhalation have shown an increase in cancers of the gastrointestinal tract. When swallowed, hexavalent chromium can upset the gastrointestinal tract and damage the liver and kidneys. In recent scientific studies of laboratory animals, hexavalent chromium has been linked to cancer when ingested, although it is rapidly converted to Cr<sup>3</sup> after entering the stomach and coming into contact with organic matter.

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**Active and Standby Public Water Wells with at least one detection of Chromium 6 above 1 µg/L (DLR), 3,798 wells. (Source: Public Well data using GeoTracker GAMA).**