Borax

Borax, also known as **sodium borate**, **sodium tetraborate**, or **disodium tetraborate**, is an important <u>boron compound</u>, a <u>mineral</u>, and a <u>salt of boric acid</u>. Powdered borax is white, consisting of soft colorless crystals that dissolve in water. A number of closely related minerals or chemical compounds that differ in their <u>crystal water</u> content are referred to as *borax*, but the word is usually used to refer to the <u>decahydrate</u>. [1] Commercially sold borax is partially dehydrated.

Borax is a component of many <u>detergents</u>, <u>cosmetics</u>, and <u>enamel</u> glazes. It is used to make <u>buffer solutions</u> in <u>biochemistry</u>, as a <u>fire retardant</u>, as an <u>anti-fungal</u> compound, in the manufacture of <u>fiberglass</u>, as a <u>flux</u> in <u>metallurgy</u>, neutron-capture shields for radioactive sources, a texturing agent in cooking, as a <u>precursor</u> for other boron compounds, and along with its inverse, boric acid, is useful as an insecticide.

In <u>artisanal gold mining</u>, <u>borax</u> is sometimes used as part of a process (as a flux) meant to eliminate the need for toxic mercury in the <u>gold extraction</u> process, although it cannot directly replace mercury. Borax was reportedly used by gold miners in parts of the <u>Philippines</u> in the 1900s.^[6]

Borax was first discovered in dry lake beds in <u>Tibet</u> and was imported via the <u>Silk Road</u> to the Arabian Peninsula in the 8th century AD.^[7] Borax first came into common use in the late 19th century when <u>Francis Marion Smith's Pacific Coast Borax Company</u> began to market and popularize a large variety of applications under the <u>20 Mule Team Borax trademark</u>, named for <u>the method</u> by which borax was originally hauled out of the <u>California and Nevada deserts</u> in large enough quantities to make it cheap and commonly available.^{[8][9]}

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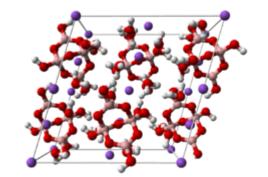
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Names

IUPAC name

Sodium tetraborate decahydrate

Other names

Borax decahydrate

Identifiers			
CAS Number	1303-96-4 (http://www. commonchemistry.org/ ChemicalDetail.aspx?r ef=1303-96-4) (decahydrate) ✓		
3D model (JSmol)	Interactive image (http s://chemapps.stolaf.ed u/jmol/jmol.php?model =%5BNa%2B%5D.%5 BNa%2B%5D.%5BO- %5DB1OB2OB%28% 5BO-%5D%29OB%28		

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Chemistry

The term *borax* is often used for a number of closely related minerals or chemical compounds that differ in their crystal water content:

- anhydrous sodium tetraborate, Na₂B₄O₇
- sodium tetraborate pentahydrate, Na₂B₄O₇·5H₂O
- sodium tetraborate decahydrate, Na₂B₄O₇·10H₂O

Borax is generally described as $Na_2B_4O_7\cdot 10H_2O$. However, it is better formulated as $Na_2[B_4O_5(OH)_4]\cdot 8H_2O$, since borax contains the $[B_4O_5(OH)_4]^{2-}$ ion. In this structure, there are two four-coordinate boron atoms (two BO_4 tetrahedra) and two three-coordinate boron atoms (two BO_3 triangles).

Borax is also easily converted to <u>boric acid</u> and other <u>borates</u>, which have many applications. Its reaction with hydrochloric acid to form boric acid is:

$$Na_2B_4O_7 \cdot 10H_2O$$
 + 2 HCl \rightarrow 4 H_3BO_3 + 2 NaCl + 5 H_2O

The "decahydrate" is sufficiently stable to find use as a primary standard for acid base <u>titrimetry</u>.^[10]

When borax is added to a flame, it produces a <u>yellow green</u> color.^[11] Borax is not used for this purpose in <u>fireworks</u> due to the overwhelming yellow color of sodium. Boric acid is used to color <u>methanol</u> flames a transparent green.

Borax is very soluble in ethylene glycol, moderately soluble in diethylene glycol and methanol, slightly soluble in acetone.^[12] It is poorly soluble in cold water, but its solubility increases significantly with temperature.^{[12][13]}

Etymology

	O1%29O2.O.O.O.O. O.O.O.O.O.O)			
ChEBI	CHEBI:86222 (https://www.ebi.ac.uk/chebi/searchId.do?chebiId=86222) ✗			
ChEMBL	ChEMBL1076681 (htt ps://www.ebi.ac.uk/ch embldb/index.php/com pound/inspect/ChEMB L1076681)			
ChemSpider	17339255 (http://www. chemspider.com/Che mical-Structure.17339 255.html) •			
ECHA InfoCard	100.014.129 (https://e cha.europa.eu/substa nce-information/-/subs tanceinfo/100.014.12 9)			
EC Number	215-540-4			
E number	E285 (preservatives)			
KEGG	D03243 (http://www.ke gg.jp/entry/D03243) *			
PubChem <u>CID</u>	11954323 (https://pub chem.ncbi.nlm.nih.go v/compound/1195432 3)			
UNII	91MBZ8H3QO (http s://fdasis.nlm.nih.gov/ srs/srsdirect.jsp?regno =91MBZ8H3QO)			
InChI				
SMILES	SMILES			
Properties				
Chemical formula	Na ₂ B ₄ O ₇ ·10H ₂ O or Na ₂ [B ₄ O ₅ (OH) ₄]·8H ₂ O			
Molar mass	381.38 (decahydrate) 201.22 (anhydrate)			
Appearance	white solid			
Density	1.73 g/cm ³			

The English word *borax* is Latinized: the <u>Middle English</u> form was *boras*, from <u>Old French</u> *boras*, *bourras*. That may have been from medieval Latin *baurach* (another English spelling), *borac(-/um/em)*, *borax*, along with Spanish *borrax* (> *borraj*) and Italian *borrace*, in the 9th century. Another name for borax is **tincal**, from Sanskrit.^[7]

The word <code>tincal / tinkel/</code> "tinkle", or <code>tincar / tinker/</code> "tinker", refers to crude borax, before it is purified, as mined from lake deposits in Tibet, Persia, and other parts of Asia. The word was adopted in the 17th century from Malay <code>tingkal</code> and from <code>Urdu/Persian/Arabic</code> נוֹל tinkār/tankār; thus the two forms in English. These all appear to be related to the Sanskrit ਫੋਰਿਆ tānkana.^[14]

Natural sources

Borax occurs naturally in <u>evaporite</u> deposits produced by the repeated evaporation of <u>seasonal lakes</u>. The most commercially important deposits are found in: <u>Turkey</u>; <u>Boron</u>, <u>California</u>; and <u>Searles Lake</u>, <u>California</u>. Also, borax has been found at many other locations in the <u>Southwestern United States</u>, the <u>Atacama desert</u> in <u>Chile</u>, newly discovered deposits in <u>Bolivia</u>, and in <u>Tibet</u> and <u>Romania</u>. Borax can also be produced synthetically from other boron compounds.

Naturally occurring borax (known by the trade name Rasorite–46 in the United States and many other countries) is refined by a process of recrystallization.^[15]

Uses

Household products

Borax is used in various household laundry and cleaning products,^[16] including the "20 Mule Team Borax" laundry booster, "Boraxo" powdered hand soap, and some tooth bleaching formulas.^[17]

pH buffer

Borate ions (commonly supplied as boric acid) are used in biochemical and chemical laboratories to make buffers, e.g. for polyacrylamide gel electrophoresis of DNA and RNA, such as TBE buffer (borate buffered tris-hydroxymethylaminomethonium)^{[18][19][20]} or the newer SB buffer or BBS buffer (borate buffered saline) in coating procedures. Borate buffers (usually at pH 8) are also used as preferential equilibration solution in dimethyl pimelimidate (DMP) based crosslinking reactions.

Co-complexing agent

	(decahydrate, solid) ^[1] 2.4 g/cm ³ (anhydrate, solid) ^[1]		
Melting point	743 °C (1,369 °F; 1,016 K) (anhydrate) ^[1] 75 °C (decahydrate, decomposes) ^[1]		
Boiling point	1,575 °C (2,867 °F; 1,848 K) (anhydrate) ^{[1}		
Magnetic susceptibility (χ)	-85.0·10 ⁻⁶ cm ³ /mol		
St	ructure		
Crystal structure	Monoclinic Prismatic		
Space group	C2/c		
Point group	2/m		
Phar	macology		
ATC code	S01AX07 (WHO (http s://www.whocc.no/atc _ddd_index/?code=S0 1AX07))		
H	azards		
GHS pictograms	\$		
GHS hazard statements	<u>H360</u>		
GHS precautionary statements	P201, P308+313		
NFPA 704	100		
US health exposure	e limits (NIOSH):		
PEL (Permissible)	none ^[2]		
REL (Recommended)	TWA 1 mg/m ³ (anhydrate and pentahydrate) ^{[2][3]} TWA 5 mg/m ³ (decahydrate) ^[4]		
IDLH (Immediate danger)	N.D. ^[2]		

Borax as a source of <u>borate</u> has been used to take advantage of the cocomplexing ability of borate with other agents in water to form complex ions with various substances. Borate and a suitable <u>polymer</u> bed are used to <u>chromatograph</u> non-glycosylated <u>hemoglobin</u> differentially from glycosylated hemoglobin (chiefly <u>HbA1c</u>), which is an indicator of long term <u>hyperglycemia</u> in diabetes mellitus.

Water-softening agent

Borax alone does not have a high affinity for the <u>hardness cations</u>, although it has been used for water-softening. Its chemical equation for *water-softening* is given below:

Ca ²⁺	(aq) +	Na ₂ B ₄ O ₇	$(aq) \rightarrow$	CaB ₄ O ₇	(s)↓ + 2	<u>Na</u> ⁺	(aq)
Mg ²⁺	(aq) +	Na ₂ B ₄ O ₇	$(aq) \rightarrow$	MgB ₄ O ₇	$(s)^{\downarrow} + 2$	Na ⁺	(aq)

The <u>sodium</u> ions introduced do not make water 'hard'. This method is suitable for removing both temporary and permanent types of hardness.

Flux

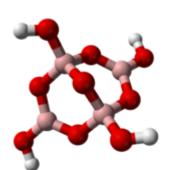
A mixture of borax and <u>ammonium chloride</u> is used as a <u>flux</u> when <u>welding iron</u> and <u>steel</u>. It lowers the melting point of the unwanted <u>iron oxide</u> (<u>scale</u>), allowing it to run off. Borax is also used mixed with water as a flux when <u>soldering jewelry metals</u> such as <u>gold</u> or <u>silver</u>, where it allows the molten <u>solder</u> to <u>wet</u> the metal and flow evenly into the joint. Borax is also a good flux for "pre-tinning" <u>tungsten</u> with zinc — making the tungsten soft-solderable. [21] Borax is often used as a flux for forge welding.

Small-scale gold mining

Borax is replacing mercury as the preferred method for extracting gold in small-scale mining facilities. The method is called the borax method and is used in the Philippines.^[22]

Related compounds			
Other anions	Sodium aluminate; sodium gallate		
Other cations Potassium tetraborate			
Related compounds	Boric acid, sodium perborate		
Except where otherwise noted, data are given for materials in their standard state (at 25 °C [77 °F], 100 kPa).			
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Infobox references



The structure of the anion $[B_4O_5(OH)_4]^{2-}$ in borax



Borax "cottonball"



Borax-based laundry detergent

Flubber

A rubbery polymer sometimes called Slime, Flubber, 'gluep' or 'glurch' (or erroneously called Silly Putty, which is based on silicone polymers), can be made by cross-linking polyvinyl alcohol with borax. Making flubber from polyvinyl acetate-based glues, such as Elmer's Glue, and borax is a common elementary-science demonstration. [23][24]



Traction steam engine hauling borax, Death Valley National Park, California, 1904

Food additive

Borax, given the <u>E number</u> E285, is used as a <u>food additive</u> in some countries, but is banned in some countries, like the U.S., and <u>Thailand</u>. As a consequence, certain foods, such as <u>caviar</u>, produced for sale in the US contain higher levels of <u>salt</u> to assist preservation. [25] Its use as a cooking ingredient is to add a firm rubbery texture to the food, or as a preservative. In Asian cooking it is mostly used for its texturing properties. In <u>China</u>, borax (<u>Chinese</u>: $\mathfrak{M}\mathfrak{P}$; <u>pinyin</u>: $p\acute{e}ng$ $sh\bar{a}$ or <u>Chinese</u>: $\mathfrak{H}\mathfrak{T}$; <u>pinyin</u>: $yu\grave{e}$ $sh\acute{i}$) was found to have been added to some Chinese foods like hand-pulled noodles <u>lamian</u> and some <u>rice noodles</u> like <u>shahe fen</u>, <u>kway teow</u>, and <u>chee cheong fun</u> recipes. [26] In <u>Indonesia</u> it is a common, but forbidden, additive to such foods as noodles, bakso (meatballs),



Old steam tractor and borax wagons, Death Valley National Park

and steamed rice. The country's Directorate of Consumer Protection warns of the risk of $\underline{liver\ cancer}$ with high consumption over a period of 5–10 years.^[27]

Other uses

- Ingredient in enamel glazes
- Component of glass, pottery, and ceramics
- Used as an additive in ceramic slips and <u>glazes</u> to improve fit on wet, greenware, and bisque
- Fire retardant
- Anti-fungal compound for cellulose insulation
- Moth proofing 10% solution for wool^[28]
- Pulverized for the prevention of stubborn pests (e.g. <u>German cockroaches</u>) in closets, pipe and cable inlets, wall <u>panelling</u> gaps, and inaccessible locations where ordinary pesticides are undesirable
- Anti-fungal foot soak
- Precursor for sodium perborate monohydrate that is used in detergents, as well as for boric acid and other borates
- Tackifier ingredient in casein, starch and dextrin based adhesives
- Precursor for boric acid, a tackifier ingredient in polyvinyl acetate, polyvinyl alcohol based adhesives
- Fluoride detoxification
- Treatment for thrush in horses' hooves
- To make indelible ink for dip pens by dissolving shellac into heated borax
- Curing agent for snake skins
- Curing agent for salmon eggs, for use in sport fishing for salmon
- Swimming pool buffering agent to control pH
- Neutron absorber, used in <u>nuclear reactors</u> and spent fuel pools to control reactivity and to shut down a nuclear chain reaction
- As a micronutrient fertilizer to correct boron-deficient soils. [29][30]
- Preservative in taxidermy
- To color fires with a green tint^[31]
- Was traditionally used to coat dry-cured meats such as hams to protect them from becoming <u>fly-blown</u> during further storage
- Is found in some commercial vitamin supplements
- For stopping car radiator and engine block leaks^[32]
- Used by blacksmiths in forge welding
- Used as a woodworm treatment (diluted in water)
- Used in the treatment or prevention of wood rot in classic wood boats
- Used to enrich beets with Boron when leaves are wilting



Rio Tinto Borax Mine Pit, Boron, California

Toxicity

Borax, sodium tetraborate decahydrate, according to one study, is not acutely toxic. [33] Its \underline{LD}_{50} (median lethal dose) score is tested at 2.66 g/kg in rats, [34] meaning that a significant dose of the chemical is needed to cause severe symptoms or death. The lethal dose is not necessarily the same for humans. On pesticide information websites it is listed as a non-lethal compound and of no hazardous concerns. [35]

Borax has been in use as an insecticide in the United States with various restrictions since 1946. All restrictions were removed in February 1986 due to the low toxicity of borax, as reported in two EPA documents relating to boric acid and borax. [36][37]

EPA has determined that, because they are of low toxicity and occur naturally, boric acid and its sodium salts should be exempted from the requirement of a tolerance (maximum residue limit) for all raw agricultural commodities.^[36]

Although it cited inconclusive data, a re-evaluation in 2006 by the EPA still found that "There were no signs of toxicity observed during the study and no evidence of cytotoxicity to the target organ." In the reevaluation, a study of toxicity due to overexposure was checked and the findings were that "The residential handler inhalation risks due to boric acid and its sodium salts as active ingredients are not a risk concern and do not exceed the level of concern..." but that there could be some risk of irritation to children inhaling it if used as a powder for cleaning rugs.

Sodium tetraborate decahydrate has no known hazard issues.^[39]

Conditions defined as "over-exposure" to borax dust can cause respiratory irritation, while no skin irritation is known to exist due to borax. Ingestion may cause gastrointestinal distress including <u>nausea</u>, persistent vomiting, abdominal pain, and <u>diarrhea</u>. Effects on the vascular system and <u>human brain</u> include headaches and lethargy, but are less frequent. "In severe poisonings, a beefy red skin rash affecting palms, soles, buttocks and scrotum has been described. With severe poisoning, erythematous and exfoliative rash, unconsciousness, respiratory depression, and renal failure."^[33]

A draft risk assessment released by <u>Health Canada</u> in July 2016 has found that overexposure to boric acid has the potential to cause developmental and reproductive health effects. Since people are already exposed to boric acid naturally through their diets and water, Health Canada advised that exposure from other sources should be reduced as much as possible, especially for children and pregnant women. The concern is not with any one product, but rather multiple exposures from a variety of sources. With this in mind, the department also announced that registrations for certain pesticides that contain boric acid, which are commonly used in homes, will have their registrations cancelled and be phased out of the marketplace. As well, new, more protective label directions are being introduced for other boric acid pesticides that continue to be registered in Canada (for example, enclosed bait stations and spot treatments using gel formulations). [40]

Possible carcinogen

The Indonesian Directorate of Consumer Protection warns of the risk of <u>liver cancer</u> with high consumption of borax over a period of 5–10 years.^[27]

Risk to fertility and pregnancy

Borax was added to the <u>Substance of Very High Concern (SVHC)</u> candidate list on 16 December 2010. The SVHC candidate list is part of the <u>EU</u> Regulations on the <u>Registration</u>, <u>Evaluation</u>, <u>Authorisation</u> and <u>Restriction</u> of Chemicals 2006 (REACH), and the addition was based on the revised classification of borax as *toxic for reproduction* category 1B under the CLP Regulations. Substances and mixtures imported into the EU which contain borax are now required to be

labelled with the warnings "May damage fertility" and "May damage the unborn child".^[41] It was proposed for addition to REACH Annex XIV by the ECHA on 1 July 2015.^[42] If this recommendation is approved, all imports and uses of borax in the EU will have to be authorized by the ECHA. Review of the boron toxicity (as boric acid and borates) published 2012 in Journal of Toxicology and Environmental Health concluded: "It clearly appears that human B [boron] exposures, even in the highest exposed cohorts, are too low to reach the blood (and target tissue) concentrations that would be required to exert adverse effects on reproductive functions."^[43]

See also

- Borax bead test
- John Veatch
- List of cleaning agents

- Sodium borohydride
- Ulexite

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External links

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- International Chemical Safety Card 1229 (http://www.inchem.org/documents/icsc/icsc/eics1229.htm) (fused borax)
- National Pollutant Inventory Boron and compounds (https://web.archive.org/web/20060209040519/http://www.npi.g ov.au/database/substance-info/profiles/15.html)
- NIOSH Pocket Guide to Chemical Hazards (https://www.cdc.gov/niosh/npg/npgd0058.html)
- Sodium Borate in sefsc.noaa.gov (https://web.archive.org/web/20020821230321/http://www.sefsc.noaa.gov/HTMLdocs/SodiumBorate.htm)

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