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Boric acid

Boric acid, also called hydrogen borate, boracic acid, orthoboric acid and acidum boricum, is a weak, monobasic Lewis acid of boron, which is often used as an antiseptic, insecticide, flame retardant, neutron absorber, or precursor to other chemical compounds. It has the chemical formula $\underline{H_3BO_3}$ (sometimes written $B(OH)_3$), and exists in the form of colorless crystals or a white powder that dissolves in water. When occurring as a mineral, it is called sassolite.

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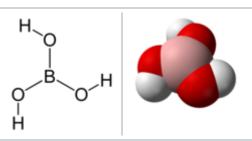
Further reading

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Occurrence

Boric acid, or <u>sassolite</u>, is found mainly in its free state in some volcanic districts, for example, in the Italian region of <u>Tuscany</u>, the <u>Lipari Islands</u> and the US state of <u>Nevada</u>. In these volcanic settings it issues, mixed with steam, from fissures in the ground. It is also found as a constituent of many naturally occurring minerals – <u>borax</u>, <u>boracite</u>, <u>ulexite</u> (boronatrocalcite) and <u>colemanite</u>. Boric acid and its salts are found in seawater. It is also found in plants, including almost all fruits.^[2]

Boric acid





Names

IUPAC names

Boric acid

Trihydrooxidoboron

Other names

Orthoboric acid,

Boracic acid,

Sassolite,

Optibor,

Borofax,

Trihydroxyborane,

Boron(III) hydroxide,

Boron Trihvdroxide

Boron minyaroxiao	
Identifiers	
CAS Number	10043-35-3 (http://w ww.commonchemistr y.org/ChemicalDetail. aspx?ref=10043-35- 3) ✓
3D model (JSmol)	Interactive image (htt ps://chemapps.stolaf. edu/jmol/jmol.php?m odel=OB%28O%29

Boric acid was first prepared by Wilhelm Homberg (1652–1715) from borax, by the action of mineral acids, and was given the name *sal sedativum Hombergi* ("sedative salt of Homberg"). However borates, including boric acid, have been used since the time of the ancient Greeks for cleaning, preserving food, and other activities.

Preparation

Boric acid may be prepared by reacting <u>borax</u> (sodium tetraborate decahydrate) with a mineral acid, such as hydrochloric acid:

$$Na_2B_4O_7 \cdot 10H_2O + 2 \ HCI \rightarrow 4 \ B(OH)_3 \ [or \ H_3BO_3] + 2 \ NaCI + 5 \ H_2O$$

It is also formed as a by product of hydrolysis of boron trihalides and diborane:^[3]

$$B_2H_6 + 6 H_2O \rightarrow 2 B(OH)_3 + 6 H_2$$

$$BX_3 + 3 H_2O \rightarrow B(OH)_3 + 3 HX (X = CI, Br, I)$$

Properties

Boric acid is soluble in boiling water. When heated above 170 $^{\circ}$ C, it dehydrates, forming metaboric acid (HBO₂):

$$H_3BO_3 \rightarrow HBO_2 + H_2O$$

<u>Metaboric acid</u> is a white, cubic crystalline solid and is only slightly soluble in water. Metaboric acid melts at about 236 °C, and when heated above about 300 °C further dehydrates, forming <u>tetraboric acid</u>, also called pyroboric acid $(H_2B_4O_7)$:

$$4 \text{ HBO}_2 \rightarrow \text{H}_2\text{B}_4\text{O}_7 + \underline{\text{H}_2\text{O}}$$

The term *boric acid* may sometimes refer to any of these compounds. Further heating (to about 330 °C)^[4] leads to boron trioxide.

$$H_2B_4O_7 \rightarrow 2 B_2O_3 + \underline{H_2O}$$

There are conflicting interpretations for the origin of the acidity of aqueous boric acid solutions. Raman spectroscopy of strongly alkaline solutions has shown the presence of $B(OH)_4^-$ ion,^[5] leading some to conclude that the acidity is exclusively due to the abstraction of OH^- from water:^{[5][6][7][8]}

$$B(OH)_3 + \underline{H_2O} \stackrel{\rightharpoonup}{\longleftarrow} B(OH)_4^- + \underline{H^+} (K = 7.3 \times 10^{-10}; pK = 9.14)$$

or more properly expressed in the aqueous solution:

	Interactive image (htt ps://chemapps.stolaf. edu/jmol/jmol.php?m odel=%5BOH%2B% 5D%3D%5BB-%5 D%28O%29O)	
ChEBI	CHEBI:33118 (http s://www.ebi.ac.uk/ch ebi/searchId.do?che bild=33118) ✓	
ChEMBL	ChEMBL42403 (http s://www.ebi.ac.uk/ch embldb/index.php/co mpound/inspect/ChE MBL42403) ✓	
ChemSpider	7346 (http://www.che mspider.com/Chemic al-Structure.7346.ht ml) ✓	
ECHA InfoCard	100.030.114 (https://echa.europa.eu/substance-information/-/substanceinfo/100.030.114)	
EC Number	233-139-2	
E number	E284 (preservatives)	
KEGG	D01089 (http://www.k egg.jp/entry/D01089)	
PubChem <u>CID</u>	7628 (https://pubche m.ncbi.nlm.nih.gov/c ompound/7628)	
UNII	R57ZHV85D4 (http s://fdasis.nlm.nih.go v/srs/srsdirect.jsp?re gno=R57ZHV85D4)	
InChl		
SMILES		
Properties		
Chemical formula	BH ₃ O ₃	

$$B(OH)_3 + 2 H_2O \stackrel{\rightharpoonup}{\longleftarrow} B(OH)_4^- + H_3O^+$$

This may be characterized^{[6][7][8]} as <u>Lewis acidity</u> of boron toward OH⁻, rather than as Brønsted acidity.

<u>Polyborate</u> anions are formed at <u>pH</u> 7–10 if the boron concentration is higher than about 0.025 mol/L. The best known of these is the 'tetraborate' ion, found in the mineral borax:

$$4[B(OH)_4]^- + 2H^+ \rightleftharpoons [B_4O_5(OH)_4]^{2-} + 7H_2O$$

Boric acid makes an important contribution to the absorption of low frequency sound in seawater.^[9]

Reactions

With <u>polyols</u> such as <u>glycerol</u> and <u>mannitol</u> the acidity of the solution is increased. With <u>mannitol</u> for example the pK decreases to 5.15. This is due to the formation of a chelate, $[((OH)_4C_6H_8O_2)_2B]^-$, and this feature is used in analytical chemistry.^[10]

Boric acid dissolves in anhydrous sulfuric acid:[10]

$$B(OH)_3 + 6H_2SO_4 \rightarrow 3H_3O^+ + 2HSO_4^- + B(HSO_4)_4^-$$

Boric acid reacts with alcohols to form <u>borate esters</u>, $B(OR)_3$ where R is <u>alkyl</u> or <u>aryl</u>. A dehydrating agent, such as concentrated <u>sulfuric acid</u> is typically added:^[11]

$$B(OH)_3 + 3 ROH \rightarrow B(OR)_3 + 3 H_2O$$

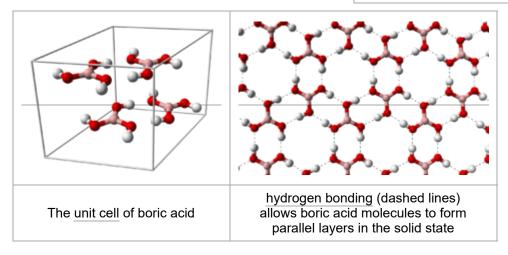
Molecular and crystal structure

The three oxygen atoms form a <u>trigonal planar geometry</u> around the boron. The B-O bond length is 136 pm and the O-H is 97 pm. The <u>molecular point</u> group is C_{3h} .

Crystalline boric acid consists of layers of $B(OH)_3$ molecules held together by hydrogen bonds of length 272 pm. The distance between two adjacent layers is 318 pm.^[10]

Molar mass	61.83 g⋅mol ⁻¹	
Appearance	White crystalline solid	
Density	1.435 g/cm ³	
Melting point	170.9 °C (339.6 °F; 444.0 K)	
Boiling point	300 °C (572 °F; 573 K)	
Solubility in water	2.52 g/100 mL (0 °C) 4.72 g/100 mL (20 °C) 5.7 g/100 mL (25 °C) 19.10 g/100 mL (80 °C) 27.53 g/100 mL (100 °C)	
Solubility in other solvents	Soluble in lower alcohols moderately soluble in pyridine very slightly soluble in acetone	
log P	-0.29 ^[1]	
Acidity (pK _a)	9.24, 12.4, 13.3	
Conjugate base	Borate	
Magnetic susceptibility (χ)	-34.1·10 ⁻⁶ cm ³ /mol	
St	ructure	
Molecular shape	Trigonal planar	
Dipole moment	Zero	
Pharmacology		
ATC code	S02AA03 (WHO (htt ps://www.whocc.no/a tc_ddd_index/?code =S02AA03)) D08AD (WHO (https://www.w hocc.no/atc_ddd_ind ex/?code=D08AD))	
Hazards		
Safety data sheet EU classification	See: data page	
(DSD) (outdated)		

	. <u></u>	
	Xn	
R-phrases	R60 R61	
(outdated)		
S-phrases	S53 S45	
(outdated)		
NFPA 704	100	
Flash point	Non-flammable	
Lethal dose or concentration (LD, LC):		
LD ₅₀ (median dose)	2660 mg/kg, oral (rat)	
Related compounds		
Related compounds	Boron trioxide Borax	
Supplementary data page		
Structure and properties	Refractive index (n) , Dielectric constant (ε_r) , etc.	
Thermodynamic data	Phase behaviour solid–liquid–gas	
Spectral data	UV, IR, NMR, MS	
Except where otherwise noted, data are given for materials in their standard state (at 25 °C [77 °F], 100 kPa).		
✓ verify	(what is ✓ ?)	
Infobox references		



Toxicology

Based on mammalian median lethal dose (LD₅₀) rating of 2,660 mg/kg body mass, boric acid is only poisonous if taken internally or inhaled in large quantities. The Fourteenth Edition of the $\underline{Merck\ Index}$ indicates that the LD₅₀ of boric acid is 5.14 g/kg for oral dosages given to rats, and that 5 to 20 g/kg has produced death in adult humans. For comparison's sake, the LD₅₀ of \underline{salt} is reported to be 3.75 g/kg in rats according to the $\underline{Merck\ Index}$. According to the $\underline{Agency\ for\ Toxic}$ Substances and Disease Registry, "The minimal lethal dose of ingested boron (as boric acid) was reported to be 2–3 g in infants, 5–6 g in children, and 15–20 g in adults. [...] However, a review of 784 human poisonings with boric acid (10–88 g) reported no fatalities, with 88% of cases being asymptomatic." [12]

Long-term exposure to boric acid may be of more concern, causing kidney damage and eventually kidney failure (see links below). Although it does not appear to be <u>carcinogenic</u>, studies in dogs have reported <u>testicular atrophy</u> after exposure to 32 mg/kg bw/day for 90 days. This level is far lower than the LD₅₀.^[13]

According to the CLH report for boric acid published by the Bureau for Chemical Substances Lodz, Poland, boric acid in high doses shows significant developmental toxicity and <u>teratogenicity</u> in rabbit, rat, and mouse fetuses as well as cardiovascular defects, skeletal variations, and mild kidney lesions. [14] As a consequence in the 30th ATP to EU directive 67/548/EEC of August 2008, the European Commission decided to amend its classification as <u>reprotoxic</u> category 2, and to apply the risk phrases R60 (may impair fertility) and R61 (may cause harm to the unborn child). [15][16][17][18][19]

At a 2010 European Diagnostics Manufacturing Association (EDMA) Meeting, several new additions to the Substance of Very High Concern (SVHC) candidate list in relation to the Registration, Evaluation, Authorisation and Restriction of Chemicals Regulations 2007 (REACH) were discussed. Following the registration and review completed as part of REACH, the classification of Boric Acid CAS 10043-35-3 / 11113-50-1 is listed from 1 December 2010 is H360FD (May damage fertility. May damage the unborn child.)^{[20][21]}

Uses

Industrial

The primary industrial use of boric acid is in the manufacture of monofilament <u>fiberglass</u> usually referred to as textile fiberglass. Textile fiberglass is used to reinforce plastics in applications that range from boats, to industrial piping to computer circuit boards.^[22]

In the jewelry industry, boric acid is often used in combination with <u>denatured alcohol</u> to reduce surface <u>oxidation</u> and firescale from forming on metals during annealing and soldering operations.

Boric acid is used in the production of the glass in LCD flat panel displays.

In electroplating, boric acid is used as part of some proprietary formulas. One such known formula calls for about a 1 to 10 ratio of H_3BO_3 to $NiSO_4$, a very small portion of sodium lauryl sulfate and a small portion of H_2SO_4 .

Boric acid, mixed with borax (sodium tetraborate decahydrate) at the weight ratio of 4:5, is highly soluble in water, though they are not so soluble separately.^[23] The solution is used for fire retarding agent of wood by impregnation.^[24]

It is also used in the manufacturing of ramming mass, a fine <u>silica</u>-containing powder used for producing <u>induction</u> furnace linings and ceramics.

Boric acid is one of the most commonly used substances that can neutralize active <u>hydrofluoric acid</u> (HF). It works by forcing the free F⁻ anions into complex salts. This process defeats the extreme toxicity of hydrofluoric acid, particularly its ability to sequester ionic calcium from blood serum which can lead to cardiac arrest and bone decomposition; such an

event can occur from just minor skin contact with HF.[25]

Boric acid is added to borax for use as welding flux by blacksmiths.^[26]

Boric acid, in combination with silicone oil, is used to manufacture Silly Putty.^[27]

Boric acid may be used in Marcellus Shale hydraulic fracturing in Pennsylvania. [28]

Medical

Boric acid can be used as an <u>antiseptic</u> for minor burns or cuts and is sometimes used in salves and <u>dressings</u>, such as <u>boracic lint</u>. Boric acid is applied in a very dilute solution as an eye wash. Dilute boric acid can be used as a vaginal douche to treat <u>bacterial vaginosis</u> due to excessive alkalinity,^[29] as well as <u>candidiasis</u> due to non-albicans candida.^[30] As an <u>antibacterial</u> compound, boric acid can also be used as an <u>acne</u> treatment. It is also used as prevention of <u>athlete's foot</u>, by inserting powder in the socks or stockings, and in alcohol solution can be used to treat some kinds of <u>otitis externa</u> (ear infection) in both humans and animals. The preservative in urine sample bottles in the UK is boric acid.

Boric acid solutions used as an eye wash or on abraded skin are known to be toxic, particularly to infants, especially after repeated use; this is because of its slow elimination rate.^[31]

Insecticidal

Boric acid was first registered in the US as an insecticide in 1948 for control of <u>cockroaches</u>, <u>termites</u>, <u>fire ants</u>, <u>fleas</u>, <u>silverfish</u>, and many other <u>insects</u>. The product is generally considered to be safe to use in household kitchens to control cockroaches and ants.^[32] It acts as a stomach poison affecting the insects' <u>metabolism</u>, and the dry powder is <u>abrasive</u> to the insects' <u>exoskeletons</u>. Boric acid also has the reputation as "the gift that keeps on killing" in that roaches that cross over lightly dusted areas do not die immediately, but that the effect is like shards of glass cutting them apart. This often allows a roach to go back to the nest where it soon dies. Cockroaches, being <u>cannibalistic</u>, eat others killed by contact or consumption of boric acid, consuming the powder trapped in the dead roach and killing them, too.

Preservation

In combination with its use as an insecticide, boric acid also prevents and destroys existing wet and dry rot in timbers. It can be used in combination with an <u>ethylene glycol</u> carrier to treat external wood against fungal and insect attack. It is possible to buy borate-impregnated rods for insertion into wood via drill holes where dampness and moisture is known to collect and sit. It is available in a gel form and injectable paste form for treating rot affected wood without the need to replace the timber. Concentrates of borate-based treatments can be used to prevent slime, mycelium, and algae growth, even in marine environments.

Boric acid is added to salt in the curing of cattle hides, <u>calfskins</u>, and <u>sheepskins</u>. This helps to control bacterial development, and helps to control insects.

pH buffer

Boric acid in equilibrium with its conjugate base the borate ion is widely used (in the concentration range 50 - 100 ppm boron equivalents) as a primary or adjunct pH buffer system in swimming pools. Boric acid is a weak acid, with pKa (the pH at which buffering is strongest because the free acid and borate ion are in equal concentrations) of 9.24 in pure water at 25 °C. But apparent pKa is substantially lower in swimming pool or ocean waters because of interactions with various

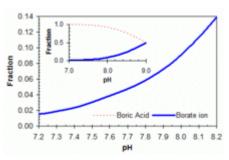
12/25/2018 Boric acid - Wikipedia

other molecules in solution. It will be around 9.0 in a salt-water pool. No matter which form of soluble boron is added, within the acceptable range of pH and boron concentration for swimming pools, boric acid is the predominant form in aqueous solution, as shown in the accompanying figure. The boric acid - borate system can be useful as a primary buffer system (substituting for the bicarbonate system with pKa1 = 6.0 and pKa2 = 9.4 under typical salt-water pool conditions) in pools with salt-water chlorine generators that tend to show upward drift in pH from a working range of pH 7.5 - 8.2. Buffer capacity is greater against rising pH (towards the pKa around 9.0), as illustrated in the accompanying graph. The use of boric acid in this concentration range does not allow any reduction in free HOCl concentration needed for pool sanitation, but it may add marginally to the photo-protective effects of cyanuric acid and confer other benefits through anti-corrosive activity or perceived water softness, depending on overall pool solute composition. [33]

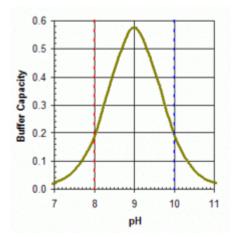
Lubrication

Colloidal suspensions of nanoparticles of boric acid dissolved in petroleum or vegetable oil can form a remarkable lubricant on ceramic or metal surfaces^[34] with a coefficient of sliding friction that decreases with increasing pressure to a value ranging from 0.10 to 0.02. Self-lubricating H_3BO_3 films result from a spontaneous chemical reaction between water molecules and B_2O_3 coatings in a humid environment. In bulk-scale, an inverse relationship exists between friction coefficient and Hertzian contact pressure induced by applied load.

Boric acid is used to lubricate <u>carrom</u> and <u>novuss</u> boards, allowing for faster play.^[35]



Boric acid predominates in solution below pH 9



Boric acid buffers against rising pH in swimming pools

Nuclear power

Boric acid is used in some nuclear power plants as a <u>neutron poison</u>. The boron in boric acid reduces the probability of thermal fission by absorbing some thermal neutrons. Fission chain reactions are generally driven by the probability that free neutrons will result in fission and is determined by the material and geometric properties of the reactor. Natural boron consists of approximately 20% boron-10 and 80% boron-11 isotopes. Boron-10 has a high cross-section for absorption of low energy (thermal) neutrons. By increasing boric acid concentration in the reactor coolant, the probability that a neutron will cause fission is reduced. Changes in boric acid concentration can effectively regulate the rate of fission taking place in the reactor. Boric acid is used only in pressurized water reactors (PWRs) whereas boiling water reactors (BWRs) employ control rod pattern and coolant flow for power control. BWRs use an aqueous solution of boric acid and borax or Sodium Pentaborate for an emergency shut down system. Boric acid may be dissolved in spent fuel pools used to store spent fuel elements. The concentration is high enough to keep neutron multiplication at a minimum. Boric acid was dumped over Reactor 4 of the <u>Chernobyl</u> Nuclear Power Plant after its meltdown to prevent another reaction from occurring.

Pyrotechnics

Boron is used in <u>pyrotechnics</u> to prevent the <u>amide</u>-forming reaction between <u>aluminum</u> and <u>nitrates</u>. A small amount of boric acid is added to the composition to neutralize alkaline amides that can react with the aluminum.

Boric acid can be used as a colorant to make fire green. For example, when dissolved in <u>methanol</u> it is popularly used by fire jugglers and fire spinners to create a deep green flame.^[36]

Agriculture

Boric acid is used to treat or prevent <u>boron deficiencies</u> in plants. It is also used in preservation of grains such as rice and wheat.

Recreational

For the game of <u>carrom</u>, boric acid is sprinkled onto the board of play to act as a <u>dry lubricant</u> to increase the speed of the pieces when struck.^[37]

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External links

- Media related to Boric acid at Wikimedia Commons
- Boric Acid Technical Fact Sheet National Pesticide Information Center (http://npic.orst.edu/factsheets/borictech.pdf)
- Boric Acid General Fact Sheet National Pesticide Information Center (http://npic.orst.edu/factsheets/boricgen.pdf)
- International Chemical Safety Card 0991 (http://www.inchem.org/documents/icsc/icsc/eics0991.htm)
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- ChemSub Online: Boric acid (http://chemsub.online.fr/name/Boric acid.html)

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