

Diborane

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Diborane is the chemical compound consisting of boron and hydrogen with the formula B₂H₆. It is a colorless gas at room temperature with a repulsively sweet odor. Diborane mixes well with air, easily forming explosive mixtures. Diborane will ignite spontaneously in moist air at room temperature. Synonyms include boroethane, boron hydride, and diboron hexahydride.

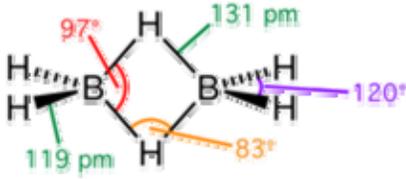
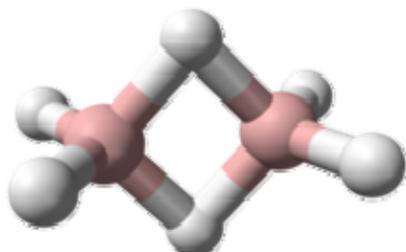
Diborane is a key boron compound with a variety of applications. The compound is classified as "endothermic," meaning that its heat of formation, ΔH[°]_f is positive (36 kJ/mol). Despite its thermodynamic instability, diborane is kinetically robust and exhibits an extensive chemistry, much of it entailing loss of hydrogen.

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Structure and bonding

Diborane adopts a D_{2h} structure containing four terminal and two bridging hydrogen atoms. The model determined by molecular orbital theory indicates that the bonds between boron and the terminal hydrogen atoms are conventional 2-center, 2-electron covalent bonds. The bonding between the boron atoms and the bridging hydrogen atoms is, however, different from that in molecules such as hydrocarbons. Having used two electrons in bonding

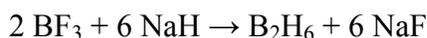
Diborane	
	
	
IUPAC name	
Diborane(6)	
Identifiers	
CAS number	19287-45-7 ✔
PubChem	29529
ChemSpider	27451
RTECS number	HQ9275000
SMILES	BB
InChI	1/B2H4/c1-2/h1-2H2
InChI key	Q SJRRLWJRLPVID-UHFFFAOYAD
Properties	
Molecular formula	B ₂ H ₆
Molar mass	27.67 g/mol
Appearance	colorless gas
Density	1.18 g/L (15 °C) 1.216 g/L (25 °C)
Melting point	−165.5 °C, 108 K, −266 °F
Boiling point	−92.4 °C, 181 K, −134 °F
Solubility in water	Reacts
Solubility	soluble in CS ₂ , diglyme and ethyl acetate ^[1]

to the terminal hydrogen atoms, each boron has one valence electron remaining for additional bonding. The bridging hydrogen atoms provide one electron each. Thus the B₂H₂ ring is held together with four electrons, an example of 3-center-2-electron bonding. This type of bond is sometimes called a 'banana bond'. The lengths of the B-H_{bridge} bonds and the B-H_{terminal} bonds are 1.33 and 1.19 Å respectively, and this difference in the lengths of these bonds reflects the difference in their strengths, the B-H_{bridge} bonds being relatively weaker. The structure is isoelectronic with C₂H₆²⁺, which would arise from the diprotonation of the planar molecule ethene.^[2] Diborane is one of many compounds with such unusual bonding.^[3]

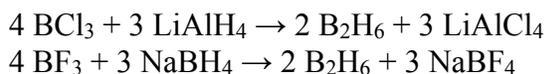
Of the other elements in Group 13, gallium is known to form a similar compound, digallane, Ga₂H₆. Aluminium forms a polymeric hydride, (AlH₃)_n, although unstable Al₂H₆ has been isolated in solid hydrogen and is isostructural with diborane.^[4] No hydrides of indium and thallium have yet been found.^[5]

Production and synthesis

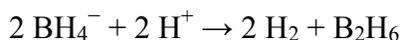
Diborane is so central and has been studied so often that many syntheses exist. Most preparations entail reactions of hydride donors with boron halides or alkoxides. The industrial synthesis involves the reduction of BF₃ by sodium hydride:



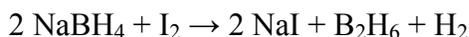
Two laboratory methods start from boron trichloride with lithium aluminium hydride or from boron trifluoride ether solution with sodium borohydride. Both methods yield in up to 30% of diborane:

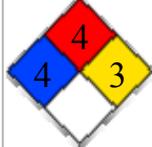


Older methods entail the direct reaction of borohydride salts with a non-oxidizing acid, such as phosphoric acid or dilute sulfuric acid.



Similarly, oxidation of borohydride salts has been demonstrated and remains convenient for small scale preparations. For example, using iodine as oxidizer:



Structure	
Coordination geometry	Tetrahedral (for boron)
Molecular shape	<i>see text</i>
Dipole moment	0 D
Hazards	
MSDS	External MSDS
EU classification	not listed
NFPA 704	
Flash point	flammable gas
Autoignition temperature	38 °C
Related compounds	
Related boron compounds	Decaborane BF ₃
Supplementary data page	
Structure and properties	<i>n</i> , ε _r , etc.
Thermodynamic data	Phase behaviour Solid, liquid, gas
Spectral data	UV, IR, NMR, MS
<div style="display: flex; align-items: center; justify-content: center;"> ✓ (what is this?) (verify) </div> <div style="text-align: center; margin-top: 0.5em;"> (http://en.wikipedia.org/w/index.php?title=Diborane&diff=cur&oldid=389085732) </div> <div style="text-align: center; margin-top: 0.5em;"> Except where noted otherwise, data are given for materials in their standard state (at 25 °C, 100 kPa) </div>	
Infobox references	

Reactions

Diborane is a highly reactive and versatile reagent that has a large number of applications.^[6] Its dominating reaction pattern involves formation of adducts with Lewis bases. Often such initial adducts proceed rapidly to give other products. It reacts with ammonia to form ammonia borane or the diammoniate of diborane, DADB, depending on the conditions used. Diborane also reacts readily with alkynes to form substituted alkene products which will readily undergo further addition reactions.

Diborane reacts with water to form hydrogen and boric acid.

The compound forms complexes with Lewis bases. Notable are the complexes with THF and dimethyl sulfide, both liquid compounds that are popular reducing agents in organic chemistry. In these 1:1 complexes, boron assumes a tetrahedral geometry, being bound to three hydrides and the Lewis base (THF or Me₂S). The THF adduct is usually prepared as a 1:5 solution in THF. The latter is indefinitely stable when stored under nitrogen at room temperature.

Reagent in organic synthesis

Diborane is the central organic synthesis reagent for hydroboration, whereby alkenes add across the B-H bonds to give trialkylboranes:



This reaction is regioselective, and the product trialkylboranes can be converted to useful organic derivatives. With bulky alkenes one can prepare species such as [HBR₂]₂, which are also useful reagents in more specialized applications.

Diborane is used as a reducing agent roughly complementary to the reactivity of lithium aluminium hydride. The compound readily reduces carboxylic acids to the corresponding alcohols, whereas ketones react only sluggishly.

History

Diborane was first synthesised in the 19th century by hydrolysis of metal borides, but it was never analysed. From 1912 to 1936, the major pioneer in the chemistry of boron hydrides, Alfred Stock, undertook his research that led to the methods for the synthesis and handling of the highly reactive, volatile, and often toxic boron hydrides. He proposed the first ethane-like structure of diborane.^[7] Electron diffraction measurements by S. H. Bauer initially appeared to support his proposed structure.^{[8][9]}

Because of a personal communication with L. Pauling (who supported the ethane-like structure), H. I. Schlessinger did not specifically discuss 3-center-2-electron bonding in his then classic review in the early 1940s.^[10] The review does, however, discuss the C_{2v} structure in some depth, "It is to be recognized that this formulation easily accounts for many of the chemical properties of diborane..."

In 1943 an undergraduate student at Balliol College, Oxford, H. Christopher Longuet-Higgins, published the currently accepted structure together with R. P. Bell.^[11] This structure had already been described in 1921.^{[12][13][14]} The years following the Longuet-Higgins/Bell proposal witnessed a colorful discussion about the correct structure. The debate ended with the electron diffraction measurement in 1951 by K.

Hedberg and V. Schomaker, with the confirmation of the structure shown in the schemes on this page.
[15]

William Nunn Lipscomb, Jr. further confirmed the molecular structure of boranes using X-ray crystallography in the 1950s, and developed theories to explain its bonding. Later, he applied the same methods to related problems, including the structure of carboranes on which he directed the research of future Nobel Prize winner Roald Hoffmann. Lipscomb himself received the Nobel Prize in Chemistry in 1976 for his efforts.

Other uses

Diborane is used in rocket propellants, as a rubber vulcaniser, as a catalyst for hydrocarbon polymerisation, as a flame-speed accelerator, and as a doping agent for the production of semiconductors. It is also an intermediate in the production of highly pure boron for semiconductor production.

Safety

The toxic effects of diborane are primarily due to its irritant properties. Short-term exposure to diborane can cause a sensation of tightness of the chest, shortness of breath, cough, and wheezing. These signs and symptoms can occur immediately or be delayed for up to 24 hours. Skin and eye irritation can also occur. Studies in animals have shown that diborane causes the same type of effects observed in humans.
[*citation needed*]

People exposed for a long time to low amounts of diborane have experienced respiratory irritation, seizures, fatigue, drowsiness, confusion, and occasional transient tremors.

References

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Further reading

H. C. Brown "Organic Synthesis via Boranes" John Wiley, New York, 1975. ISBN 0-471-11280-1.

External links

- International Chemical Safety Card 0432 (http://www.ilo.org/public/english/protection/safework/cis/products/icsc/dtasht/_icsc04/icsc0432.h)
- Computational Chemistry Wiki (<http://box27.bluehost.com/~edsanvil/wiki/index.php?title=Diborane>)
- National Pollutant Inventory - Boron and compounds (<http://www.npi.gov.au/database/substance-info/profiles/15.html>)
- NIOSH Pocket Guide to Chemical Hazards (<http://www.cdc.gov/niosh/npg/npgd0183.html>)
- U.S. EPA Acute Exposure Guideline Levels (<http://www.epa.gov/oppt/aegl/results65.htm>)

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