[Template:About](/wiki/Template:About" \o "Template:About) [Template:Pp-vandalism](/wiki/Template:Pp-vandalism) [Template:Good article](/wiki/Template:Good_article) [Template:Infobox lithium](/wiki/Template:Infobox_lithium) **Lithium** (from [Template:Lang-el](/wiki/Template:Lang-el), "stone") is a [chemical element](/wiki/Chemical_element) with the symbol **Li** and [atomic number](/wiki/Atomic_number) 3. It is a soft, silver-white [metal](/wiki/Metal) belonging to the [alkali metal](/wiki/Alkali_metal) [group](/wiki/Group_(periodic_table)) of chemical elements. Under [standard conditions](/wiki/Standard_conditions_for_temperature_and_pressure), it is the lightest metal and the least dense solid element. Like all alkali metals, lithium is highly reactive and flammable. For this reason, it is typically stored in [mineral oil](/wiki/Mineral_oil). When cut open, it exhibits a metallic [luster](/wiki/Luster_(mineralogy)), but contact with moist air [corrodes](/wiki/Corrosion) the surface quickly to a dull silvery gray, then black tarnish. Because of its high [reactivity](/wiki/Reactivity_(chemistry)), lithium never occurs freely in nature, and instead, appears only in [compounds](/wiki/Chemical_compound), which are usually [ionic](/wiki/Ionic_compound). Lithium occurs in a number of [pegmatitic](/wiki/Pegmatite) minerals, but due to its solubility as an ion, is present in ocean water and is commonly obtained from [brines](/wiki/Brine) and [clays](/wiki/Clay). On a commercial scale, lithium is isolated [electrolytically](/wiki/Electrolysis) from a mixture of [lithium chloride](/wiki/Lithium_chloride) and [potassium chloride](/wiki/Potassium_chloride).

The [nucleus](/wiki/Atomic_nucleus) of the lithium atom verges on instability, since the two stable lithium [isotopes](/wiki/Isotope) found in nature have among the lowest [binding energies](/wiki/Nuclear_binding_energy) per [nucleon](/wiki/Nucleon) of all stable [nuclides](/wiki/Nuclide). Because of its relative nuclear instability, lithium is less common in the solar system than 25 of the first 32 chemical elements even though the nuclei are very light in atomic weight.[[1]](#cite_note-1) For related reasons, lithium has important links to [nuclear physics](/wiki/Nuclear_physics). The [transmutation](/wiki/Nuclear_transmutation) of lithium atoms to [helium](/wiki/Helium) in 1932 was the first fully man-made [nuclear reaction](/wiki/Nuclear_reaction), and [lithium-6 deuteride](/wiki/Lithium_deuteride) serves as a [fusion](/wiki/Nuclear_fusion) fuel in [staged thermonuclear weapons](/wiki/Teller-Ulam_design).[[2]](#cite_note-2) Lithium and its compounds have several industrial applications, including heat-resistant glass and [ceramics](/wiki/Ceramic), [lithium grease](/wiki/Lithium_grease) lubricants, flux additives for iron, steel and aluminium production, [lithium batteries](/wiki/Lithium_battery), and [lithium-ion batteries](/wiki/Lithium-ion_battery). These uses consume more than three quarters of lithium production.

Trace amounts of lithium are present in all organisms. The element serves no apparent vital biological function, since animals and plants survive in good health without it, though non-vital functions have not been ruled out. The lithium [ion](/wiki/Ion) Li+ administered as any of several lithium [salts](/wiki/Salt_(chemistry)) has proved to be useful as a [mood-stabilizing](/wiki/Mood_stabilizer) drug in the treatment of [bipolar disorder](/wiki/Bipolar_disorder) in humans. [Template:TOC limit](/wiki/Template:TOC_limit)

## Contents

* 1 Properties[[edit](/index.php?title=(none)&action=edit&section=1)]
  + 1.1 Atomic and physical[[edit](/index.php?title=(none)&action=edit&section=2)]
  + 1.2 Chemistry and compounds[[edit](/index.php?title=(none)&action=edit&section=3)]
  + 1.3 Isotopes[[edit](/index.php?title=(none)&action=edit&section=5)]
* 2 Occurrence[[edit](/index.php?title=(none)&action=edit&section=6)]
  + 2.1 Astronomical[[edit](/index.php?title=(none)&action=edit&section=7)]
  + 2.2 Terrestrial[[edit](/index.php?title=(none)&action=edit&section=8)]
  + 2.3 Biological[[edit](/index.php?title=(none)&action=edit&section=9)]
* 3 History[[edit](/index.php?title=(none)&action=edit&section=10)]
* 4 Production[[edit](/index.php?title=(none)&action=edit&section=11)]
  + 4.1 Reserves[[edit](/index.php?title=(none)&action=edit&section=12)]
  + 4.2 Pricing[[edit](/index.php?title=(none)&action=edit&section=13)]
  + 4.3 Sources[[edit](/index.php?title=(none)&action=edit&section=14)]
    - 4.3.1 Geothermal wells[[edit](/index.php?title=(none)&action=edit&section=15)]
    - 4.3.2 Seawater[[edit](/index.php?title=(none)&action=edit&section=16)]
  + 4.4 Extraction[[edit](/index.php?title=(none)&action=edit&section=17)]
    - 4.4.1 Brine evaporation[[edit](/index.php?title=(none)&action=edit&section=18)]
    - 4.4.2 Dialysis[[edit](/index.php?title=(none)&action=edit&section=19)]
    - 4.4.3 Reverse osmosis[[edit](/index.php?title=(none)&action=edit&section=20)]
* 5 Uses[[edit](/index.php?title=(none)&action=edit&section=21)]
  + 5.1 Ceramics and glass[[edit](/index.php?title=(none)&action=edit&section=22)]
  + 5.2 Electrical and electronics[[edit](/index.php?title=(none)&action=edit&section=23)]
  + 5.3 Lubricating greases[[edit](/index.php?title=(none)&action=edit&section=24)]
  + 5.4 Metallurgy[[edit](/index.php?title=(none)&action=edit&section=25)]
  + 5.5 Silicon nano-welding[[edit](/index.php?title=(none)&action=edit&section=26)]
  + 5.6 Other chemical and industrial uses[[edit](/index.php?title=(none)&action=edit&section=27)]
    - 5.6.1 Pyrotechnics[[edit](/index.php?title=(none)&action=edit&section=28)]
    - 5.6.2 Air purification[[edit](/index.php?title=(none)&action=edit&section=29)]
    - 5.6.3 Optics[[edit](/index.php?title=(none)&action=edit&section=30)]
    - 5.6.4 Organic and polymer chemistry[[edit](/index.php?title=(none)&action=edit&section=31)]
    - 5.6.5 Military applications[[edit](/index.php?title=(none)&action=edit&section=32)]
  + 5.7 Nuclear[[edit](/index.php?title=(none)&action=edit&section=33)]
  + 5.8 Medicine[[edit](/index.php?title=(none)&action=edit&section=34)]
* 6 Precautions[[edit](/index.php?title=(none)&action=edit&section=35)]
  + 6.1 Regulation[[edit](/index.php?title=(none)&action=edit&section=36)]
* 7 See also[[edit](/index.php?title=(none)&action=edit&section=37)]
* 8 Notes[[edit](/index.php?title=(none)&action=edit&section=38)]
* 9 References[[edit](/index.php?title=(none)&action=edit&section=39)]
* 10 External links[[edit](/index.php?title=(none)&action=edit&section=40)]

## Properties[[edit](/index.php?title=(none)&action=edit&section=1)]

[Template:Main](/wiki/Template:Main)

### Atomic and physical[[edit](/index.php?title=(none)&action=edit&section=2)]

[Template:Multiple image](/wiki/Template:Multiple_image) Like the other [alkali metals](/wiki/Alkali_metal), lithium has a single [valence electron](/wiki/Valence_electron) that is easily given up to form a [cation](/wiki/Cation).<ref name=krebs/> Because of this, lithium is a good conductor of heat and electricity as well as a highly reactive element, though it is the least reactive of the alkali metals. Lithium's low reactivity is due to the proximity of its valence electron to its [nucleus](/wiki/Atomic_nucleus) (the remaining [two electrons](/wiki/Two-electron_atom) are in the [1s orbital](/wiki/S-orbital), much lower in energy, and do not participate in chemical bonds).<ref name=krebs/>

Lithium metal is soft enough to be cut with a knife. When cut, it possesses a silvery-white color that quickly changes to gray as it oxidizes to [lithium oxide](/wiki/Lithium_oxide).<ref name=krebs/> While it has one of the [lowest melting points](/wiki/List_of_elements_by_melting_point) among all metals (180 °C), it has the highest melting and boiling points of the alkali metals.[[3]](#cite_note-3) Lithium has a very low density (0.534 g/cm3), comparable with pine wood. It is the least dense of all elements that are solids at room temperature; the next lightest solid element (potassium, at 0.862 g/cm3) is more than 60% denser. Furthermore, apart from [helium](/wiki/Helium) and [hydrogen](/wiki/Hydrogen), it is less dense than any liquid element, being only 2/3 as dense as [liquid nitrogen](/wiki/Liquid_nitrogen) (0.808 g/cm3).[[4]](#cite_note-4) Lithium can float on the lightest hydrocarbon oils and is one of only three metals that can float on water, the other two being [sodium](/wiki/Sodium) and [potassium](/wiki/Potassium). [thumb|left|150px|Lithium floating in oil](/wiki/File:Lithium_element.jpg) Lithium's [coefficient of thermal expansion](/wiki/Coefficient_of_thermal_expansion) is twice that of [aluminium](/wiki/Aluminium) and almost four times that of [iron](/wiki/Iron).[[5]](#cite_note-5) Lithium is [superconductive](/wiki/Superconductive) below 400 [μK](/wiki/Microkelvin) at standard pressure[[6]](#cite_note-6) and at higher temperatures (more than 9 K) at very high pressures (>20 GPa).[[7]](#cite_note-7) At temperatures below 70 K, lithium, like sodium, undergoes [diffusionless phase change transformations](/wiki/Diffusionless_transformations). At 4.2 K it has a [rhombohedral crystal system](/wiki/Rhombohedral_crystal_system) (with a nine-layer repeat spacing); at higher temperatures it transforms to [face-centered cubic](/wiki/Face-centered_cubic) and then [body-centered cubic](/wiki/Body-centered_cubic). At liquid-helium temperatures (4 K) the rhombohedral structure is prevalent.[[8]](#cite_note-8) Multiple allotropic forms have been identified for lithium at high pressures.[[9]](#cite_note-9) Lithium has a mass [specific heat capacity](/wiki/Specific_heat_capacity) of 3.58 kilojoules per kilogram-kelvin, the highest of all solids.<ref name= CRC/>[[10]](#cite_note-10) Because of this, lithium metal is often used in [coolants](/wiki/Coolant) for [heat transfer](/wiki/Heat_transfer) applications.<ref name=CRC/>

### Chemistry and compounds[[edit](/index.php?title=(none)&action=edit&section=3)]

Lithium reacts with water easily, but with noticeably less energy than other alkali metals. The reaction forms [hydrogen](/wiki/Hydrogen) gas and [lithium hydroxide](/wiki/Lithium_hydroxide) in aqueous solution.<ref name=krebs/> Because of its reactivity with water, lithium is usually stored in a hydrocarbon sealant, often [petroleum jelly](/wiki/Petroleum_jelly). Though the heavier alkali metals can be stored in more dense substances, such as [mineral oil](/wiki/Mineral_oil), lithium is not dense enough to be fully submerged in these liquids.<ref name=emsley/> In moist air, lithium rapidly tarnishes to form a black coating of [lithium hydroxide](/wiki/Lithium_hydroxide) (LiOH and LiOH·H2O), [lithium nitride](/wiki/Lithium_nitride) (Li3N) and [lithium carbonate](/wiki/Lithium_carbonate) (Li2CO3, the result of a secondary reaction between LiOH and [CO2](/wiki/Carbon_dioxide)).[[11]](#cite_note-11) [thumb|left|Hexameric structure of the](/wiki/File:Butyllithium-hexamer-from-xtal-3D-balls-A.png) [n-butyllithium](/wiki/N-butyllithium) fragment in a crystal When placed over a flame, lithium compounds give off a striking crimson color, but when it burns strongly the flame becomes a brilliant silver. Lithium will ignite and burn in oxygen when exposed to water or water vapors.[[12]](#cite_note-12) Lithium is [flammable](/wiki/Flammable), and it is potentially explosive when exposed to air and especially to water, though less so than the other [alkali metals](/wiki/Alkali_metals). The lithium-water reaction at normal temperatures is brisk but nonviolent because the hydrogen produced does not ignite on its own. As with all alkali metals, lithium fires are difficult to extinguish, requiring dry powder fire extinguishers ([Class D](/wiki/Class_D_fire_extinguisher) type). Lithium is the only metal which reacts with [nitrogen](/wiki/Nitrogen) under [normal conditions](/wiki/Standard_conditions_for_temperature_and_pressure).[[13]](#cite_note-13)[[14]](#cite_note-14) Lithium has a [diagonal relationship](/wiki/Diagonal_relationship) with [magnesium](/wiki/Magnesium), an element of similar atomic and [ionic radius](/wiki/Ionic_radius). Chemical resemblances between the two metals include the formation of a [nitride](/wiki/Nitride) by reaction with N2, the formation of an [oxide](/wiki/Lithium_oxide) ([Template:Chem](/wiki/Template:Chem)) and peroxide ([Template:Chem](/wiki/Template:Chem)) when burnt in O2, [salts](/wiki/Salt_(chemistry)) with similar [solubilities](/wiki/Solubility), and thermal instability of the [carbonates](/wiki/Carbonate) and nitrides.[[11]](#cite_note-11)[[15]](#cite_note-15) The metal reacts with hydrogen gas at high temperatures to produce [lithium hydride](/wiki/Lithium_hydride) (LiH).[[16]](#cite_note-16) Other known [binary compounds](/wiki/Binary_compound) include [halides](/wiki/Halide) ([LiF](/wiki/Lithium_fluoride), [LiCl](/wiki/Lithium_chloride), [LiBr](/wiki/Lithium_bromide), [LiI](/wiki/Lithium_iodide)), [sulfide](/wiki/Sulfide) ([Template:Chem](/wiki/Template:Chem)), [superoxide](/wiki/Superoxide) ([Template:Chem](/wiki/Template:Chem)), and [carbide](/wiki/Carbide) ([Template:Chem](/wiki/Template:Chem)). Many other inorganic compounds are known in which lithium combines with [anions](/wiki/Anion) to form salts: [borates](/wiki/Lithium_borate), [amides](/wiki/Lithium_amide), [carbonate](/wiki/Lithium_carbonate), [nitrate](/wiki/Lithium_nitrate), or [borohydride](/wiki/Borohydride) ([Template:Chem](/wiki/Template:Chem)). [Lithium aluminium hydride](/wiki/Lithium_aluminium_hydride) ([Template:Chem](/wiki/Template:Chem)) is commonly used as a reducing agent in organic synthesis.

Multiple [organolithium reagents](/wiki/Organolithium_reagent) are known in which there is a direct [bond](/wiki/Covalent_bond) between [carbon](/wiki/Carbon) and lithium atoms, effectively creating a [carbanion](/wiki/Carbanion). These are extremely powerful [bases](/wiki/Base_(chemistry)) and [nucleophiles](/wiki/Carbon_nucleophile). In many of these organolithium compounds, the lithium ions tend to aggregate into high-symmetry clusters by themselves, which is relatively common for alkali cations.[[17]](#cite_note-17) [LiHe](/wiki/LiHe), a very weakly interacting [van der Waals compound](/wiki/Van_der_Waals_compound), has been detected at very low temperatures.[[18]](#cite_note-18)===Ferromagnetism in Lithium-6 as a cryogenic gas=== Experimental evidence suggests that, in its gaseous form at temperatures of about 150 nano[Kelvin](/wiki/Kelvin), lithium-6 exhibits ferromagnetism; if fully confirmed, this is the first example of ferromagnetism observed in any gas under any conditions. Reservations on the reality of the demonstration are tenuous however, largely the desirability of direct observation that the spins of the atoms were parallel.[[19]](#cite_note-19)

### Isotopes[[edit](/index.php?title=(none)&action=edit&section=5)]

[Template:Main](/wiki/Template:Main) Naturally occurring lithium is composed of two stable [isotopes](/wiki/Isotope), 6Li and 7Li, the latter being the more abundant (92.5% [natural abundance](/wiki/Natural_abundance)).<ref name=krebs/><ref name=emsley/><ref name=isotopesproject>[Template:Cite web](/wiki/Template:Cite_web)</ref> Both natural isotopes have anomalously low [nuclear binding energy](/wiki/Nuclear_binding_energy) per nucleon (compared to the neighboring elements on the [periodic table](/wiki/Periodic_chart_of_the_elements), [helium](/wiki/Helium) and [beryllium](/wiki/Beryllium)); lithium is the only low numbered element that can produce net energy through [nuclear fission](/wiki/Nuclear_fission). The two lithium nuclei have lower binding energies per nucleon than any other stable nuclides other than [deuterium](/wiki/Deuterium) and [helium-3](/wiki/Helium-3).<ref name=bind>[:File:Binding energy curve - common isotopes.svg](/wiki/File:Binding_energy_curve_-_common_isotopes.svg) shows binding energies of stable nuclides graphically; the source of the data-set is given in the figure background.</ref> As a result of this, though very light in atomic weight, lithium is less common in the Solar System than 25 of the first 32 chemical elements.[[1]](#cite_note-1) Seven [radioisotopes](/wiki/Radioisotope) have been characterized, the most stable being 8Li with a [half-life](/wiki/Half-life) of 838 [ms](/wiki/Millisecond) and 9Li with a half-life of 178 ms. All of the remaining [radioactive](/wiki/Radioactive) isotopes have half-lives that are shorter than 8.6 ms. The shortest-lived isotope of lithium is 4Li, which decays through [proton emission](/wiki/Proton_emission) and has a half-life of 7.6 × 10−23 s.<ref name=nuclidetable>[Template:Cite web](/wiki/Template:Cite_web)</ref>

7Li is one of the [primordial elements](/wiki/Primordial_elements) (or, more properly, primordial [nuclides](/wiki/Nuclide)) produced in [Big Bang nucleosynthesis](/wiki/Big_Bang_nucleosynthesis). A small amount of both 6Li and 7Li are produced in stars, but are thought to be "[burned](/wiki/Lithium_burning)" as fast as produced.[[20]](#cite_note-20) Additional small amounts of lithium of both 6Li and 7Li may be generated from solar wind, cosmic rays hitting heavier atoms, and from early solar system 7[Be](/wiki/Beryllium) and 10Be radioactive decay.[[21]](#cite_note-21) While lithium is created in stars during [stellar nucleosynthesis](/wiki/Stellar_nucleosynthesis), it is further burned. 7Li can also be generated in [carbon stars](/wiki/Carbon_star).[[22]](#cite_note-22) Lithium isotopes fractionate substantially during a wide variety of natural processes,[[23]](#cite_note-23) including mineral formation (chemical precipitation), [metabolism](/wiki/Metabolism), and [ion exchange](/wiki/Ion_exchange). Lithium ions substitute for [magnesium](/wiki/Magnesium) and iron in octahedral sites in [clay](/wiki/Clay) minerals, where 6Li is preferred to 7Li, resulting in enrichment of the light isotope in processes of hyperfiltration and rock alteration. The exotic 11Li is known to exhibit a [nuclear halo](/wiki/Nuclear_halo). The process known as [laser isotope separation](/wiki/Atomic_vapor_laser_isotope_separation) can be used to separate lithium isotopes, in particular 7Li from 6Li.[[24]](#cite_note-24) Nuclear weapons manufacture and other nuclear physics applications are a major source of artificial lithium fractionation, with the light isotope 6Li being retained by industry and military stockpiles to such an extent that it has caused slight but measurable change in the 6Li to 7Li ratios in natural sources, such as rivers. This has led to unusual uncertainty in the standardized [atomic weight](/wiki/Atomic_weight) of lithium, since this quantity depends on the natural abundance ratios of these naturally-occurring stable lithium isotopes, as they are available in commercial lithium mineral sources.[[25]](#cite_note-25)

## Occurrence[[edit](/index.php?title=(none)&action=edit&section=6)]

[thumb|350px|Lithium is about as common as](/wiki/File:Elemental_abundances.svg) [chlorine](/wiki/Chlorine) in the Earth's upper continental [crust](/wiki/Crust_(geology)), on a per-atom basis.

### Astronomical[[edit](/index.php?title=(none)&action=edit&section=7)]

[Template:Main](/wiki/Template:Main) According to modern cosmological theory, lithium—in both stable isotopes (lithium-6 and lithium-7)—was one of the 3 elements [synthesized](/wiki/Nucleosynthesis) in the [Big Bang](/wiki/Big_Bang).[[26]](#cite_note-26) Though the amount of lithium generated in [Big Bang nucleosynthesis](/wiki/Big_Bang_nucleosynthesis) is dependent upon the number of [photons](/wiki/Photon) per [baryon](/wiki/Baryon), for accepted values the lithium abundance can be calculated, and there is a "cosmological lithium discrepancy" in the Universe: older stars seem to have less lithium than they should, and some younger stars have much more. The lack of lithium in older stars is apparently caused by the "mixing" of lithium into the interior of stars, where it is destroyed,<ref name=cld>[Template:Cite news](/wiki/Template:Cite_news)</ref> while lithium is produced in younger stars. Though it transmutes into two atoms of [helium](/wiki/Helium) due to collision with a [proton](/wiki/Proton) at temperatures above 2.4 million degrees Celsius (most stars easily attain this temperature in their interiors), lithium is more abundant than current computations would predict in later-generation stars.<ref name=emsley/> [thumb|](/wiki/File:Nova_Centauri_2013_ESO.jpg)[Nova Centauri 2013](/wiki/Nova_Centauri_2013) is the first in which evidence of lithium has been found.[[27]](#cite_note-27) Though it was one of the three first elements to be synthesized in the Big Bang, lithium, together with [beryllium](/wiki/Beryllium) and [boron](/wiki/Boron) are markedly less abundant than other elements. This is a result of the low temperature necessary to destroy lithium, and a lack of common processes to produce it.<ref name=wesleyan>[Template:Cite web](/wiki/Template:Cite_web)</ref>

Lithium is also found in [brown dwarf](/wiki/Brown_dwarf) substellar objects and certain anomalous orange stars. Because lithium is present in cooler, less-massive brown dwarfs, but is destroyed in hotter [red dwarf](/wiki/Red_dwarf) stars, its presence in the stars' spectra can be used in the "lithium test" to differentiate the two, as both are smaller than the Sun.<ref name=emsley/>[[28]](#cite_note-28)[[29]](#cite_note-29) Certain orange stars can also contain a high concentration of lithium. Those orange stars found to have a higher than usual concentration of lithium (such as [Centaurus X-4](/wiki/Centaurus_X-4)) orbit massive objects—neutron stars or black holes—whose gravity evidently pulls heavier lithium to the surface of a hydrogen-helium star, causing more lithium to be observed.<ref name=emsley/>

### Terrestrial[[edit](/index.php?title=(none)&action=edit&section=8)]

|  |  |  |
| --- | --- | --- |
| Lithium mine production (2015) and reserves in tonnes[[30]](#cite_note-30) | | |
| **Country** | **Production** | **Reserves<ref group=note name=res/>** |
| [Template:Flag](/wiki/Template:Flag) | 3,800 | 2,000,000 |
| [Template:Flag](/wiki/Template:Flag) | 13,400 | 1,500,000 |
| [Template:Flag](/wiki/Template:Flag) | 160 | 48,000 |
| [Template:Flag](/wiki/Template:Flag) (2010) | 480 | 180,000 |
| [Template:Flag](/wiki/Template:Flag) | 11,700 | 7,500,000 |
| [Template:Flag](/wiki/Template:Flag) | 2,200 | 3,200,000 |
| [Template:Flag](/wiki/Template:Flag) | 300 | 60,000 |
| [Template:Flag](/wiki/Template:Flag) | 900 | 23,000 |
| **World total** | **32,500** | **14,000,000** |

[Template:See also](/wiki/Template:See_also) Although lithium is widely distributed on Earth, it does not naturally occur in elemental form due to its high reactivity.<ref name=krebs>[Template:Cite book](/wiki/Template:Cite_book)</ref> The total lithium content of seawater is very large and is estimated as 230 billion tonnes, where the element exists at a relatively constant concentration of 0.14 to 0.25 parts per million (ppm),[[31]](#cite_note-31)<ref name=enc/> or 25 [micromolar](/wiki/Micromolar);[[32]](#cite_note-32) higher concentrations approaching 7 ppm are found near [hydrothermal vents](/wiki/Hydrothermal_vents).<ref name=enc/>

Estimates for the Earth's [crustal](/wiki/Crust_(geology)) content range from 20 to 70 ppm by weight.[[11]](#cite_note-11) In keeping with its name, lithium forms a minor part of [igneous rocks](/wiki/Igneous_rock), with the largest concentrations in [granites](/wiki/Granite). Granitic [pegmatites](/wiki/Pegmatite) also provide the greatest abundance of lithium-containing minerals, with [spodumene](/wiki/Spodumene) and [petalite](/wiki/Petalite) being the most commercially viable sources.[[11]](#cite_note-11) Another significant mineral of lithium is [lepidolite](/wiki/Lepidolite).[[33]](#cite_note-33) A newer source for lithium is [hectorite](/wiki/Hectorite) clay, the only active development of which is through the Western Lithium Corporation in the United States.[[34]](#cite_note-34) At 20 mg lithium per kg of Earth's crust,[[35]](#cite_note-35) lithium is the 25th most abundant element.

According to the *Handbook of Lithium and Natural Calcium*, "Lithium is a comparatively rare element, although it is found in many rocks and some brines, but always in very low concentrations. There are a fairly large number of both lithium mineral and brine deposits but only comparatively few of them are of actual or potential commercial value. Many are very small, others are too low in grade."[[36]](#cite_note-36) The [US Geological Survey](/wiki/US_Geological_Survey) estimates that in 2010, Chile had the largest reserves by far (7.5 million tonnes)[[37]](#cite_note-37) and the highest annual production (8,800 tonnes). One of the largest *reserve bases*<ref group=note name=res>[Apendixes](http://minerals.usgs.gov/minerals/pubs/mcs/2011/mcsapp2011.pdf). By USGS definitions, the reserve base "may encompass those parts of the resources that have a reasonable potential for becoming economically available within planning horizons beyond those that assume proven technology and current economics. The reserve base includes those resources that are currently economic (reserves), marginally economic (marginal reserves), and some of those that are currently subeconomic (subeconomic resources)."</ref> of lithium is in the [Salar de Uyuni](/wiki/Salar_de_Uyuni) area of [Bolivia](/wiki/Bolivia), which has 5.4 million tonnes. Other major suppliers include Australia, Argentina and China.[[30]](#cite_note-30)<ref name=meridian>[Template:Cite web](/wiki/Template:Cite_web)</ref>

In June 2010, the [*New York Times*](/wiki/New_York_Times) reported that American geologists were conducting ground surveys on [dry](/wiki/Dry_lake) [salt lakes](/wiki/Salt_lake) in western [Afghanistan](/wiki/Afghanistan) believing that large deposits of lithium are located there. "Pentagon officials said that their initial analysis at one location in [Ghazni Province](/wiki/Ghazni_Province) showed the potential for lithium deposits as large as those of Bolivia, which now has the world's largest known lithium reserves."[[38]](#cite_note-38) These estimates are "based principally on old data, which was gathered mainly by the Soviets during their occupation of Afghanistan from 1979–1989". Stephen Peters, the head of the USGS's Afghanistan Minerals Project, said that he was unaware of [USGS](/wiki/USGS) involvement in any new surveying for minerals in Afghanistan in the past two years. 'We are not aware of any discoveries of lithium,' he said."[[39]](#cite_note-39)

### Biological[[edit](/index.php?title=(none)&action=edit&section=9)]

Lithium is found in trace amount in numerous plants, plankton, and invertebrates, at concentrations of 69 to 5,760 parts per billion (ppb). In vertebrates the concentration is slightly lower, and nearly all vertebrate tissue and body fluids contain lithium ranging from 21 to 763 ppb.<ref name=enc/> Marine organisms tend to bioaccumulate lithium more than terrestrial organisms.[[40]](#cite_note-40) Whether lithium has a physiological role in any of these organisms is unknown.<ref name=enc>[Template:Cite web](/wiki/Template:Cite_web)</ref>

## History[[edit](/index.php?title=(none)&action=edit&section=10)]

[thumb|Johan August Arfwedson is credited with the discovery of lithium in 1817](/wiki/File:Arfwedson_Johan_A.jpg) [Petalite](/wiki/Petalite) (LiAlSi4O10) was discovered in 1800 by the [Brazilian](/wiki/Brazil) chemist and statesman [José Bonifácio de Andrada e Silva](/wiki/José_Bonifácio_de_Andrada_e_Silva) in a mine on the island of [Utö, Sweden](/wiki/Utö,_Sweden).[[41]](#cite_note-41)<ref name=mindat>[Template:Cite web](/wiki/Template:Cite_web)</ref><ref name=webelementshistory>[Template:Cite web](/wiki/Template:Cite_web)</ref><ref name=discovery>[Template:Cite book](/wiki/Template:Cite_book)</ref> However, it was not until 1817 that [Johan August Arfwedson](/wiki/Johan_August_Arfwedson), then working in the laboratory of the chemist [Jöns Jakob Berzelius](/wiki/Jöns_Jakob_Berzelius), [detected](/wiki/Discovery_of_the_chemical_elements) the presence of a new element while analyzing petalite ore.[[42]](#cite_note-42)<ref name=berzelius>[Template:Cite web](/wiki/Template:Cite_web)</ref><ref name=uwis>[Template:Cite web](/wiki/Template:Cite_web)</ref><ref name=vanderkrogt>[Template:Cite web](/wiki/Template:Cite_web)</ref> This element formed compounds similar to those of [sodium](/wiki/Sodium) and [potassium](/wiki/Potassium), though its [carbonate](/wiki/Lithium_carbonate) and [hydroxide](/wiki/Lithium_hydroxide) were less [soluble in water](/wiki/Solubility) and more [alkaline](/wiki/Base_(chemistry)).<ref name=compounds>[Template:Cite web](/wiki/Template:Cite_web)</ref> Berzelius gave the alkaline material the name "*lithion*/*lithina*", from the [Greek](/wiki/Greek_language) word *λιθoς* (transliterated as *lithos*, meaning "stone"), to reflect its discovery in a solid mineral, as opposed to potassium, which had been discovered in plant ashes, and sodium which was known partly for its high abundance in animal blood. He named the metal inside the material "*lithium*".<ref name=krebs/><ref name=webelementshistory/><ref name=vanderkrogt/>

Arfwedson later showed that this same element was present in the minerals [spodumene](/wiki/Spodumene) and [lepidolite](/wiki/Lepidolite).<ref name=webelementshistory/> In 1818, [Christian Gmelin](/wiki/Christian_Gmelin) was the first to observe that lithium salts give a bright red color to flame.<ref name=webelementshistory/>[[43]](#cite_note-43) However, both Arfwedson and Gmelin tried and failed to isolate the pure element from its salts.[[44]](#cite_note-44) It was not isolated until 1821, when [William Thomas Brande](/wiki/William_Thomas_Brande) obtained it by [electrolysis](/wiki/Electrolysis) of [lithium oxide](/wiki/Lithium_oxide), a process that had previously been employed by the chemist Sir [Humphry Davy](/wiki/Humphry_Davy) to isolate the alkali metals potassium and sodium.<ref name=emsley>[Template:Cite book](/wiki/Template:Cite_book)</ref>[[44]](#cite_note-44)[[45]](#cite_note-45)[[46]](#cite_note-46)[[47]](#cite_note-47) Brande also described some pure salts of lithium, such as the chloride, and, estimating that lithia ([lithium oxide](/wiki/Lithium_oxide)) contained about 55% metal, estimated the atomic weight of lithium to be around 9.8 g/mol (modern value ~6.94 g/mol).[[48]](#cite_note-48) In 1855, larger quantities of lithium were produced through the electrolysis of [lithium chloride](/wiki/Lithium_chloride) by [Robert Bunsen](/wiki/Robert_Bunsen) and [Augustus Matthiessen](/wiki/Augustus_Matthiessen).<ref name=webelementshistory/>[[49]](#cite_note-49) The discovery of this procedure led to commercial production of lithium in 1923 by the German company [Metallgesellschaft AG](/wiki/Metallgesellschaft_AG), which performed an electrolysis of a liquid mixture of lithium chloride and [potassium chloride](/wiki/Potassium_chloride).<ref name=webelementshistory/>[[50]](#cite_note-50)[[51]](#cite_note-51) The production and use of lithium underwent several drastic changes in history. The first major application of lithium was in high-temperature [lithium greases](/wiki/Lithium_grease) for aircraft engines and similar applications in [World War II](/wiki/World_War_II) and shortly after. This use was supported by the fact that lithium-based [soaps](/wiki/Soap) have a higher melting point than other alkali soaps, and are less corrosive than calcium based soaps. The small market for lithium soaps and lubricating greases was supported by several small mining operations mostly in the United States.

The demand for lithium increased dramatically during the [Cold War](/wiki/Cold_War) with the production of [nuclear fusion weapons](/wiki/Nuclear_weapon_design). Both lithium-6 and lithium-7 produce [tritium](/wiki/Tritium) when irradiated by neutrons, and are thus useful for the production of tritium by itself, as well as a form of solid fusion fuel used inside hydrogen bombs in the form of [lithium deuteride](/wiki/Lithium_deuteride). The United States became the prime producer of lithium in the period between the late 1950s and the mid-1980s. At the end, the stockpile of lithium was roughly 42,000 tonnes of lithium hydroxide. The stockpiled lithium was depleted in lithium-6 by 75%, which was enough to affect the measured [atomic weight](/wiki/Atomic_weight) of lithium in many standardized chemicals, and even the atomic weight of lithium in some "natural sources" of lithium ion which had been "contaminated" by lithium salts discharged from isotope separation facilities, which had found its way into ground water.[[25]](#cite_note-25)[[52]](#cite_note-52) Lithium was used to decrease the melting temperature of glass and to improve the melting behavior of [aluminium oxide](/wiki/Aluminium_oxide) when using the [Hall-Héroult process](/wiki/Hall-Héroult_process).[[53]](#cite_note-53)[[54]](#cite_note-54) These two uses dominated the market until the middle of the 1990s. After the end of the [nuclear arms race](/wiki/Nuclear_arms_race), the demand for lithium decreased and the sale of Department of Energy stockpiles on the open market further reduced prices.[[52]](#cite_note-52) But in the mid-1990s, several companies started to extract lithium from [brine](/wiki/Brine) which proved to be a less expensive method than underground or even open-pit mining. Most of the mines closed or shifted their focus to other materials because only the ore from zoned pegmatites could be mined for a competitive price. For example, the US mines near [Kings Mountain, North Carolina](/wiki/Kings_Mountain,_North_Carolina) closed before the turn of the 21st century.

The development of lithium ion batteries increased the demand for lithium and became the dominant use in 2007.[[55]](#cite_note-55) With the surge of lithium demand in batteries in the 2000s, new companies have expanded brine extraction efforts to meet the rising demand.[[56]](#cite_note-56)[[57]](#cite_note-57)

## Production[[edit](/index.php?title=(none)&action=edit&section=11)]

[Template:Multiple image](/wiki/Template:Multiple_image) [thumb|World production trend of lithium](/wiki/File:Lithium_world_production.svg)

Lithium production has greatly increased since the end of [World War II](/wiki/World_War_II). The metal is separated from other elements in [igneous minerals](/wiki/Igneous_rock). Lithium salts are extracted from water in [mineral springs](/wiki/Mineral_springs), [brine](/wiki/Brine) pools and brine deposits. The metal is produced through [electrolysis](/wiki/Electrolysis) from a mixture of fused 55% [lithium chloride](/wiki/Lithium_chloride) and 45% [potassium chloride](/wiki/Potassium_chloride) at about 450 °C.[[58]](#cite_note-58) In 1998 it was about [Template:Nowrap](/wiki/Template:Nowrap) (or 43 USD/[lb](/wiki/Pound_(mass))).[[59]](#cite_note-59)

### Reserves[[edit](/index.php?title=(none)&action=edit&section=12)]

Worldwide identified reserves in 2008 were estimated by the [US Geological Survey](/wiki/US_Geological_Survey) (USGS) as 13 million [tonnes](/wiki/Tonne),[[30]](#cite_note-30) though it is difficult to accurately estimate the world's lithium reserves.[[60][60]](#cite_note-60)[[61]](#cite_note-61) Deposits are found in South America throughout the [Andes](/wiki/Andes) mountain chain. [Chile](/wiki/Chile) is the leading producer, followed by [Argentina](/wiki/Argentina). Both countries recover lithium from brine pools. In the United States lithium is recovered from brine pools in [Nevada](/wiki/Nevada).[[62]](#cite_note-62) However, half the world's known reserves are located in [Bolivia](/wiki/Bolivia), along the central eastern slope of the Andes. In 2009, Bolivia negotiated with Japanese, French and Korean firms to begin extraction.[[63]](#cite_note-63) According to USGS, Bolivia's [Uyuni](/wiki/Uyuni) Desert has 5.4 million tonnes of lithium.[[63]](#cite_note-63)[[64]](#cite_note-64) A newly discovered deposit in [Wyoming's](/wiki/Wyoming) [Rock Springs Uplift](/wiki/Rock_Springs_Uplift) is estimated at 228,000 tons. Additional deposits in the same formation were extrapolated to be as much as 18 million tons.[[65]](#cite_note-65) Opinions differ about potential growth. A 2008 study concluded that "realistically achievable lithium carbonate production will be sufficient for only a small fraction of future [PHEV](/wiki/PHEV) and [EV](/wiki/Electric_vehicle) global market requirements", that "demand from the portable electronics sector will absorb much of the planned production increases in the next decade", and that "mass production of lithium carbonate is not environmentally sound, it will cause irreparable ecological damage to ecosystems that should be protected and that [LiIon](/wiki/LiIon) propulsion is incompatible with the notion of the 'Green Car'".[[66]](#cite_note-66) However, according to a 2011 study conducted at [Lawrence Berkeley National Laboratory](/wiki/Lawrence_Berkeley_National_Laboratory) and the [University of California, Berkeley](/wiki/University_of_California,_Berkeley), the currently estimated reserve base of lithium should not be a limiting factor for large-scale battery production for electric vehicles because an estimated 1 billion 40 [kWh](/wiki/KWh) Li-based batteries could be built with current reserves[[67]](#cite_note-67) - about 10 kg of lithium per car.[[68]](#cite_note-68) Another 2011 study by researchers from the [University of Michigan](/wiki/University_of_Michigan) and [Ford Motor Company](/wiki/Ford_Motor_Company) found sufficient resources to support global demand until 2100, including the lithium required for the potential widespread transportation use. The study estimated global reserves at 39 million tons, and total demand for lithium during the 90-year period analyzed at 12–20 million tons, depending on the scenarios regarding economic growth and recycling rates.[[69]](#cite_note-69) On June 9, 2014, the *Financialist* stated that demand for lithium was growing at more than 12 percent a year; according to Credit Suisse, this rate exceeds projected availability by 25 percent. The publication compared the 2014 lithium situation with oil, whereby "higher oil prices spurred investment in expensive deepwater and oil sands production techniques"; that is, the price of lithium will continue to rise until more expensive production methods that can boost total output receive the attention of investors.[[70]](#cite_note-70)

### Pricing[[edit](/index.php?title=(none)&action=edit&section=13)]

After the [2007 financial crisis](/wiki/Financial_crisis_of_2007–08), major suppliers such as [Sociedad Química y Minera](/wiki/Sociedad_Química_y_Minera) (SQM) dropped [lithium carbonate](/wiki/Lithium_carbonate) pricing by 20%.[[71]](#cite_note-71) Prices rose in 2012. A 2012 [Business Week](/wiki/Bloomberg_Businessweek) article outlined the [oligopoly](/wiki/Oligopoly) in the lithium space: "SQM, controlled by billionaire Julio Ponce, is the second-largest, followed by Rockwood, which is backed by Henry Kravis’s KKR & Co., and Philadelphia-based FMC". Global consumption may jump to 300,000 metric tons a year by 2020 from about 150,000 tons in 2012, to match demand for lithium batteries that has been growing at about 25 percent a year, outpacing the 4 percent to 5 percent overall gain in lithium.[[72]](#cite_note-72)

### Sources[[edit](/index.php?title=(none)&action=edit&section=14)]

#### Geothermal wells[[edit](/index.php?title=(none)&action=edit&section=15)]

A potential source is [geothermal wells](/wiki/Geothermal_electricity). Geothermal fluids carry leachates to the surface;[[73]](#cite_note-73) recovery of lithium has been demonstrated in the field.[[74]](#cite_note-74) The lithium is separated by simple filtration. The process and environmental costs are primarily those of the already-operating well; net environmental impacts may thus be positive.[[75]](#cite_note-75)

#### Seawater[[edit](/index.php?title=(none)&action=edit&section=16)]

Lithium is present in seawater, but practical extraction techniques have yet to be commercialized.[[76]](#cite_note-76)

### Extraction[[edit](/index.php?title=(none)&action=edit&section=17)]

#### Brine evaporation[[edit](/index.php?title=(none)&action=edit&section=18)]

As of 2015 most of the world’s lithium production is in South America, where lithium-containing brine is extracted from underground pools and concentrated by solar evaporation. In 2010, Simbol Materials received a $3 million grant from the [U.S. Department of Energy](/wiki/U.S._Department_of_Energy) for a pilot project to show the financial feasibility of extracting high-quality lithium from [geothermal](/wiki/Geothermal) brine. It uses brine from the 49.9-megawatt Featherstone geothermal power plant in California's [Imperial Valley](/wiki/Imperial_Valley). The extracted fluid is passed through a series of membranes, filters and adsorption materials to extract lithium.[[77]](#cite_note-77) The standard extraction technique is to evaporate water from brine. Each batch takes from 18 to 24 months.[[76]](#cite_note-76)

#### Dialysis[[edit](/index.php?title=(none)&action=edit&section=19)]

In 2015 researchers announced a new seawater extraction process using [dialysis](/wiki/Dialysis_(biochemistry)). The dialysis cell uses a superconducting [membrane](/wiki/Membrane). Lithium is the only ion in the seawater that can pass through the membrane.[[76]](#cite_note-76)

#### Reverse osmosis[[edit](/index.php?title=(none)&action=edit&section=20)]

[Reverse osmosis](/wiki/Reverse_osmosis) has been proposed as an alternative, but has not been commercialized.[[76]](#cite_note-76)

## Uses[[edit](/index.php?title=(none)&action=edit&section=21)]

[thumb|Estimates of global lithium uses in 2011](/wiki/File:Global_Lithium_Uses.svg)[[78]](#cite_note-78)[Template:Legend](/wiki/Template:Legend) [Template:Legend](/wiki/Template:Legend) [Template:Legend](/wiki/Template:Legend) [Template:Legend](/wiki/Template:Legend) [Template:Legend](/wiki/Template:Legend) [Template:Legend](/wiki/Template:Legend) [Template:Legend](/wiki/Template:Legend) [Template:Legend](/wiki/Template:Legend) [Template:Legend](/wiki/Template:Legend)

### Ceramics and glass[[edit](/index.php?title=(none)&action=edit&section=22)]

Lithium oxide is widely used as a [flux](/wiki/Flux_(metallurgy)) for processing [silica](/wiki/Silica), reducing the [melting point](/wiki/Melting_point) and [viscosity](/wiki/Viscosity) of the material and leading to [glazes](/wiki/Ceramic_glaze) with improved physical properties including low coefficients of thermal expansion. Worldwide, this is the single largest use for lithium compounds.[[78]](#cite_note-78)[[79]](#cite_note-79) Glazes containing lithium oxides are used for ovenware. [Lithium carbonate](/wiki/Lithium_carbonate) (Li2CO3) is generally used in this application because it converts to the oxide upon heating.[[80]](#cite_note-80)

### Electrical and electronics[[edit](/index.php?title=(none)&action=edit&section=23)]

Late in the 20th century, lithium became an important component of battery electrolytes and electrodes, because of its high [electrode potential](/wiki/Electrode_potential). Because of its low [atomic mass](/wiki/Atomic_mass), it has a high charge- and power-to-weight ratio. A typical [lithium-ion battery](/wiki/Lithium-ion_battery) can generate approximately 3 [volts](/wiki/Volt) per cell, compared with 2.1 volts for [lead-acid](/wiki/Lead–acid_battery) or 1.5 volts for [zinc-carbon cells](/wiki/Zinc-carbon_cell). Lithium-ion batteries, which are rechargeable and have a high energy density, should not be confused with [lithium batteries](/wiki/Lithium_batteries), which are [disposable](/wiki/Disposable) ([primary](/wiki/Primary_cell)) [batteries](/wiki/Battery_(electricity)) with lithium or its compounds as the [anode](/wiki/Anode).[[81]](#cite_note-81)[[82]](#cite_note-82) Other rechargeable batteries that use lithium include the [lithium-ion polymer battery](/wiki/Lithium-ion_polymer_battery), [lithium iron phosphate battery](/wiki/Lithium_iron_phosphate_battery), and the [nanowire battery](/wiki/Nanowire_battery).

### Lubricating greases[[edit](/index.php?title=(none)&action=edit&section=24)]

[Template:Main](/wiki/Template:Main) The third most common use of lithium is in greases. Lithium hydroxide is a strong [base](/wiki/Base_(chemistry)) and, when heated with a fat, produces a soap made of lithium [stearate](/wiki/Stearate). Lithium soap has the ability to [thicken](/wiki/Thickening_agent) oils, and it is used to manufacture all-purpose, high-temperature [lubricating greases](/wiki/Grease_(lubricant)).<ref name=CRC/>[[83]](#cite_note-83)[[84]](#cite_note-84)

### Metallurgy[[edit](/index.php?title=(none)&action=edit&section=25)]

Lithium (e.g. as lithium carbonate) is used as an additive to [continuous casting](/wiki/Continuous_casting) mould flux slags where it increases fluidity,[[85]](#cite_note-85)[[86]](#cite_note-86) a use which accounts for 5% of global lithium use (2011).[[30]](#cite_note-30) Lithium compounds are also used as additives (fluxes) to [foundry sand](/wiki/Foundry_sand) for iron casting to reduce veining.[[87]](#cite_note-87) Lithium (as [lithium fluoride](/wiki/Lithium_fluoride)) is used as an additive to aluminium smelters ([Hall–Héroult process](/wiki/Hall–Héroult_process)), reducing melting temperature and increasing electrical resistance,[[88]](#cite_note-88) a use which accounts for 3% of production (2011).[[30]](#cite_note-30) When used as a [flux](/wiki/Flux_(metallurgy)) for [welding](/wiki/Welding) or [soldering](/wiki/Soldering), metallic lithium promotes the fusing of metals during the process[[89]](#cite_note-89) and eliminates the forming of [oxides](/wiki/Oxide) by absorbing impurities.[Template:Citation needed](/wiki/Template:Citation_needed) [Alloys](/wiki/Alloy) of the metal with aluminium, [cadmium](/wiki/Cadmium), copper and [manganese](/wiki/Manganese) are used to make high-performance aircraft parts (see also [Lithium-aluminium alloys](/wiki/Al-Li)).[[90]](#cite_note-90)

### Silicon nano-welding[[edit](/index.php?title=(none)&action=edit&section=26)]

Lithium has been found effective in assisting the perfection of silicon nano-welds in electronic components for electric batteries and other devices.[[91]](#cite_note-91)

### Other chemical and industrial uses[[edit](/index.php?title=(none)&action=edit&section=27)]

[thumb|left||100px|Lithium use in flares and](/wiki/File:FlammenfärbungLi.png) [pyrotechnics](/wiki/Pyrotechnics) is due to its rose-red flame.[[92]](#cite_note-92)

#### Pyrotechnics[[edit](/index.php?title=(none)&action=edit&section=28)]

Lithium compounds are used as [pyrotechnic colorants](/wiki/Pyrotechnic_colorant) and oxidizers in red [fireworks](/wiki/Fireworks) and [flares](/wiki/Flare_(pyrotechnic)).<ref name=CRC/>[[93]](#cite_note-93)

#### Air purification[[edit](/index.php?title=(none)&action=edit&section=29)]

[Lithium chloride](/wiki/Lithium_chloride) and [lithium bromide](/wiki/Lithium_bromide) are [hygroscopic](/wiki/Hygroscopic) and are used as [desiccants](/wiki/Desiccant) for gas streams.<ref name=CRC/> Lithium hydroxide and [lithium peroxide](/wiki/Lithium_peroxide) are the salts most used in confined areas, such as aboard [spacecraft](/wiki/Spacecraft) and [submarines](/wiki/Submarine), for carbon dioxide removal and air purification. Lithium hydroxide absorbs [carbon dioxide](/wiki/Carbon_dioxide) from the air by forming lithium carbonate, and is preferred over other alkaline hydroxides for its low weight.

Lithium peroxide (Li2O2) in presence of moisture not only reacts with carbon dioxide to form lithium carbonate, but also releases oxygen.[[94]](#cite_note-94)[[95]](#cite_note-95) The reaction is as follows:

2 Li2O2 + 2 CO2 → 2 Li2CO3 + O2.

Some of the aforementioned compounds, as well as [lithium perchlorate](/wiki/Lithium_perchlorate), are used in [oxygen candles](/wiki/Chemical_oxygen_generator#Oxygen_candle) that supply [submarines](/wiki/Submarine) with [oxygen](/wiki/Oxygen). These can also include small amounts of [boron](/wiki/Boron), [magnesium](/wiki/Magnesium), [aluminum](/wiki/Aluminum), [silicon](/wiki/Silicon), [titanium](/wiki/Titanium), [manganese](/wiki/Manganese), and [iron](/wiki/Iron).[[96]](#cite_note-96)

#### Optics[[edit](/index.php?title=(none)&action=edit&section=30)]

[Lithium fluoride](/wiki/Lithium_fluoride), artificially grown as [crystal](/wiki/Crystal), is clear and transparent and often used in specialist optics for [IR](/wiki/Infrared), [UV](/wiki/Ultraviolet) and VUV ([vacuum UV](/wiki/Vacuum_UV)) applications. It has one of the lowest [refractive indexes](/wiki/Refractive_index) and the farthest transmission range in the deep UV of most common materials.[[97]](#cite_note-97) Finely divided lithium fluoride powder has been used for [thermoluminescent radiation dosimetry](/wiki/Thermoluminescent_Dosimeter) (TLD): when a sample of such is exposed to radiation, it accumulates [crystal defects](/wiki/Crystal_defect) which, when heated, resolve via a release of bluish light whose intensity is proportional to the [absorbed dose](/wiki/Absorbed_dose), thus allowing this to be quantified.[[98]](#cite_note-98) Lithium fluoride is sometimes used in focal lenses of [telescopes](/wiki/Telescope).<ref name=CRC/>[[99]](#cite_note-99) The high non-linearity of [lithium niobate](/wiki/Lithium_niobate) also makes it useful in [non-linear optics applications](/wiki/Nonlinear_optics). It is used extensively in telecommunication products such as mobile phones and [optical modulators](/wiki/Optical_modulator), for such components as [resonant crystals](/wiki/Crystal_oscillator). Lithium applications are used in more than 60% of mobile phones.[[100]](#cite_note-100)

#### Organic and polymer chemistry[[edit](/index.php?title=(none)&action=edit&section=31)]

[Organolithium compounds](/wiki/Organolithium_compound) are widely used in the production of polymer and fine-chemicals. In the polymer industry, which is the dominant consumer of these reagents, alkyl lithium compounds are [catalysts](/wiki/Catalyst)/[initiators](/wiki/Radical_initiator).[[101]](#cite_note-101) in [anionic polymerization](/wiki/Anionic_addition_polymerization) of [unfunctionalized](/wiki/Functional_group) [olefins](/wiki/Olefin).[[102]](#cite_note-102)[[103]](#cite_note-103)[[104]](#cite_note-104) For the production of fine chemicals, organolithium compounds function as strong bases and as reagents for the formation of [carbon-carbon bonds](/wiki/Carbon-carbon_bond). Organolithium compounds are prepared from lithium metal and alkyl halides.[[105]](#cite_note-105) Many other lithium compounds are used as reagents to prepare organic compounds. Some popular compounds include [lithium aluminium hydride](/wiki/Lithium_aluminium_hydride) (LiAlH4), [lithium triethylborohydride](/wiki/Lithium_triethylborohydride), [n-Butyllithium](/wiki/N-Butyllithium) and [tert-butyllithium](/wiki/Tert-butyllithium) are commonly used as extremely strong bases called [superbase](/wiki/Superbase).

#### Military applications[[edit](/index.php?title=(none)&action=edit&section=32)]

Metallic lithium and its complex [hydrides](/wiki/Hydride), such as [Li[AlH4](/wiki/Lithium_aluminium_hydride)], are used as high-energy additives to [rocket propellants](/wiki/Rocket_propellant).<ref name=emsley/> Lithium aluminum hydride can also be used by itself as a [solid fuel](/wiki/Solid_fuel).[[106]](#cite_note-106) [thumb|The launch of a torpedo using lithium as fuel](/wiki/File:US_Navy_040626-N-5319A-006_An_Anti-Submarine_Warfare_(ASW)_MK-50_Torpedo_is_launched_from_guided_missile_destroyer_USS_Bulkeley_(DDG_84).jpg)

The [Mark 50 torpedo](/wiki/Mark_50_torpedo) stored chemical energy propulsion system (SCEPS) uses a small tank of [sulfur hexafluoride](/wiki/Sulfur_hexafluoride) gas, which is sprayed over a block of solid lithium. The reaction generates heat, creating [steam](/wiki/Steam) to propel the torpedo in a closed [Rankine cycle](/wiki/Rankine_cycle).[[107]](#cite_note-107) [Lithium hydride](/wiki/Lithium_hydride) containing lithium-6 is used in [thermonuclear weapons](/wiki/Thermonuclear_weapon), where it encases the core of the [bomb](/wiki/Nuclear_bomb).[[108]](#cite_note-108)

### Nuclear[[edit](/index.php?title=(none)&action=edit&section=33)]

Lithium-6 is valued as a source material for [tritium](/wiki/Tritium) production and as a [neutron absorber](/wiki/Neutron_absorber) in [nuclear fusion](/wiki/Nuclear_fusion). Natural lithium contains about 7.5% lithium-6 from which large amounts of lithium-6 have been produced by [isotope separation](/wiki/Isotope_separation) for use in [nuclear weapons](/wiki/Nuclear_weapon).[[109]](#cite_note-109) Lithium-7 gained interest for use in [nuclear reactor](/wiki/Nuclear_reactor) [coolants](/wiki/Coolant).[[110]](#cite_note-110) [thumb|Lithium deuteride was used as fuel in the](/wiki/File:Castle_Bravo_Blast.jpg) [Castle Bravo](/wiki/Castle_Bravo) nuclear device. [Lithium deuteride](/wiki/Lithium_deuteride) was the [fusion fuel](/wiki/Nuclear_fusion) of choice in early versions of the [hydrogen bomb](/wiki/Nuclear_weapon). When bombarded by [neutrons](/wiki/Neutron), both 6Li and 7Li produce [tritium](/wiki/Tritium) — this reaction, which was not fully understood when [hydrogen bombs](/wiki/Teller-Ulam_design) were first tested, was responsible for the runaway yield of the [Castle Bravo](/wiki/Castle_Bravo) [nuclear test](/wiki/Nuclear_test). Tritium fuses with [deuterium](/wiki/Deuterium) in a [fusion](/wiki/Nuclear_fusion) reaction that is relatively easy to achieve. Although details remain secret, lithium-6 deuteride apparently still plays a role in modern [nuclear weapons](/wiki/Nuclear_weapons) as a fusion material.[[111]](#cite_note-111) [Lithium fluoride](/wiki/Lithium_fluoride), when highly enriched in the lithium-7 isotope, forms the basic constituent of the fluoride salt mixture LiF-[BeF2](/wiki/Beryllium_fluoride) used in [liquid fluoride nuclear reactors](/wiki/Molten_salt_reactor). Lithium fluoride is exceptionally chemically stable and LiF-BeF2 mixtures have low melting points. In addition, 7Li, Be, and F are among the few [nuclides](/wiki/Nuclide) with low enough [thermal neutron capture cross-sections](/wiki/Neutron_cross-section) not to poison the fission reactions inside a nuclear fission reactor.<ref group=note>Beryllium and fluorine occur only as one isotope, 9Be and 19F respectively. These two, together with 7Li, as well as [2H](/wiki/Deuterium), 11B, 15N, 209Bi, and the stable isotopes of C, and O, are the only nuclides with low enough thermal neutron capture cross sections aside from [actinides](/wiki/Actinide) to serve as major constituents of a molten salt breeder reactor fuel.</ref>[[112]](#cite_note-112) In conceptualized (hypothetical) nuclear [fusion power](/wiki/Fusion_power) plants, lithium will be used to produce tritium in [magnetically confined reactors](/wiki/Magnetic_confinement_fusion) using [deuterium](/wiki/Deuterium) and [tritium](/wiki/Tritium) as the fuel. Naturally occurring tritium is extremely rare, and must be synthetically produced by surrounding the reacting [plasma](/wiki/Plasma_(physics)) with a 'blanket' containing lithium where neutrons from the deuterium-tritium reaction in the plasma will fission the lithium to produce more tritium:

6Li + n → 4He + 3T.

Lithium is also used as a source for [alpha particles](/wiki/Alpha_particle), or [helium](/wiki/Helium) nuclei. When 7Li is bombarded by accelerated [protons](/wiki/Proton) 8[Be](/wiki/Beryllium) is formed, which undergoes fission to form two alpha particles. This feat, called "splitting the atom" at the time, was the first fully man-made [nuclear reaction](/wiki/Nuclear_reaction). It was produced by [Cockroft](/wiki/John_Douglas_Cockcroft) and [Walton](/wiki/Ernest_Walton) in 1932.[[113]](#cite_note-113)[[114]](#cite_note-114) In 2013, the US [Government Accountability Office](/wiki/Government_Accountability_Office) said a shortage of lithium-7 critical to the operation of 65 out of 100 American nuclear reactors “places their ability to continue to provide electricity at some risk”. The problem stems from the decline of US nuclear infrastructure. The equipment needed to separate lithium-6 from lithium-7 is mostly a cold war leftover. The US shut down most of this machinery in 1963, when it had a huge surplus of separated lithium, mostly consumed during the twentieth century. The report said it would take five years and $10 million to $12 million to reestablish the ability to separate lithium-6 from lithium-7.<ref name=nyt1013/>

Reactors that use lithium-7 heat water under high pressure and transfer heat through heat exchangers that are prone to corrosion. The reactors use lithium to counteract the corrosive effects of [boric acid](/wiki/Boric_acid), which is added to the water to absorb excess neutrons.<ref name=nyt1013>[Template:Cite news](/wiki/Template:Cite_news)</ref>

### Medicine[[edit](/index.php?title=(none)&action=edit&section=34)]

[Template:Main](/wiki/Template:Main) Lithium is useful in the treatment of [bipolar disorder](/wiki/Bipolar_disorder).[[115]](#cite_note-115) Lithium salts may also be helpful for related diagnoses, such as [schizoaffective disorder](/wiki/Schizoaffective_disorder) and cyclic [major depression](/wiki/Major_depression). The active part of these salts is the lithium ion Li+.[[115]](#cite_note-115) They may increase the risk of developing [Ebstein's cardiac anomaly](/wiki/Ebstein's_anomaly) in infants born to women who take lithium during the first trimester of pregnancy.[[116]](#cite_note-116) Lithium has also been researched as a possible treatment for [cluster headaches](/wiki/Cluster_headache).[[117]](#cite_note-117)

## Precautions[[edit](/index.php?title=(none)&action=edit&section=35)]

[Template:NFPA 704](/wiki/Template:NFPA_704) Lithium is [corrosive](/wiki/Corrosive) and requires special handling to avoid skin contact. Breathing lithium dust or lithium compounds (which are often alkaline) initially [irritate](/wiki/Irritation) the [nose](/wiki/Human_nose) and throat, while higher exposure can cause a buildup of fluid in the [lungs](/wiki/Lung), leading to [pulmonary edema](/wiki/Pulmonary_edema). The metal itself is a handling hazard because contact with moisture produces the [caustic](/wiki/Corrosive_substance) lithium hydroxide. Lithium is safely stored in non-reactive compounds such as [naphtha](/wiki/Naphtha).[[118]](#cite_note-118)

### Regulation[[edit](/index.php?title=(none)&action=edit&section=36)]

Some jurisdictions limit the sale of [lithium batteries](/wiki/Lithium_battery), which are the most readily available source of lithium for ordinary consumers. Lithium can be used to reduce [pseudoephedrine](/wiki/Pseudoephedrine) and [ephedrine](/wiki/Ephedrine) to [methamphetamine](/wiki/Methamphetamine) in the [Birch reduction](/wiki/Birch_reduction) method, which employs solutions of alkali metals dissolved in anhydrous [ammonia](/wiki/Ammonia).[[119]](#cite_note-119)[[120]](#cite_note-120) Carriage and shipment of some kinds of lithium batteries may be prohibited aboard certain types of transportation (particularly aircraft) because of the ability of most types of lithium batteries to fully discharge very rapidly when [short-circuited](/wiki/Short_circuit), leading to overheating and possible [explosion](/wiki/Explosion) in a process called [thermal runaway](/wiki/Thermal_runaway). Most consumer lithium batteries have built-in thermal overload protection to prevent this type of incident, or are otherwise designed to limit short-circuit currents. Internal shorts from manufacturing defect or physical damage can lead to spontaneous thermal runaway.[[121]](#cite_note-121)[[122]](#cite_note-122)

## See also[[edit](/index.php?title=(none)&action=edit&section=37)]

* [Dilithium](/wiki/Dilithium)
* [List of countries by lithium production](/wiki/List_of_countries_by_lithium_production)
* [Lithium compounds](/wiki/Category:Lithium_compounds)
* [Lithium (medication)](/wiki/Lithium_(medication))
* [Lithium soap](/wiki/Lithium_soap)
* [Lithium-ion battery](/wiki/Lithium-ion_battery)
* [Lithium–air battery](/wiki/Lithium–air_battery)

[Template:Subject bar](/wiki/Template:Subject_bar)

## Notes[[edit](/index.php?title=(none)&action=edit&section=38)]

<references group=note/>

## References[[edit](/index.php?title=(none)&action=edit&section=39)]

[Template:Reflist](/wiki/Template:Reflist)

## External links[[edit](/index.php?title=(none)&action=edit&section=40)]

* [Lithium](http://www.periodicvideos.com/videos/003.htm) at [*The Periodic Table of Videos*](/wiki/The_Periodic_Table_of_Videos) (University of Nottingham)
* [International Lithium Alliance](http://www.lithiumalliance.org/)
* [USGS: Lithium Statistics and Information](http://minerals.usgs.gov/minerals/pubs/commodity/lithium/)
* [Lithium Supply & Markets 2009 IM Conference 2009 Sustainable lithium supplies through 2020 in the face of sustainable market growth](http://trugroup.com/whitepapers/TRU-Lithium-Outlook-2020.pdf)
* [University of Southampton, Mountbatten Centre for International Studies, Nuclear History Working Paper No5.](http://www.mcis.soton.ac.uk/Site_Files/pdf/nuclear_history/Working_Paper_No_5.pdf)

[Template:Compact periodic table](/wiki/Template:Compact_periodic_table) [Template:Lithium compounds](/wiki/Template:Lithium_compounds) [Template:Alkalimetals](/wiki/Template:Alkalimetals)

[Template:Use dmy dates](/wiki/Template:Use_dmy_dates)

[Template:Authority control](/wiki/Template:Authority_control)

[Category:Lithium](/wiki/Category:Lithium) [Category:Alkali metals](/wiki/Category:Alkali_metals) [Category:Chemical elements](/wiki/Category:Chemical_elements) [Category:Reducing agents](/wiki/Category:Reducing_agents)