= Caesium =

Caesium or cesium is a chemical element with symbol Cs and atomic number 55 . It is a soft , silvery @-@ gold alkali metal with a melting point of 28 @.@ 5 ° C (83 @.@ 3 ° F) , which makes it one of only five elemental metals that are liquid at or near room temperature . Caesium is an alkali metal and has physical and chemical properties similar to those of rubidium and potassium . The metal is extremely reactive and pyrophoric , reacting with water even at ? 116 ° C (? 177 ° F) . It is the least electronegative element . It has only one stable isotope , caesium @-@ 133 . Caesium is mined mostly from pollucite , while the radioisotopes , especially caesium @-@ 137 , a fission product , are extracted from waste produced by nuclear reactors .

The German chemist Robert Bunsen and physicist Gustav Kirchhoff discovered caesium in 1860 by the newly developed method of flame spectroscopy . The first small @-@ scale applications for caesium were as a "getter" in vacuum tubes and in photoelectric cells . In 1967 , based on Einstein defining the speed of light as the most constant dimension in the universe , the International System of Units isolated two specific wave counts from an emission spectrum of caesium @-@ 133 to co @-@ define the second and the meter . Since then , caesium has been widely used in highly accurate atomic clocks .

Since the 1990s , the largest application of the element has been as caesium formate for drilling fluids . It has a range of applications in the production of electricity , in electronics , and in chemistry . The radioactive isotope caesium @-@ 137 has a half @-@ life of about 30 years and is used in medical applications , industrial gauges , and hydrology . Although the element is only mildly toxic , it is a hazardous material as a metal and its radioisotopes present a high health risk if released into the environment .

= = Characteristics = =

= = = Physical properties = = =

Caesium is a very soft (it has the lowest hardness of all elements , 0 @.@ 2 Mohs) , very ductile , pale metal , which darkens in the presence of trace amounts of oxygen . It has a melting point of 28 @.@ 4 ° C (83 @.@ 1 ° F) , making it one of the few elemental metals that are liquid near room temperature . Mercury is the only elemental metal with a known melting point lower than caesium . In addition , the metal has a rather low boiling point , 641 ° C (1 @,@ 186 ° F) , the lowest of all metals other than mercury . Its compounds burn with a blue or violet colour .

Caesium forms alloys with the other alkali metals , as well as with gold , and amalgams with mercury . At temperatures below 650 ° C (1 @,@ 202 ° F) , it does not alloy with cobalt , iron , molybdenum , nickel , platinum , tantalum or tungsten . It forms well @-@ defined intermetallic compounds with antimony , gallium , indium and thorium , which are photosensitive . It mixes with all the other alkali metals (except lithium) ; the alloy with a molar distribution of 41 % caesium , 47 % potassium , and 12 % sodium has the lowest melting point of any known metal alloy , at ? 78 ° C (? 108 ° F) . A few amalgams have been studied : CsHg

2 is black with a purple metallic lustre, while CsHg is golden @-@ coloured, also with a metallic lustre.

= = = Chemical properties = = =

Caesium metal is highly reactive and very pyrophoric . In addition to igniting spontaneously in air , it reacts explosively with water even at low temperatures , more so than other members of the first group of the periodic table . The reaction with solid water occurs at temperatures as low as ? 116 ° C (? 177 ° F) . Because of its high reactivity , the metal is classified as a hazardous material . It is stored and shipped in dry saturated hydrocarbons , such as mineral oil . Similarly , it must be handled under inert gas , such as argon . However , a caesium @-@ water explosion is often less

powerful than a sodium @-@ water explosion with a similar amount of sodium . This is because caesium explodes instantly upon contact with water , leaving little time for hydrogen to accumulate . Caesium can be stored in vacuum @-@ sealed borosilicate glass ampoules . In quantities of more than about 100 grams (3 @.@ 5 oz) , caesium is shipped in hermetically sealed , stainless steel containers .

The chemistry of caesium is similar to that of other alkali metals , but is more closely similar to that of rubidium , the element above caesium in the periodic table . As expected for an alkali metal , its only common oxidation state is $+\ 1$. Some small differences arise from the fact that it has a higher atomic mass and is more electropositive than other (nonradioactive) alkali metals . Caesium is the most electropositive chemical element . The caesium ion is also larger and less " hard " than those of the lighter alkali metals .

= = = Compounds = = =

The vast majority of caesium compounds contain the element as the cation Cs + , which binds ionically to a wide variety of anions . One noteworthy exception is provided by the caeside anion (Cs?). Other exceptions include the several suboxides (see section on oxides below).

Returning to more normal compounds , salts of Cs + are almost invariably colourless unless the anion itself is coloured . Many of the simple salts are hygroscopic , but less so than the corresponding salts of the lighter alkali metals . The phosphate , acetate , carbonate , halides , oxide , nitrate , and sulfate salts are water @-@ soluble . Double salts are often less soluble , and the low solubility of caesium aluminium sulfate is exploited in the purification of Cs from its ores . The double salt with antimony (such as CsSbCl

4), bismuth, cadmium, copper, iron, and lead are also poorly soluble.

Caesium hydroxide (CsOH) is hygroscopic and a very strong base . It rapidly etches the surface of semiconductors such as silicon . CsOH has been previously regarded by chemists as the " strongest base " , reflecting the relatively weak attraction between the large Cs + ion and OH?; it is indeed the strongest Arrhenius base , but a number of compounds that cannot exist in aqueous solution , such as n @-@ butyllithium and sodium amide , are more basic .

A stoichiometric mixture of caesium and gold will react to form yellow caesium auride (Cs + Au?) upon heating. The auride anion here behaves as a pseudohalogen. The compound reacts violently with water, yielding caesium hydroxide, metallic gold, and hydrogen gas; in liquid ammonia it can be reacted with a caesium @-@ specific ion exchange resin to produce tetramethylammonium auride. The analogous platinum compound, the red caesium platinide (Cs2Pt), contains the platinide ion that behaves as a pseudochalcogen.

= = = = Complexes = = =

Like all metal cations , Cs + forms complexes with Lewis bases in solution . Because of its large size , Cs + usually adopts coordination numbers greater than six @-@ coordination , which is typical for the lighter alkali metal cations . This trend is already apparent by the 8 @-@ coordination in CsCl , vs. the halite motif adopted by the other alkali metal chlorides . Its high coordination number and softness (tendency to form covalent bonds) are the basis of the separation of Cs + from other cations , as is practiced in the remediation of nuclear wastes , where 137Cs + is separated from large amounts of nonradioactive K + .

= = = = Halides = = =

Caesium fluoride (CsF) is a hygroscopic white solid that is widely used in organofluorine chemistry as a source of the fluoride anion . Caesium fluoride has the halite structure , which means that the Cs + and F ? pack in a cubic closest packed array as do Na + and Cl ? in sodium chloride . It is noteworthy as caesium and fluorine have the lowest and highest electronegativities respectively among all the known elements .

Caesium chloride (CsCl) crystallizes in the simple cubic crystal system . Also called the " caesium chloride structure " , this structural motif is composed of a primitive cubic lattice with a two @-@ atom basis , each with an eightfold coordination ; the chloride atoms lie upon the lattice points at the edges of the cube , while the caesium atoms lie in the holes in the center of the cubes . This structure is shared with CsBr and Csl , and many other compounds that do not contain Cs . In contrast , most other alkaline halides adopt the sodium chloride (NaCl) structure . The CsCl structure is preferred because Cs + has an ionic radius of 174 pm and Cl ? 181 pm .

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= = = = Oxides = = = = =
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More so than the other alkali metals, caesium forms numerous binary compounds with oxygen. When caesium burns in air, the superoxide CsO

2 is the main product . The "normal "caesium oxide (Cs

2O) forms yellow @-@ orange hexagonal crystals , and is the only oxide of the anti @-@ CdCl 2 type . It vaporizes at 250 $^{\circ}$ C (482 $^{\circ}$ F) , and decomposes to caesium metal and the peroxide Cs 2O

2 at temperatures above 400 $^{\circ}$ C (752 $^{\circ}$ F) . Aside from the superoxide and the ozonide CsO 3 , several brightly coloured suboxides have also been studied . These include Cs

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70 , Cs
40 , Cs
110
3 , Cs
30 (dark @-@ green), CsO, Cs
30
2 , as well as Cs
70
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2. The latter may be heated under vacuum to generate Cs

20 . Binary compounds with sulfur, selenium, and tellurium also exist.

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Caesium has a total of 39 known isotopes that range in their mass number (i.e. number of nucleons in its nucleus) from 112 to 151 . Several of these are synthesized from lighter elements by the slow neutron capture process (S @-@ process) inside old stars , as well as inside supernova explosions (R @-@ process) . However , the only stable caesium isotope is 133Cs , which has 78 neutrons . Although it has a large nuclear spin (7 / 2 +) , nuclear magnetic resonance studies can be done with this isotope at a resonating frequency of 11 @.@ 7 MHz .

The radioactive 135Cs has a very long half @-@ life of about 2 @.@ 3 million years, longest of all radioactive isotopes of caesium. 137Cs and 134Cs have half @-@ lives of 30 and two years, respectively. 137Cs decomposes to a short @-@ lived 137mBa by beta decay, and then to nonradioactive barium, while 134Cs transforms into 134Ba directly. The isotopes with mass numbers of 129, 131, 132 and 136, have half @-@ lives between a day and two weeks, while most of the other isotopes have half @-@ lives from a few seconds to fractions of a second. There are at least 21 metastable nuclear isomers. Other than 134mCs (with a half @-@ life of just under 3 hours), all are very unstable and decay with half @-@ lives of a few minutes or less.

The isotope 135Cs is one of the long @-@ lived fission products of uranium which form in nuclear reactors. However, its fission product yield is reduced in most reactors because its predecessor, 135Xe, is an extremely potent neutron poison and transmutes frequently to stable 136Xe before it can decay to 135Cs.

Because of its beta decay (to 137mBa) , 137Cs is a strong emitter of gamma radiation . Its half @-@ life makes it the principal medium @-@ lived fission product along with 90Sr ? both are responsible for radioactivity of spent nuclear fuel after several years of cooling up to several hundred years after use . For example , 137Cs together with 90Sr currently generate the largest source of

radioactivity generated in the area around the Chernobyl disaster. It is not feasible to dispose of 137Cs through neutron capture (due to the low capture rate) and as a result it must be allowed to decay.

Almost all caesium produced from nuclear fission comes from beta decay of originally more neutron @-@ rich fission products , passing through various isotopes of iodine and of xenon . Because iodine and xenon are volatile and can diffuse through nuclear fuel or air , radioactive caesium is often created far from the original site of fission . With the commencement of nuclear weapons testing around 1945 , 137Cs was released into the atmosphere and then returned to the surface of the earth as a component of radioactive fallout .

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= = = Occurrence = = =
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Caesium is a relatively rare element as it is estimated to average approximately 3 parts per million in the Earth 's crust . This makes it the 45th most abundant of all elements and the 36th of all the metals . Nevertheless , it is more abundant than such elements as antimony , cadmium , tin and tungsten , and two orders of magnitude more abundant than mercury or silver , but 3 @.@ 3 % as abundant as rubidium ? with which it is so closely associated , chemically .

Due to its large ionic radius , caesium is one of the "incompatible elements". During magma crystallization , caesium is concentrated in the liquid phase and crystallizes last . Therefore , the largest deposits of caesium are zone pegmatite ore bodies formed by this enrichment process . Because caesium does not substitute for potassium as readily as does rubidium , the alkali evaporite minerals sylvite (KCI) and carnallite (KMgCI

3 · 6H

20) may contain only 0 @.@ 002 % caesium. Consequently, Cs is found in few minerals. Percentage amounts of caesium may be found in beryl (Be

3AI

2 (SiO

3)

6) and avogadrite ((K, Cs)BF

4), up to 15 wt % Cs2O in the closely related mineral pezzottaite (Cs (Be2Li) Al2Si6O18), up to 8 @.@ 4 wt % Cs2O in the rare mineral londonite ((Cs, K) Al

4Be

4 (B, Be)

120

28), and less in the more widespread rhodizite. The only economically important source mineral for caesium is pollucite Cs (AlSi

20

6), which is found in a few places around the world in zoned pegmatites, and is associated with the more commercially important lithium minerals lepidolite and petalite. Within the pegmatites, the large grain size and the strong separation of the minerals create high @-@ grade ore for mining.

One of the world 's most significant and richest sources of the metal is the Tanco Mine at Bernic Lake in Manitoba , Canada . The deposits there are estimated to contain 350 @,@ 000 metric tons of pollucite ore , which represent more than two @-@ thirds of the world 's reserve base . Although the stoichiometric content of caesium in pollucite is 42 @.@ 6 % , pure pollucite samples from this deposit contain only about 34 % caesium , while the average content is 24 wt % . Commercial pollucite contains over 19 % caesium . The Bikita pegmatite deposit in Zimbabwe is mined for its petalite , but it also contains a significant amount of pollucite . Notable amounts of pollucite are also mined in the Karibib Desert , Namibia . At the present rate of world mine production of 5 to 10 metric tons per year , reserves will last for thousands of years .

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= = Production = =
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The mining of pollucite ore is a selective process and is conducted on a small scale in comparison

with most metal mining operations. The ore is crushed, hand @-@ sorted, but not usually concentrated, and then ground. Caesium is then extracted from pollucite mainly by three methods: acid digestion, alkaline decomposition, and direct reduction.

In the acid digestion , the silicate pollucite rock is dissolved with strong acids , such as hydrochloric (HCl) , sulfuric (H

2SO

4), hydrobromic (HBr), or hydrofluoric (HF) acids. With hydrochloric acid, a mixture of soluble chlorides is produced, and the insoluble chloride double salts of caesium are precipitated as caesium antimony chloride (Cs

4SbCI

7), caesium iodine chloride (Cs

2ICI), or caesium hexachlorocerate (Cs

2 (CeCl

6)). After separation, the pure precipitated double salt is decomposed, and pure CsCl is obtained after evaporating the water. The method using sulfuric acid yields the insoluble double salt directly as caesium alum (CsAl (SO

4)

2 · 12H

20). The aluminium sulfate in it is converted to the insoluble aluminium oxide by roasting the alum with carbon, and the resulting product is leached with water to yield a Cs

2SO

4 solution.

The roasting of pollucite with calcium carbonate and calcium chloride yields insoluble calcium silicates and soluble caesium chloride. Leaching with water or dilute ammonia (NH

4OH) yields then a dilute chloride (CsCl) solution . This solution can be evaporated to produce caesium chloride or transformed into caesium alum or caesium carbonate . Albeit not commercially feasible , direct reduction of the ore with potassium , sodium or calcium in vacuum can produce caesium metal directly .

Most of the mined caesium (as salts) is directly converted into caesium formate (HCOO ? Cs +) for applications such as oil drilling . To supply the developing market , Cabot Corporation built a production plant in 1997 at the Tanco mine near Bernic Lake in Manitoba , with a capacity of 12 @,@ 000 barrels (1 @,@ 900 m3) per year of caesium formate solution . The primary smaller @-@ scale commercial compounds of caesium are caesium chloride and its nitrate .

Alternatively , caesium metal may be obtained from the purified compounds derived from the ore . Caesium chloride , and the other caesium halides , as well , can be reduced at 700 to 800 $^{\circ}$ C (1 @,@ 292 to 1 @,@ 472 $^{\circ}$ F) with calcium or barium , followed by distillation of the caesium metal . In the same way , the aluminate , carbonate , or hydroxide may be reduced by magnesium . The metal can also be isolated by electrolysis of fused caesium cyanide (CsCN) . Exceptionally pure and gas @-@ free caesium can be made by the thermal decomposition at 390 $^{\circ}$ C (734 $^{\circ}$ F) of caesium azide CsN

3, which is produced from aqueous caesium sulfate and barium azide. In vacuum applications, caesium dichromate can be reacted with zirconium forming pure caesium metal without other gaseous products.

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Cs
2Cr
2O
7 + 2 Zr ? 2 Cs + 2 ZrO
2 + Cr
2O
3
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The price of 99 @.@ 8 % pure caesium (metal basis) in 2009 was about US \$ 10 per gram (\$ 280 per ounce), but its compounds are significantly cheaper.

In 1860 , Robert Bunsen and Gustav Kirchhoff discovered caesium in the mineral water from Dürkheim , Germany . Due to the bright blue lines in its emission spectrum , they chose a name derived from the Latin word caesius , meaning sky @-@ blue . Caesium was the first element to be discovered spectroscopically , only one year after the invention of the spectroscope by Bunsen and Kirchhoff .

To obtain a pure sample of caesium , 44 @,@ 000 litres (9 @,@ 700 imp gal ; 12 @,@ 000 US gal) of mineral water had to be evaporated to yield 240 kilograms (530 lb) of concentrated salt solution . The alkaline earth metals were precipitated either as sulfates or oxalates , leaving the alkali metal in the solution . After conversion to the nitrates and extraction with ethanol , a sodium @-@ free mixture was obtained . From this mixture , the lithium was precipitated by ammonium carbonate . Potassium , rubidium and caesium form insoluble salts with chloroplatinic acid , but these salts show a slight difference in solubility in hot water . Therefore , the less @-@ soluble caesium and rubidium hexachloroplatinate ((Cs , Rb) 2PtCl6) could be obtained by fractional crystallization . After reduction of the hexachloroplatinate with hydrogen , caesium and rubidium could be separated by the difference in solubility of their carbonates in alcohol . The process yielded 9 @.@ 2 grams (0 @.@ 32 oz) of rubidium chloride and 7 @.@ 3 grams (0 @.@ 26 oz) of caesium chloride from the initial 44 @,@ 000 liters of mineral water .

The two scientists used the caesium chloride thus obtained to estimate the atomic weight of the new element at 123 @.@ 35 (compared to the currently accepted one of 132 @.@ 9) . They tried to generate elemental caesium by electrolysis of molten caesium chloride , but instead of a metal , they obtained a blue homogeneous substance which " neither under the naked eye nor under the microscope " showed the slightest trace of metallic substance " ; as a result , they assigned it as a subchloride (Cs

2CI) . In reality , the product was probably a colloidal mixture of the metal and caesium chloride . The electrolysis of the aqueous solution of chloride with a mercury anode produced a caesium amalgam which readily decomposed under the aqueous conditions . The pure metal was eventually isolated by the German chemist Carl Setterberg while working on his doctorate with Kekulé and Bunsen . In 1882 , he produced caesium metal by electrolysing caesium cyanide , and thus avoiding the problems with the chloride .

Historically , the most important use for caesium has been in research and development , primarily in chemical and electrical fields . Very few applications existed for caesium until the 1920s , when it came to be used in radio vacuum tubes . It had two functions ; as a getter , it removed excess oxygen after manufacture , and as a coating on the heated cathode , it increased its electrical conductivity . Caesium did not become recognized as a high @-@ performance industrial metal until the 1950s . Applications of nonradioactive caesium included photoelectric cells , photomultiplier tubes , optical components of infrared spectrophotometers , catalysts for several organic reactions , crystals for scintillation counters , and in magnetohydrodynamic power generators . Caesium also was , and still is , used as a source of positive ions in secondary ion mass spectrometry (SIMS) .

Since 1967, the International System of Measurements has based its unit of time, the second, on the properties of caesium. The International System of Units (SI) defines the second as the duration of 9 @,@ 192 @,@ 631 @,@ 770 cycles at the microwave frequency of the spectral line corresponding to the transition between two hyperfine energy levels of the ground state of caesium @-@ 133. The 13th General Conference on Weights and Measures of 1967 defined a second as: "the duration of 9 @,@ 192 @,@ 631 @,@ 770 cycles of microwave light absorbed or emitted by the hyperfine transition of caesium @-@ 133 atoms in their ground state undisturbed by external fields "

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= = Applications = = = = = = = Petroleum exploration = = = =
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The largest current end @-@ use of nonradioactive caesium is in caesium formate @-@ based drilling fluids for the extractive oil industry . Aqueous solutions of caesium formate (HCOO ? Cs +) ? made by reacting caesium hydroxide with formic acid ? were developed in the mid @-@ 1990s for use as oil well drilling and completion fluids . The function of a drilling fluid is to lubricate drill bits , to bring rock cuttings to the surface , and to maintain pressure on the formation during drilling of the well . Completion fluids assist the emplacement of control hardware after drilling but prior to production by maintaining the pressure .

The high density of the caesium formate brine (up to 2 @.@ 3 g \cdot cm ? 3 , or 19 @.@ 2 pounds per gallon) , coupled with the relatively benign nature of most caesium compounds , reduces the requirement for toxic high @-@ density suspended solids in the drilling fluid ? a significant technological , engineering and environmental advantage . Unlike the components of many other heavy liquids , caesium formate is relatively environment @-@ friendly . The caesium formate brine can be blended with potassium and sodium formates to decrease the density of the fluids down to that of water (1 @.@ 0 g \cdot cm ? 3 , or 8 @.@ 3 pounds per gallon) . Furthermore , it is biodegradable and reclaimable , and may be recycled , which is important in view of its high cost (about \$ 4 @,@ 000 per barrel in 2001) . Alkali formates are safe to handle and do not damage the producing formation or downhole metals as corrosive alternative , high @-@ density brines (such as zinc bromide ZnBr

2 solutions) sometimes do; they also require less cleanup and reduce disposal costs.

= = = Atomic clocks = = =

Caesium @-@ based atomic clocks observe electromagnetic transitions in the hyperfine structure of caesium @-@ 133 atoms and use it as a reference point . The first accurate caesium clock was built by Louis Essen in 1955 at the National Physical Laboratory in the UK . They have been improved repeatedly over the past half @-@ century , and form the basis for standards @-@ compliant time and frequency measurements , and have been regarded as " the most accurate realization of a unit that mankind has yet achieved . " These clocks measure frequency with an error of 2 to 3 parts in 1014 , which would correspond to a time measurement accuracy of 2 nanoseconds per day , or one second in 1 @.@ 4 million years . The latest versions are accurate to better than 1 part in 1015 , which means they would be off by about 1 second in 20 million years , . Caesium clocks are also used in networks that oversee the timing of cell phone transmissions and the information flow on the Internet .

= = = Electric power and electronics = = =

Caesium vapour thermionic generators are low @-@ power devices that convert heat energy to electrical energy. In the two @-@ electrode vacuum tube converter, it neutralizes the space charge that builds up near the cathode, and in doing so, it enhances the current flow.

Caesium is also important for its photoemissive properties by which light energy is converted to electron flow . It is used in photoelectric cells because caesium @-@ based cathodes , such as the intermetallic compound K

2CsSb , have low threshold voltage for emission of electrons . The range of photoemissive devices using caesium include optical character recognition devices , photomultiplier tubes , and video camera tubes . Nevertheless , germanium , rubidium , selenium , silicon , tellurium , and several other elements can substitute caesium in photosensitive materials .

Caesium iodide (CsI) , bromide (CsBr) and caesium fluoride (CsF) crystals are employed for scintillators in scintillation counters widely used in mineral exploration and particle physics research , as they are well @-@ suited for the detection of gamma and X @-@ ray radiation . Caesium , being a heavy element , provides good stopping power , contributing to better detectivity . Caesium compounds may also provide a faster response (CsF) and be less hygroscopic (CsI) .

Caesium vapor is used in many common magnetometers . The element is also used as an internal

standard in spectrophotometry. Like other alkali metals, caesium has a great affinity for oxygen and is used as a "getter" in vacuum tubes. Other uses of the metal include high @-@ energy lasers, vapour glow lamps, and vapor rectifiers.

= = = Centrifugation fluids = = =

Because of their high density , solutions of caesium chloride , caesium sulfate , and caesium trifluoroacetate (Cs (O 2CCF

3)) are commonly used in molecular biology for density gradient ultracentrifugation . This technology is primarily applied to the isolation of viral particles , subcellular organelles and fractions , and nucleic acids from biological samples .

= = = Chemical and medical use = = =

There are relatively few chemical applications for caesium. Doping with caesium compounds is used to enhance the effectiveness of several metal @-@ ion catalysts used in the production of chemicals, such as acrylic acid, anthraquinone, ethylene oxide, methanol, phthalic anhydride, styrene, methyl methacrylate monomers, and various olefins. It is also used in the catalytic conversion of sulfur dioxide into sulfur trioxide in the production of sulfuric acid.

Caesium fluoride enjoys niche use in organic chemistry as a base , or as an anhydrous source of fluoride ion . Caesium salts sometimes replace potassium or sodium salts in organic synthesis , such as cyclization , esterification , and polymerization . It has also been used in thermoluminescent radiation dosimetry (TLD) : When exposed to radiation , it acquires crystal defects that , when heated , revert with emission of light proportionate to the received dose . Thus , measuring the light pulse with a photomultiplier tube can allow the accumulated radiation dose to be quantified .

= = = Nuclear and isotope applications = = =

Caesium @-@ 137 is a very common radioisotope used as a gamma @-@ emitter in industrial applications. Its advantages include a half @-@ life of roughly 30 years, its availability from the nuclear fuel cycle, and having 137Ba as a stable end product. The high water solubility is a disadvantage which makes it incompatible with large pool irradiators for food and medical supplies. It has been used in agriculture, cancer treatment, and the sterilization of food, sewage sludge, and surgical equipment. Radioactive isotopes of caesium in radiation devices were used in the medical field to treat certain types of cancer, but emergence of better alternatives and the use of water @-@ soluble caesium chloride in the sources, which could create wide @-@ ranging contamination, gradually put some of these caesium sources out of use. Caesium @-@ 137 has been employed in a variety of industrial measurement gauges, including moisture, density, leveling, and thickness gauges. It has also been used in well logging devices for measuring the electron density of the rock formations, which is analogous to the bulk density of the formations.

Isotope 137 has also been used in hydrologic studies analogous to those using tritium . It is a daughter product of nuclear fission reactions . With the commencement of nuclear testing around 1945 , and continuing through the mid @-@ 1980s , caesium @-@ 137 was released into the atmosphere , where it is absorbed readily into solution . Known year @-@ to @-@ year variation within that period allows correlation with soil and sediment layers . Caesium @-@ 134 , and to a lesser extent caesium @-@ 135 , have also been used in hydrology as a measure of caesium output by the nuclear power industry . While they are less prevalent than either caesium @-@ 133 or caesium @-@ 137 , these isotopes have the advantage of being produced solely from anthropogenic sources .

= = = Other uses = = =

Caesium and mercury were used as a propellant in early ion engines designed for spacecraft propulsion on very long interplanetary or extraplanetary missions. The ionization method was to strip the outer electron from the propellant upon contact with a tungsten electrode that had voltage applied. Concerns about the corrosive action of caesium on spacecraft components have pushed development in the direction of the use of inert gas propellants, such as xenon; this is easier to handle in ground @-@ based tests and has less potential to interfere with the spacecraft. Eventually, xenon was used in the experimental spacecraft Deep Space 1 launched in 1998. Nevertheless, Field Emission Electric Propulsion thrusters which use a simple system of accelerating liquid metal ions such as of caesium to create thrust have been built.

Caesium nitrate is used as an oxidizer and pyrotechnic colorant to burn silicon in infrared flares , such as the LUU @-@ 19 flare , because it emits much of its light in the near infrared spectrum . Caesium has been used to reduce the radar signature of exhaust plumes in the SR @-@ 71 Blackbird military aircraft . Caesium , along with rubidium , has been added as a carbonate to glass because it reduces electrical conductivity and improves stability and durability of fibre optics and night vision devices . Caesium fluoride or caesium aluminium fluoride are used in fluxes formulated for the brazing of aluminium alloys that contain magnesium .

Magnetohydrodynamic (MHD) power @-@ generating systems were researched, but failed to gain widespread acceptance. Caesium metal has also been considered as the working fluid in high @-@ temperature Rankine cycle turboelectric generators. Caesium salts have been evaluated as antishock reagents to be used following the administration of arsenical drugs. Because of their effect on heart rhythms, however, they are less likely to be used than potassium or rubidium salts. They have also been used to treat epilepsy.

= = Health and safety hazards = =

Nonradioactive caesium compounds are only mildly toxic. Exposure to large amounts can cause hyperirritability and spasms, due to the chemical similarity of caesium to potassium, but such amounts would not ordinarily be encountered in natural sources and nonradioactive caesium is not a significant environmental hazard. The median lethal dose (LD50) value for caesium chloride in mice is 2 @.@ 3 g per kilogram, which is comparable to the LD50 values of potassium chloride and sodium chloride. However, in large quantities caesium competes with potassium in various biological processes. Accordingly, excess caesum intake can lead to hypokalemia, arrythmia, and acute cardiac arrest. The principal use of nonradioactive caesium, as caesium formate in petroleum drilling fluids, takes advantage of its low toxicity compared to less costly alternatives.

Caesium metal is one of the most reactive elements and is highly explosive when it comes in contact with water . The hydrogen gas produced by the reaction is heated by the thermal energy released at the same time , causing ignition and a violent explosion . This can occur with other alkali metals , but caesium is so potent that this explosive reaction can even be triggered by cold water . The autoignition temperature of caesium is also ? 116 ° C , so it is highly pyrophoric , and ignites explosively in air to form caesium hydroxide and various oxides . Caesium hydroxide is a very strong base , and will rapidly corrode glass .

The isotopes 134 and 137 are present in the biosphere in small amounts from human activities and which differs between locations . Radiocaesium does not accumulate in the body as effectively as many other fission products (such as radioiodine and radiostrontium) . About 10 % of absorbed radiocaesium washes out of the body relatively quickly in sweat and urine . The remaining 90 % has a biological half @-@ life between 50 and 150 days . Radiocaesium follows potassium and tends to accumulate in plant tissues , including fruits and vegetables . Plants absorb caesium differently , some do not absorb it much , and some take it large amounts , sometimes displaying great resistance to it . It is also well @-@ documented that mushrooms from contaminated forests accumulate radiocaesium (caesium @-@ 137) in their fungal sporocarps . Accumulation of caesium @-@ 137 in lakes has been a high concern after the Chernobyl disaster . Experiments with dogs showed that a single dose of 3 @.@ 8 millicuries (140 MBq , 4 @.@ 1 ?g of caesium @-@ 137) per kilogram is lethal within three weeks ; smaller amounts may cause infertility and cancer .

The International Atomic Energy Agency and other sources have warned that radioactive materials, such as caesium @-@ 137, could be used in radiological dispersion devices, or " dirty bombs ".