

Aerogel

en.wikipedia.org/wiki/Aerogel

Aerogel does not have a designated material with set chemical formula but the term is used to group all the material with a certain geometric structure.^[7]

Properties

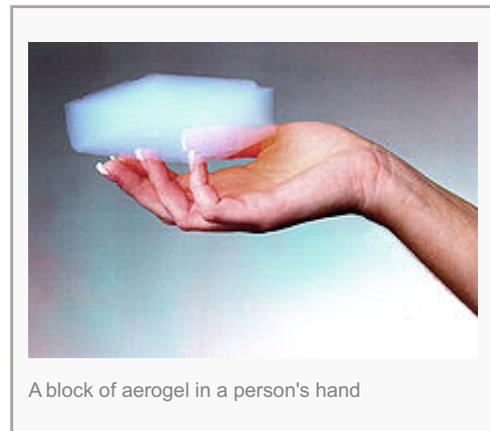
Despite their name, aerogels are solid, rigid, and dry materials that do not resemble a gel in their physical properties: The name comes from the fact that they are made from gels. Pressing softly on an aerogel typically does not leave even a minor mark; pressing more firmly will leave a permanent depression. Pressing extremely firmly will cause a catastrophic breakdown in the sparse structure, causing it to shatter like glass – a property known as *friability*; although more modern variations do not suffer from this. Despite the fact that it is prone to shattering, it is very strong structurally. Its impressive load bearing abilities are due to the *dendritic* microstructure, in which *spherical* particles of average size (2–5 nm) are fused together into clusters. These clusters form a three-dimensional highly *porous* structure of almost *fractal* chains, with pores just under 100 nm. The average size and density of the pores can be controlled during the manufacturing process.

Aerogel is a material that is 98.2% air. The lack of solid material allows aerogel to be almost weightless. The reason for the difference in the composition is the structure of the aerogel. Aerogel has a porous solid network that contains air pockets, with the air pockets taking up majority of space within the material.^[8]

Aerogels are good *thermal insulators* because they almost nullify two of the three methods of *heat transfer* (convection, conduction, and radiation). They are good *conductive* insulators because they are composed almost entirely of gas, and gases are very poor heat conductors. (Silica aerogel is especially good because silica is also a poor conductor of heat; a metallic aerogel, on the other hand, would be less effective.) They are good *convective* inhibitors because air cannot circulate through the lattice. Aerogels are poor *radiative* insulators because infrared radiation (which transfers heat) passes through them.

Owing to its *hygroscopic* nature, aerogel feels dry and acts as a strong *desiccant*. People handling aerogel for extended periods should wear gloves to prevent the appearance of dry brittle spots on their skin.

The slight color it does have is due to *Rayleigh scattering* of the shorter *wavelengths* of *visible light* by the nano-sized dendritic structure. This causes it to appear smoky blue against dark backgrounds and yellowish against bright backgrounds.



A block of aerogel in a person's hand



A flower is on a piece of aerogel which is suspended over a flame from a *Bunsen burner*. Aerogel has excellent insulating properties, and the flower is protected from the flame.

Aerogel is an open porous network. The difference between an open porous network and a closed porous network is that an open porous network allows gases to enter and leave the substance without any limitation, while a closed porous network traps the gases within the material. [9]

Aerogels by themselves are [hydrophilic](#), but chemical treatment can make them [hydrophobic](#). If they absorb moisture they usually suffer a structural change, such as contraction, and deteriorate, but degradation can be prevented by making them hydrophobic. Aerogels with hydrophobic interiors are less susceptible to degradation than aerogels with only an outer hydrophobic layer, even if a crack penetrates the surface. Hydrophobic treatment facilitates processing because it allows the use of a [water jet cutter](#).

Knudsen effect

Aerogels may have a [thermal conductivity](#) smaller than the gas they contain. This is caused by the [Knudsen effect](#). Knudsen effect is the reduction of thermal conductivity in gases when the size of the cavity encompassing the gas becomes comparable to the [mean free path](#). Effectively, the cavity restricts the movement of the gas particles, decreasing the thermal conductivity in addition to eliminating convection. For example, thermal conductivity of air is about 25 mW/m·K at STP and in a large container, but decreases to about 5 mW/m·K in a pore 30 nanometers in diameter. [10]

Structure

Aerogel structure is the result from a [sol-gel polymerization](#), which is when [monomers](#) (simple molecules) reacts with other monomers to form a sol or a substance that consists of bonded, cross-linked [macromolecules](#) with deposits of liquid solution between them. When the material is critically heated the liquid is [evaporated](#) out and the bonded, [cross-linked](#) macromolecule frame is left behind. The result of the polymerization and critical heating is the creation of a material that has a porous strong structure classified as aerogel. [11]

Waterproofing

Aerogel contains particles that are 2-5 nm in diameter. After the process of creating aerogel, it will contain a large amount of hydroxyl groups on the surface. The [hydroxyl groups](#) can cause a strong reaction when placing it in water. The aerogel will catastrophically dissolve in the water. One way to waterproof the [hydrophilic](#) aerogel is by soaking the aerogel with some chemical base that will replace hydroxyl groups with non-polar groups on the surface, the non-polar groups (-OR) is most effective when R is an [aliphatic](#) group. [12]

Porosity of aerogel

There are several ways to determine the porosity of aerogel; the three main methods are gas adsorption, Mercury Porosimetry, and Scattering Method. In gas adsorption nitrogen at its boiling point is adsorbed into the aerogel sample. The gas being absorbed is dependent on the size of the pores within the sample and on the partial pressure of the gas relative to its [saturation pressure](#). Measure the volume of the gas adsorbed by using the Brunauer, Emmet and, Teller formula (BET) gives the specific [surface area](#) of the sample. At high partial pressure in the adsorption/desorption the Kelvin equation gives the pore size distribution of the sample. In Mercury Porosimetry, the [mercury](#) is forced into the aerogel porous system to determine the pores size, but this method is highly inefficient since the solid frame of aerogel will collapse from the high compressive force. The Scattering Methods involves the angle dependent deflection of radiation within the aerogel sample. The sample can be solid particles or pores. The radiation goes into the material and determines the fractal geometry of the aerogel pore network. The best radiation wavelengths

to use are X-rays and neutrons. Aerogel is also an open porous network, the difference between an open porous network and a closed porous network is that in the open network, gasses can enter and leave the substance without any limitation. While a closed porous network traps the gases within the material forcing it to stay within the pores. [13]

Materials

Silica

Silica aerogel is the most common type of aerogel, and the most extensively studied and used. It is [silica](#)-based, derived from [silica gel](#). The lowest-density silica nanofoam weighs $1,000 \text{ g/m}^3$, [14] which is the evacuated version of the record-aerogel of $1,900 \text{ g/m}^3$. [15] The density of [air](#) is $1,200 \text{ g/m}^3$ (at $20 \text{ }^\circ\text{C}$ and 1 atm). [16] As of 2013, [aerographene](#) had a lower density at 160 g/m^3 , or 13% the density of air at room temperature. [17]

The silica solidifies into three-dimensional, intertwined clusters that comprise only 3% of the volume. Conduction through the solid is therefore very low. The remaining 97% of the volume is composed of air in extremely small nanopores. The air has little room to move, inhibiting both convection and gas-phase conduction. [18]

It has remarkable thermal insulative properties, having an extremely low [thermal conductivity](#): from $0.03 \text{ W/m}\cdot\text{K}$ [19] in atmospheric pressure down to $0.004 \text{ W/m}\cdot\text{K}$ [14] in modest vacuum, which correspond to [R-values](#) of 14 to 105 (US customary) or 3.0 to 22.2 (metric) for 3.5 in (89 mm) thickness. For comparison, typical wall insulation is 13 (US customary) or 2.7 (metric) for the same thickness. Its [melting point](#) is $1,473 \text{ K}$ ($1,200 \text{ }^\circ\text{C}$; $2,192 \text{ }^\circ\text{F}$).

Carbon

[Carbon](#) aerogels are composed of particles with sizes in the [nanometer](#) range, [covalently bonded](#) together. They have very high [porosity](#) (over 50%, with pore diameter under 100 nm) and surface areas ranging between $400\text{--}1,000 \text{ m}^2/\text{g}$. They are often manufactured as composite paper: non-woven paper made of [carbon fibers](#), impregnated with [resorcinol–formaldehyde](#) aerogel, and [pyrolyzed](#). Depending on the density, carbon aerogels may be electrically conductive, making composite aerogel paper useful for electrodes in [capacitors](#) or deionization electrodes. Due to their extremely high surface area, carbon aerogels are used to create [supercapacitors](#), with values ranging up to thousands of [farads](#) based on a capacitance density of 104 F/g and 77 F/cm^3 . Carbon aerogels are also extremely "black" in the infrared spectrum, reflecting only 0.3% of radiation between 250 nm and $14.3 \text{ }\mu\text{m}$, making them efficient for [solar energy](#) collectors.

The term "aerogel" to describe airy masses of [carbon nanotubes](#) produced through certain [chemical vapor deposition](#) techniques is incorrect. Such materials can be spun into fibers with strength greater than [Kevlar](#), and unique electrical properties. These materials are not aerogels, however, since they do not have a monolithic internal structure and do not have the regular pore structure characteristic of aerogels.

Metal oxide



[Metal oxide](#) aerogels are used as catalysts in various chemical reactions/transformations or as precursors for other materials.

Aerogels made with [aluminium oxide](#) are known as alumina aerogels. These aerogels are used as catalysts, especially when "doped" with a metal other than aluminium. [Nickel](#)–alumina aerogel is the most common combination. Alumina aerogels are also being considered by [NASA](#) for capturing hypervelocity particles; a formulation doped with [gadolinium](#) and [terbium](#) could [fluoresce](#) at the particle impact site, with the amount of fluorescence dependent on impact energy.

One of the most notable difference between silica aerogels and metal oxide aerogel is that metal oxide aerogels are often variedly colored.

Aerogel	Color
Silica, Alumina, Titania, Zirconia	Clear with Rayleigh scattering blue or white
Iron Oxide	Rust red or yellow, opaque
Chromia	Deep green or deep blue, opaque
Vanadia	Olive green, opaque
Neodymium Oxide	Purple, transparent
Samarium Oxide	Yellow, transparent
Holmium Oxide, Erbium Oxide	Pink, transparent

[23]

Other

Organic polymers can be used to create aerogels. [SEAgel](#) is made of [agar](#). Cellulose from plants can be used to create a flexible aerogel.^[24]

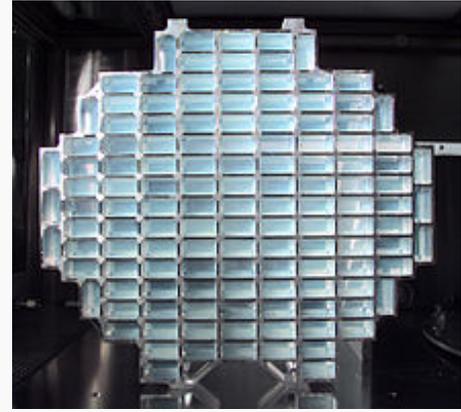
[Chalcogel](#) is an aerogel made of [chalcogens](#) (the column of elements on the periodic table beginning with oxygen) such as sulfur, selenium and other elements.^[25] Metals less expensive than platinum have been used in its creation.

Aerogel performance may be augmented for a specific application by the addition of [dopants](#), reinforcing structures and hybridizing compounds. Aspen Aerogels makes products such as Spaceloft^[27] which are composites of aerogel with some kind of fibrous batting.^[28]

Applications

Aerogels are used for a variety of applications:

- In 2004 about US\$25 million of aerogel insulation product were sold, which had risen to about US\$500 million by 2013. This represents the most substantial economic impact of these materials today. The potential to replace conventional insulation by aerogel solutions in the building and construction sector as well as in industrial insulation is quite significant.^[29]
- In granular form to add [insulation](#) to [skylights](#). Georgia Institute of Technology's 2007 [Solar Decathlon](#) House project used an aerogel as an insulator in the semi-transparent roof.^[30]
- A chemical [adsorber](#) for cleaning up spills.^[31]
- A [catalyst](#) or a catalyst carrier.
- [Thickening agents](#) in some [paints](#) and [cosmetics](#).
- Laser targets for the [National Ignition Facility](#).
- Commercial manufacture of aerogel 'blankets' began around the year 2000, combining silica aerogel and fibrous reinforcement that turns the brittle aerogel into a durable, flexible material. The mechanical and thermal properties of the product may be varied based upon the choice of reinforcing fibers, the aerogel matrix and [opacification additives](#) included in the composite.
- [NASA](#) used an aerogel to trap [space dust](#) particles aboard the [Stardust](#) spacecraft. The particles vaporize on impact with solids and pass through gases, but can be trapped in aerogels. NASA also used aerogel for [thermal insulation](#) of the [Mars Rover](#) and [space suits](#).^{[32][33]}
- The [US Navy](#) is evaluating aerogel undergarments as passive thermal protection for divers.^[34]
- In [particle physics](#) as radiators in [Cherenkov effect](#) detectors, such as the ACC system of the Belle detector, used in the [Belle Experiment](#) at [KEKB](#). The suitability of aerogels is determined by their low [index of refraction](#), filling the gap between gases and liquids, and their transparency and solid state, making them easier to use than [cryogenic](#) liquids or compressed gases. Their low mass is also advantageous for space missions.
- [Resorcinol–formaldehyde](#) aerogels (polymers chemically similar to [phenol formaldehyde resins](#)) are used as precursors for manufacture of carbon aerogels, or when an organic insulator with large surface is desired. They come as high-density material, with surface area about 600 m²/g.
- Metal–aerogel [nanocomposites](#) prepared by impregnating the hydrogel with solution containing ions of a [transition metal](#) and irradiating the result with [gamma rays](#), precipitates nanoparticles of the metal. Such composites can be used as [catalysts](#), sensors, [electromagnetic shielding](#), and in waste disposal. A prospective use of platinum-on-carbon catalysts is in [fuel cells](#).
- As a drug delivery system owing to its [biocompatibility](#). Due to its high surface area and porous structure, drugs can be adsorbed from supercritical CO₂. The release rate of the drugs can be tailored by varying the properties of the aerogel.^[35]
- Carbon aerogels are used in the construction of small electrochemical double layer [supercapacitors](#). Due to the high surface area of the aerogel, these capacitors can be 1/2000th to 1/5000th the size of similarly rated electrolytic capacitors.^[36] Aerogel supercapacitors can have a very low [impedance](#) compared to normal supercapacitors and can absorb or produce very high



The [Stardust](#) dust collector with aerogel blocks. (NASA)

peak currents. At present, such capacitors are [polarity-sensitive](#) and need to be wired in series to achieve a working voltage of greater than about 2.75 V.

- [Dunlop Sport](#) uses aerogel in some of its racquets for tennis, squash and badminton.
- In water purification, [chalcogels](#) have shown promise in absorbing the heavy metal pollutants mercury, lead, and cadmium from water.^[37]
- Aerogel can introduce disorder into [superfluid helium-3](#).^[38]
- In aircraft de-icing, a new proposal uses a [carbon nanotube](#) aerogel. A thin filament is spun on a winder to create a 10 micron-thick film, equivalent to an [A4 sheet of paper](#). The amount of material needed to cover the wings of a jumbo jet weighs 80 grams (2.8 oz). Aerogel heaters could be left on continuously at low power, to prevent ice from forming.^[39]
- [Chevrolet Corvette \(C7\)](#) transmission tunnel^[40]
- [CamelBak](#) uses aerogel as insulation in a thermal sport bottle ^[41]
- 45 North uses aerogel as palm insulation in its Sturmfiist 5 cycling gloves. ^[42]

Production

Production of aerogels is done by the [sol-gel](#) process. First a [gel](#) is created in solution and then the liquid is carefully removed to leave the aerogel intact.

When this interlinking has stopped the flow of liquid within the material, this is known as a [gel](#). This process is known as gelation. These reactions generally have moderately slow reaction rates, and as a result either acidic or basic [catalysts](#) are used to improve the processing speed. Basic catalysts tend to produce more transparent aerogels with less shrinkage.^[43]

The removal of the liquid from a true aerogel involves special processing. Gels where the liquid is allowed to evaporate normally are known as [xerogels](#). As the liquid evaporates, forces caused by [surface tensions](#) of the liquid-solid [interfaces](#) are enough to destroy the fragile gel network. As a result, xerogels cannot achieve the high porosities and instead peak at lower porosities and exhibit large amounts of shrinkage after drying.^[45]

In 1931, to develop the first aerogels, Kistler used a process known as [supercritical drying](#). By increasing the temperature and pressure he forced the liquid into a [supercritical fluid](#) state where by dropping the pressure he could instantly gasify and remove the liquid inside the aerogel, avoiding damage to the delicate three-dimensional network. While this can be done with [ethanol](#), the high temperatures and pressures lead to dangerous processing conditions. A safer, lower temperature and pressure method involves a solvent exchange. This is typically done by exchanging the initial aqueous pore liquid for a [CO2](#) miscible liquid such as [ethanol](#) or [acetone](#), then onto liquid [carbon dioxide](#) and then bringing the carbon dioxide above its [critical point](#). A variant on this process involves the direct injection of supercritical carbon dioxide into the pressure vessel containing the aerogel. The end result of either process exchanges the initial liquid from the gel with [carbon dioxide](#), without allowing the gel structure to collapse or lose volume.^[43]

Aerogel [composites](#) have been made using a variety of continuous and discontinuous [reinforcements](#). The high aspect ratio of fibers such as [fiberglass](#) have been used to reinforce aerogel composites with significantly improved mechanical properties.

[Resorcinol–formaldehyde](#) aerogel (RF aerogel) is made in a way similar to production of silica aerogel.

Carbon aerogel is made from a resorcinol–formaldehyde aerogel by its [pyrolysis](#) in [inert gas](#) atmosphere,

leaving a matrix of [carbon](#). It is commercially available as solid shapes, powders, or composite paper.

Safety

Silica-based aerogels are not known to be [carcinogenic](#) or toxic. However, they are a mechanical [irritant](#) to the eyes, skin, respiratory tract, and digestive system. Small silica particles can potentially cause [silicosis](#) when inhaled. They also can induce dryness of the skin, eyes, and mucous membranes. Therefore, it is recommended that protective gear including respiratory protection, gloves and eye goggles be worn whenever handling aerogels.^[46]

See also

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Further reading

External links

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