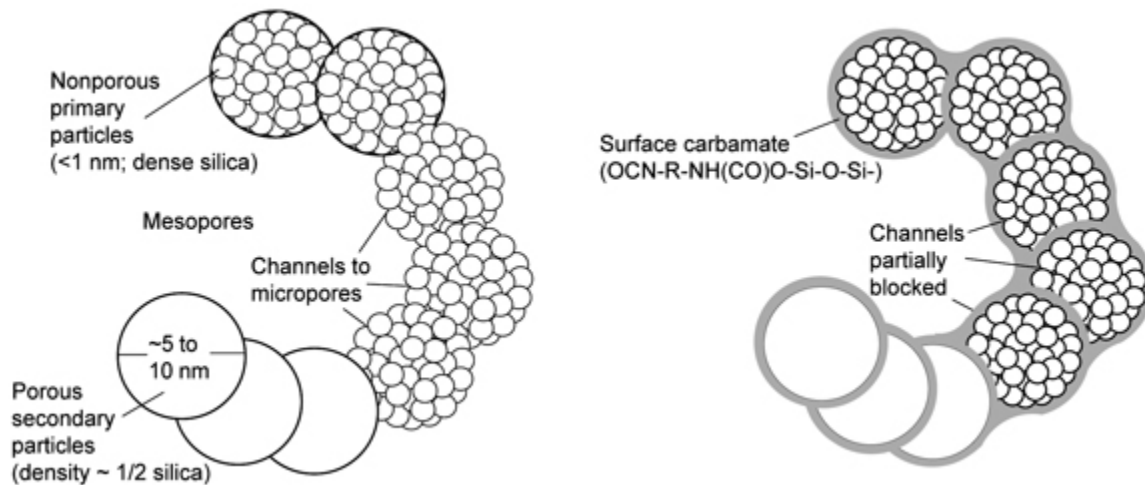


Mechanically Strong, Lightweight Porous Materials Developed (X-Aerogels)

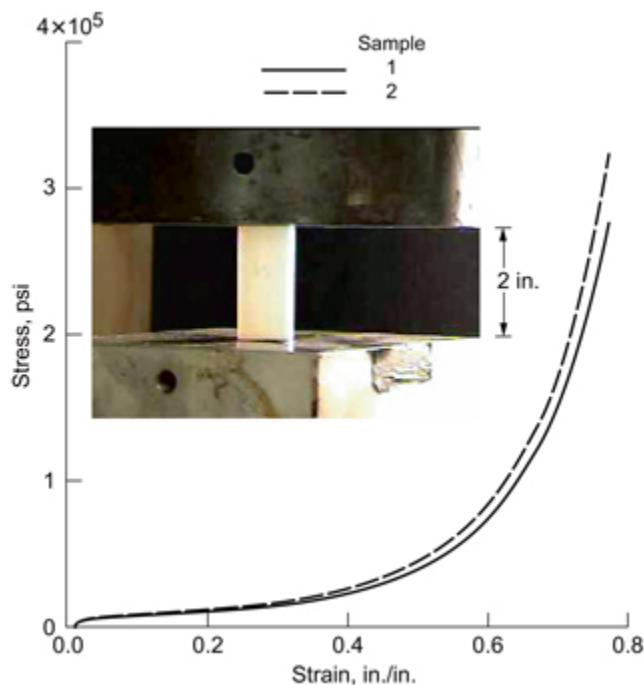
Aerogels are attractive materials for a variety of NASA missions because they are ultralightweight, have low thermal conductivity and low-dielectric constants, and can be readily doped with other materials. Potential NASA applications for these materials include lightweight insulation for spacecraft, habitats, and extravehicular activity (EVA) suits; catalyst supports for fuel cell and in situ resource utilization; and sensors for air- and water-quality monitoring for vehicles, habitats, and EVA suits. Conventional aerogels are extremely fragile and require processing via supercritical fluid extraction, which adds cost to the production of an aerogel and limits the sizes and geometries of samples that can be produced from these materials. These issues have severely hampered the application of aerogels in NASA missions.



Microstructure of a silica aerogel cross-linked with polyurethane/polyurea (X-Aerogel). Long description of figure 1. Illustrations show nonporous primary particles (less than 1 nanometer; dense silica), mesopores, channels to micropores, porous secondary particles (about 5 to 10 nanometers; density about one-half silica), surface carbamate (OCN-R-NH(CO)O-Si-O-Si-), and partially blocked channels.

Researchers at the NASA Glenn Research Center have developed a new class of strong lightweight materials, named X-Aerogels. These materials are produced by reacting the internal (mesoporous) surfaces of porous networks of inorganic nanoparticles with polymeric crosslinking agents. The preceding illustration shows a schematic of the microstructure of a typical aerogel before and after modification with an isocyanate. Reaction of the aerogel with an isocyanate monomer results in the formation of conformal polyurethane/polyurea coating around the nanoparticles.

The most striking feature of these novel aerogels is that, for a nominal increase (no more than threefold) in density, their mechanical strength can be increased by up to 300 times that of the unmodified, conventional aerogel. In addition, the specific compressive strength of these materials is ~ 10 times that of steel. The following graph shows the stress-strain curve of a typical X-Aerogel monolith under a compressive load. Under compression, these aerogels fail at a load of about 30,000 psi, where the sample has been compressed to ~ 80 percent of its original length with no buckling and only a minimal “swelling” (~ 20 percent). This occurs mostly during the last stages of compression. The thermal conductivity of these new aerogels is in the range of only 20 to 40 mW/mK, about 3 to 6 times that of a conventional aerogel. Ongoing research with these materials is focused on reducing their density and thermal conductivity while maintaining acceptable mechanical properties.



Stress-strain curves from a compressive strength test of an isocyanate cross-linked X-Aerogel monolith ($\sim 0.45 \text{ g/cm}^3$, 0.7 in. diam, 1.5 in. long). (Data obtained by Prof. Samit Roy of Oklahoma State University).

Other Glenn research is focused on extending this approach to the development of other aerogel materials. Two parallel approaches are being explored--(1) developing aerogels from other inorganic materials besides silica and (2) utilizing other polymer cross-linkers besides diisocyanates. Along these lines, this chemistry has been successfully demonstrated with more than 35 different metal and semimetal oxide aerogels utilizing isocyanates as the cross-linking agent (see the periodic table). Many of these new aerogel materials may have unique optical, magnetic, or electrical properties.

and industrial applications, thermal and acoustic insulation materials for construction, optoelectronic devices, and packaging of sensitive electronic devices.

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