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CO₂ adsorption in a hierarchically structured carbon by SANS
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TESCU, NIDIA GALLEGU, Oak Ridge National Laboratory — This contribution
investigated the high pressure adsorption behavior of CO₂ at T = 296 K in hierarchi-
cally structured carbon using small-angle neutron scattering (SANS) technique. We
observed a strong densification of CO₂ in micropores accompanied by non-monotonic
adsorption-induced pore deformation. Liquid-like density of CO₂ confined in the mi-
cropores was reached with increasing pressure to 20 bar, which corresponds to the
relative pressure of $P/P_{sat} \sim 0.3$. At $P > 20$ bar, density of confined CO₂ approached
a plateau. The size of micropores first increases with pressure, reached a maximum
at 20 bar, and then decreased with further increasing pressure. A complementary
SANS experiment carried out on the same microporous carbon saturated with argon
that is neutron-transparent and non-adsorbing inert shows no deformation of
micropores at pressures up to ~ 200 bars. This result proved that the observed de-
formation of micropores in CO₂ was an adsorption-induced phenomenon, caused by
the solvation pressure - induced strain and strong densification of confined CO₂ in
the micropores.

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