

SPONTANEOUS WATER DESALINATION IN GRAPHENE OXIDE FRAMEWORK

Luca CERVINI, John GRIFFIN, Nathan BARROW

(n.

Background: ACTIVATED CARBONS

Synthesis: carbonization of PEEK and subsequent steam-activation

Structure: amorphous but presence of nm-sized ordered graphitic domains -



Powder X-Ray Diffraction

Transmission Electron Microscopy



small pore volume created by carbonization pore volume and average pore size increase proportionally to activation time



2-Theta angle







Nuclear Magnetic Resonance

ex-pore

 δ ²³Na (ppm)

2 3 J Pore size (nm)

saturate sample with NaCl(aq) Na⁺ ions reach equilibrium distribution across the pore network position of Na⁺ in-pore peak depends on average perceived pore size

liquid N₂ populates the pores at specific pressures depending on their sizes introduce known volumes of N₂ into pores of increasing size 3 distinct pore size populations (0.8 nm, 1-2 nm, 3-4 nm)

Na⁺ enters subnanometre pores only at high concentrations

Application: SPONTANEOUS WATER DESALINATION

- Pore requirement: hydrophobic/ionophobic, subnanometre size, open-ended, stable in saline water - Target material: graphene-based framework, apolar pillars, e.g. alkyl-ether





CONTACT PERSON

Luca Cervini, I.cervini@lancaster.ac.uk

REFERENCES

L. Cervini, O. D. Lynes, G. R. Akien, A. Kerridge, N. S. Barrow, J. M. Griffin; Energy Storage Materials 2019, 21, 335 S. Eigler; Chem. Commun. 2015, 51, 3162

S. Eigler, Y. Hu, Y. Ishii, A. Hirsch; Nanoscale 2013, 5, 12136

raphene Conline 2020