

A water-based and high space-time yield synthetic route to MOF Ni₂(dhtp) and its linker 2,5-dihydroxyterephthalic acid.

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DOI: 10.1039/b000000x/

Supplementary information

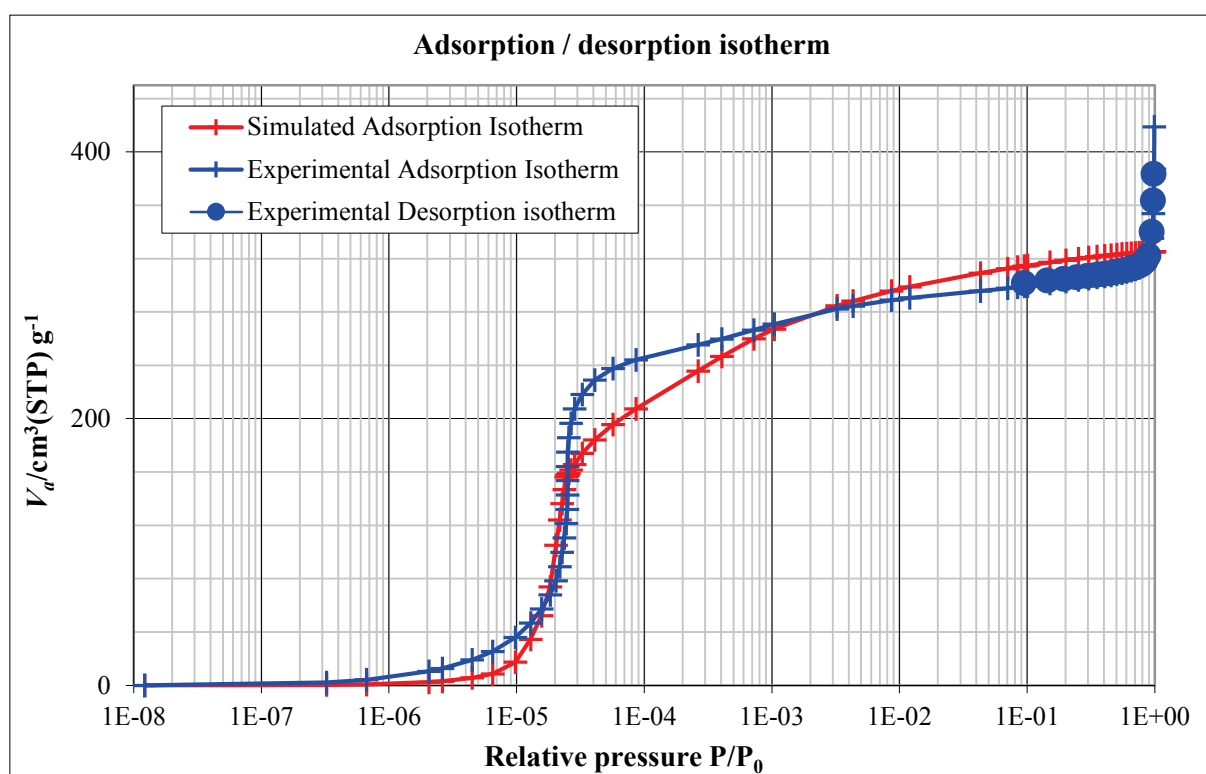


Figure S1: Experimental nitrogen adsorption (blue crosses) and desorption (blue dots) isotherms at 77K for MOF Ni₂(dhtp) synthesized in water, and simulated adsorption isotherm (red curve) obtained using the NLDFIT simulation method^{S1-S11}.

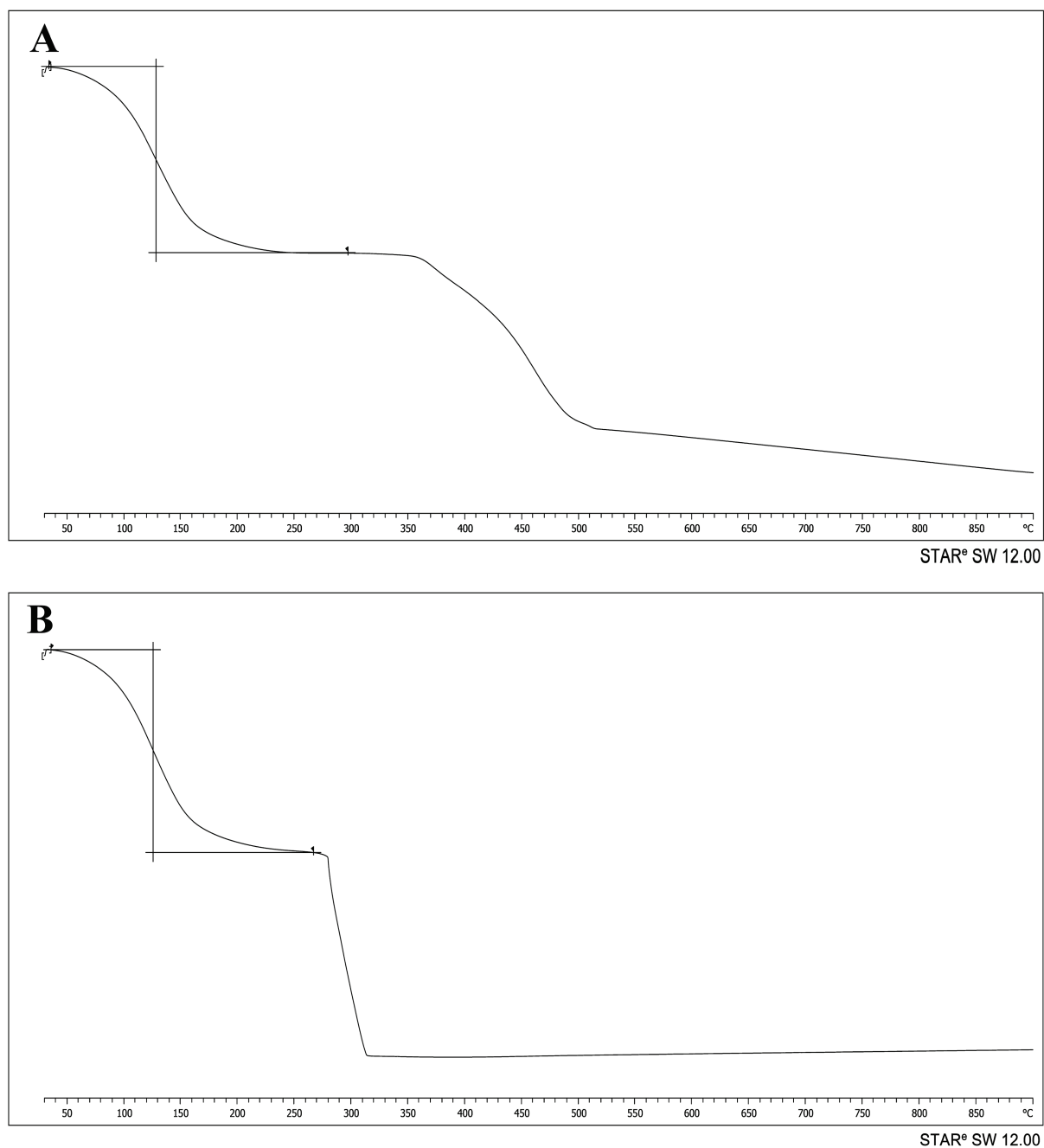


Figure S2: Thermogravimetric analysis of MOF $\text{Ni}_2(\text{dhtp})$ synthesized in water measured under N_2 (A) and under air (B).

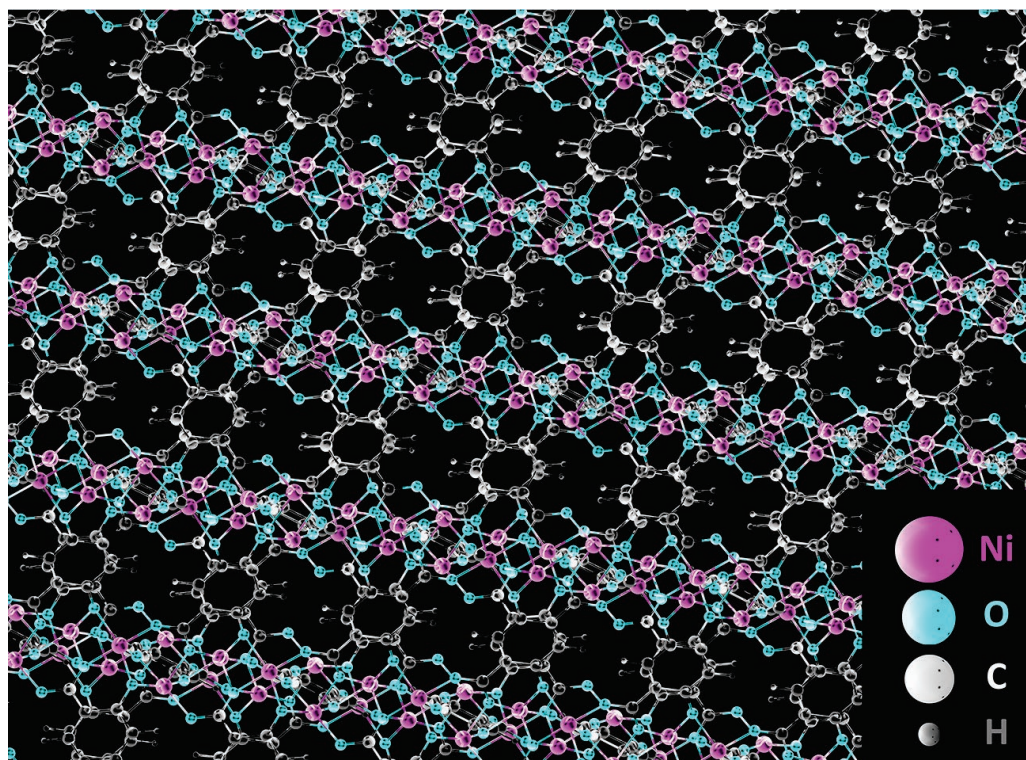


Figure S3: Chemical Structure of $\text{Ni}_2(\text{dhtp})$ drawn using the available crystallographic data^{S12} and oriented toward the (-120) plane, matching the observed HRTEM micrograph reported in **Figure 4A)**

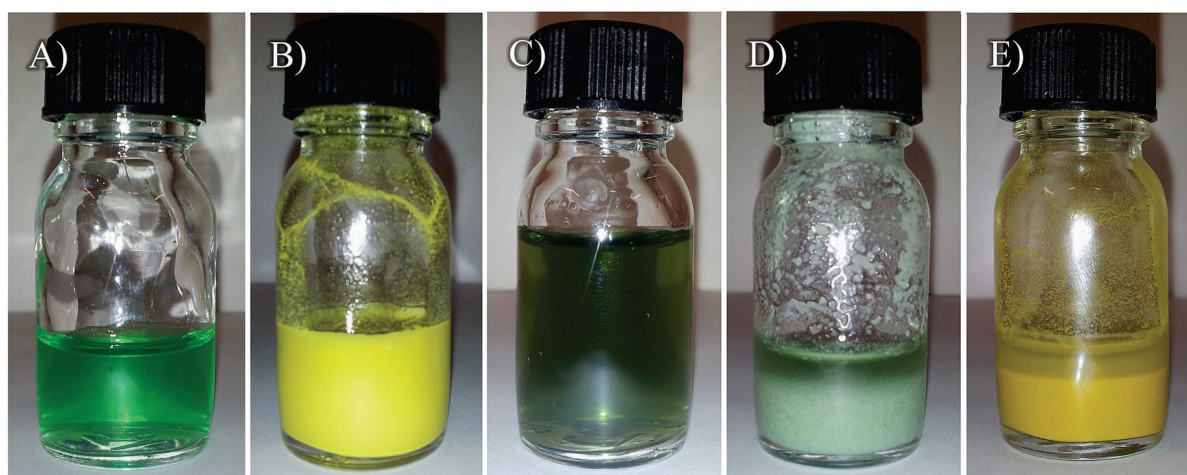


Figure S4: Photographs of samples taken at different steps of the $\text{Ni}_2(\text{dhtp})$ synthesis in water (**A**: 1M aqueous solution of nickel (II) acetate; **B**: suspension of 2,5-dihydroxyterephthalic acid in water; **C**: homogeneous solution obtained by mixing A and B; **D**: precipitate obtained from solution C at room temperature; **E**: $\text{Ni}_2(\text{dhtp})$ obtained from solution C after 1h under reflux.

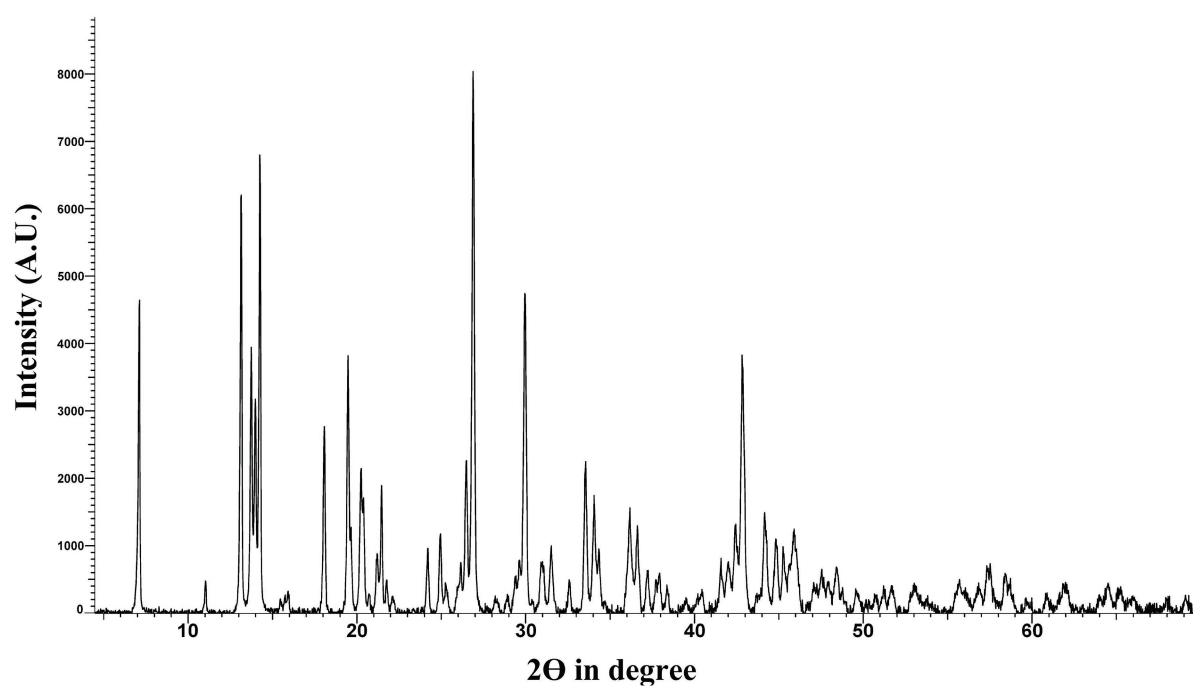


Figure S5: X-ray diffraction pattern of the intermediate which precipitate at room temperature from the solution obtained by mixing nickel acetate and 2,5-dihydroxyterephthalic acid in water (corresponding to **Figure S4 D**).

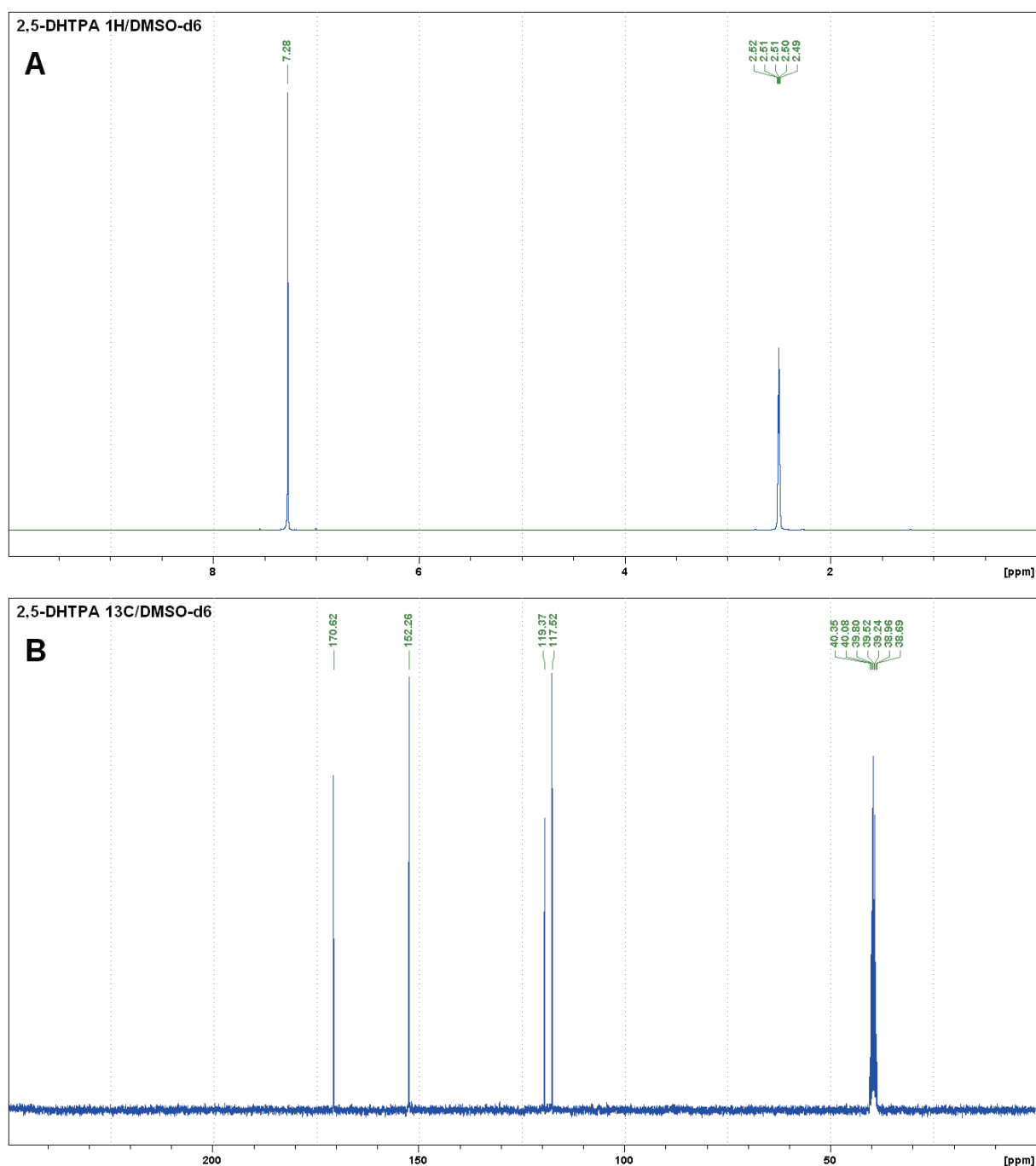


Figure S6: **A:** ¹H NMR and **B:** ¹³C NMR spectra (in DMSO-d₆) of the 2,5-dihydroxyterephthalic acid synthesized by hydroquinone dicarboxylation (see **Scheme 2**).

Notes and references

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