

Supporting Information

A family of microporous carbons prepared via a simple metal salt carbonization route with high selectivity for exceptional gravimetric and volumetric post-combustion CO₂ uptake

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Supporting Table S1. Textural properties and CO₂ uptake of metal salt (potassium hydrogen phthalate, KHP) derived carbon materials via carbonisation at 900 °C for 1 to 4 h.

Sample	Surface area ^a (m ² /g)	Pore volume ^b (cm ³ /g)	Pore size ^c (Å)	CO ₂ uptake ^d (mmol/g)		
				0.15 bar	1 bar	20 bar
CKHP900-1	1601 (1417)	0.77 (0.63)	5/6/8/12/20	1.0	3.7	13.0
CKHP900-2	2104 (1805))	1.11 (0.81)	8.5/12/16/22	0.8	3.0	12.7
CKHP900-3	1600 (1360)	0.81 (0.61)	6.5/8.5/12/21	1.1	3.8	13.7
CKHP900-4	1561 (1452)	0.81 (0.69)	6.8/8.5/12	0.9	3.5	12.6

The values in the parenthesis refer to: ^amicropore surface area and ^bmicropore volume. ^cpore size distribution maxima obtained from NLDFT analysis. ^dCO₂ uptake at 25 °C and various pressures (i.e., 0.15 bar, 1 bar and 20 bar).

Supporting Table S2. Textural properties and CO₂ uptake of metal salt (potassium hydrogen phthalate, KHP) derived carbon materials via carbonisation at 700 °C for 1 to 4 h.

Sample	Surface area ^a (m ² /g)	Pore volume ^b (cm ³ /g)	Pore size ^c (Å)	CO ₂ uptake ^d (mmol/g)		
				0.15 bar	1 bar	20 bar
CKHP700-1	932 (900)	0.44 (0.39)	7/8.5	1.6	4.2	7.9
CKHP700-2	795 (760))	0.44 (0.34))	6	1.7	4.1	6.9
CKHP700-3	1019 (991)	0.47 (0.43)	7/8.5	1.5	4.2	8.8
CKHP700-4	1014 (983)	0.47 (0.43)	6/8.5	1.5	4.4	8.9

The values in the parenthesis refer to: ^amicropore surface area and ^bmicropore volume. ^cpore size distribution maxima obtained from NLDFT analysis. ^dCO₂ uptake at 25 °C and various pressures (i.e., 0.15 bar, 1 bar and 20 bar).

Supporting Table S3. CO₂ uptake of various porous carbons at 25 °C and 0.15 bar or 1 bar.

	CO ₂ uptake (mmol/g)		Reference
	1 bar	0.15 bar	
Sawdust-derived activated carbon	4.8	1.2	1
KOH-activated templated carbons	3.4	~1.0	2
Hierarchical porous carbon (HPC)	3.0	~0.9	3
Petroleum pitch-derived activated carbon	4.55	~1.0	4
Activated carbon spheres	4.55	~1.1	5
Phenolic resin activated carbon spheres	4.5	~1.2	6
Poly(benzoxazine-co-resol)-derived carbon	3.3	1.0	7
Fungi-derived activated carbon	3.5	~1.0	8
Chitosan-derived activated carbon	3.86	~1.1	9
Polypyrrole derived activated carbon	3.9	~1.0	10
Soya bean derived N-doped activated carbon	4.24	1.2	11
N-doped ZTCs	4.4	~1.0	12
Activated templated N-doped carbon	4.5	1.4	13
Polyaniline derived activated carbon	4.3	1.38	14
N-doped activated carbon monoliths	5.14	1.25	15
Activated N-doped carbon	3.2	1.5	16
Activated hierarchical N-doped carbon	4.8	1.4	17
Activated N-doped carbon from algae	4.5	~1.1	18

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Supporting Table S4. CO₂ uptake at 0 °C of metal salt (potassium hydrogen phthalate, KHP) derived carbon materials via carbonisation at 800 °C for 1 to 4 h.

Sample	CO ₂ uptake at 0 °C (mmol/g)			
	1 bar	5 bar	10 bar	20 bar
CKHP800-1	6.4	10.4	11.6	12.1
CKHP800-2	6.5	11.0	12.5	13.3
CKHP800-3	5.5	10.7	12.9	13.9
CKHP800-4	6.5	10.6	11.7	12.3

Supporting Table S5. CO₂ uptake at 0 °C of metal salt (potassium hydrogen phthalate, KHP) derived carbon materials via carbonisation at 900 °C for 1 to 4 h.

Sample	CO ₂ uptake at 0 °C (mmol/g)			
	1 bar	5 bar	10 bar	20 bar
CKHP900-1	5.3	10.8	13.3	15.3
CKHP900-2	4.8	10.3	13.2	15.7
CKHP900-3	5.1	11.2	13.2	15.5
CKHP900-4	5.3	10.5	13.0	14.8

Supporting Table S6. CO₂ uptake at 0 °C of metal salt (potassium hydrogen phthalate, KHP) derived carbon materials via carbonisation at 700 °C for 1 to 4 h.

Sample	CO ₂ uptake at 0 °C (mmol/g)			
	1 bar	5 bar	10 bar	20 bar
CKHP700-1	5.7	7.9	8.5	8.8
CKHP700-2	5.1	6.6	7.1	7.4
CKHP700-3	6.2	8.8	9.5	9.8
CKHP700-4	6.3	8.8	9.5	9.9

Supporting Table S7. Textural properties of unwashed CKHP carbon products

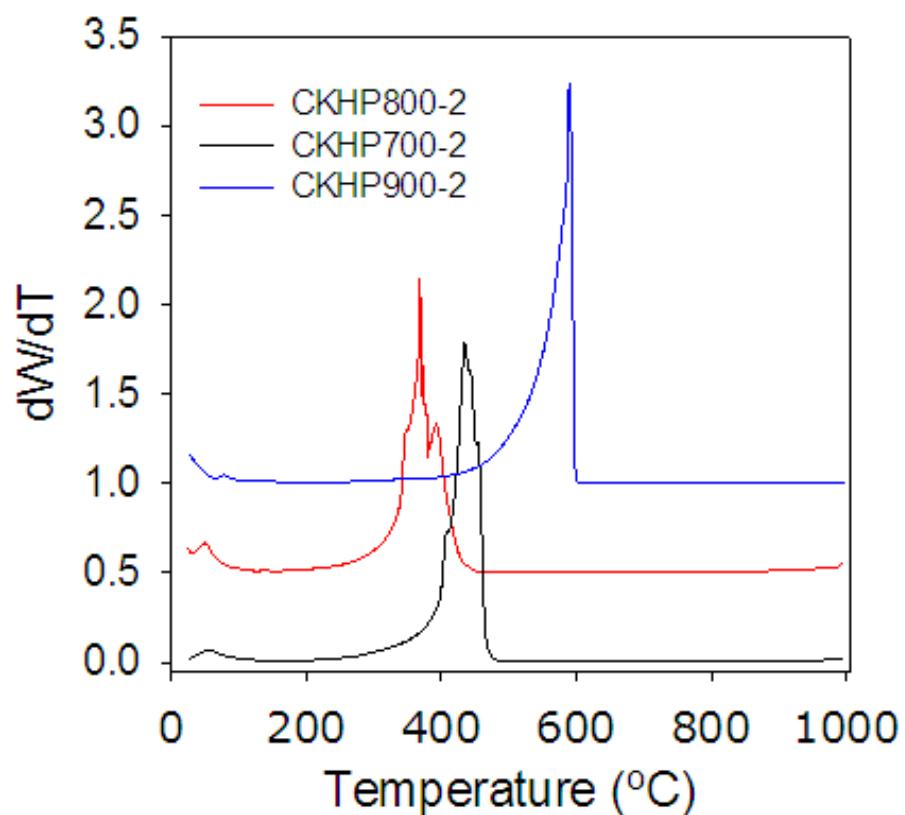
Sample	Surface area ^a (m ² /g)	Pore volume ^b (cm ³ /g)	Pore size ^c (Å)
CKHP700-2-UW	2 (1)		
CKHP800-2-UW	69 (68))	0.20 (0.03)	
CKHP900-2-UW	597 (467)	0.39 (0.22)	6.4/8.5/11.5
CKHP1000-2-UW	461 (395)	0.26(0.19)	6.5/8.5/11.5

The values in the parenthesis refer to: ^amicropore surface area and ^bmicropore volume. ^cpore size distribution maxima obtained from NLDFT analysis.

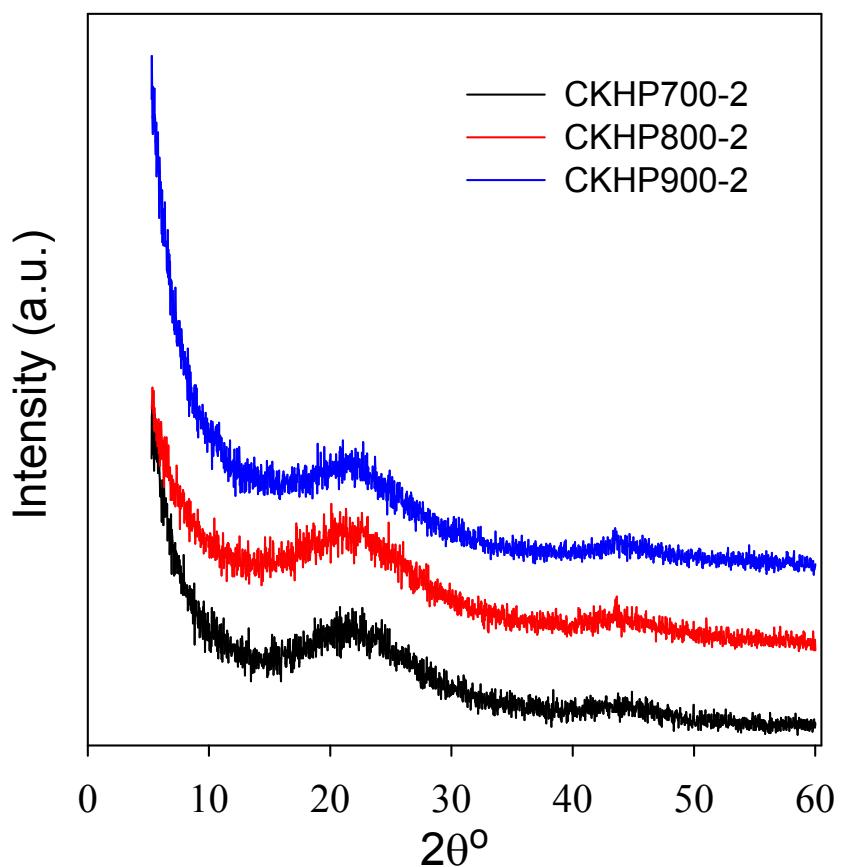
Supporting Table S8. Textural properties and CO₂ uptake of various CKHP carbons before and after compaction at a load of 5 tons (i.e. 370 MPa)

Sample	Surface area ^a (m ² /g)	Pore volume ^b (cm ³ /g)	Pore size ^c (Å)	CO ₂ uptake ^d (mmol/g)		
				0.15 bar	1 bar	20 bar
CKHP600-2	513 (494)	0.27 (0.24)	6.8	1.2	2.8	4.6
CKHP600-2-C5	516 (496)	0.28 (0.24)	5.8	1.4	3.3	4.9
CKHP700-1	932 (900)	0.44 (0.39)	7/8.5	1.6	4.2	7.9
CKHP700-1-C5	803 (775)	0.43 (0.37)	6/8	1.5	4.1	7.7
CKHP800-1	1255 (1200)	0.59 (0.52)	7/8.5/12	1.3	4.4	11.1
CKHP800-1-C5	1072 (1028)	0.56 (0.49)	6.5/8.5/12	1.3	4.2	10.8
CKHP900-1	1601 (1417)	0.77 (0.63)	5/6/8/12/20	1.0	3.7	13.0
CKHP900-1-C5	1508 (1403)	0.77 (0.66)	6/8/12	1.0	3.5	12.7
CKHP900-2-UW	597 (467)	0.39 (0.22)	6.4/8.5/11.5	1.2	2.9	8.6
CKHP900-2-UW-C5	789 (664)	0.48 (0.31)	6.4/8.5/11.5	1.4	3.0	7.4

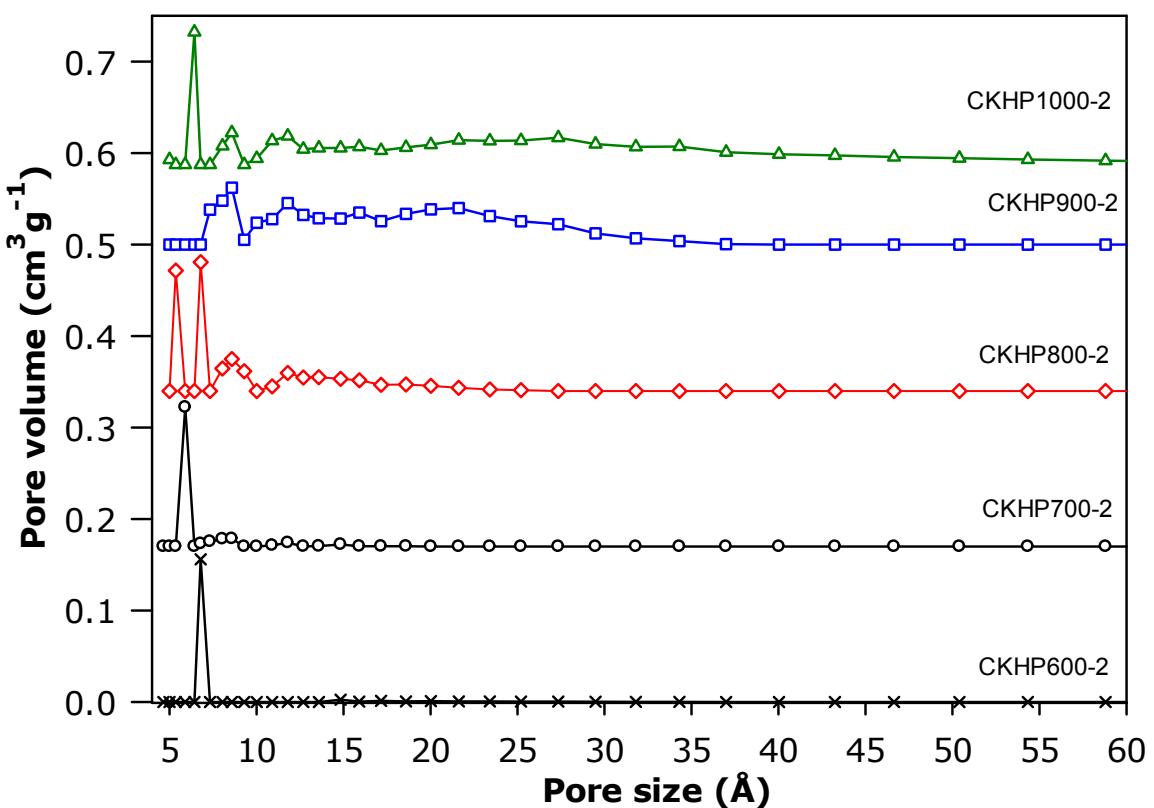
The values in the parenthesis refer to: ^amicropore surface area and ^bmicropore volume. ^cpore size distribution maxima obtained from NLDFT analysis. ^dCO₂ uptake at 25 °C and various pressures (i.e., 0.15 bar, 1 bar and 20 bar).



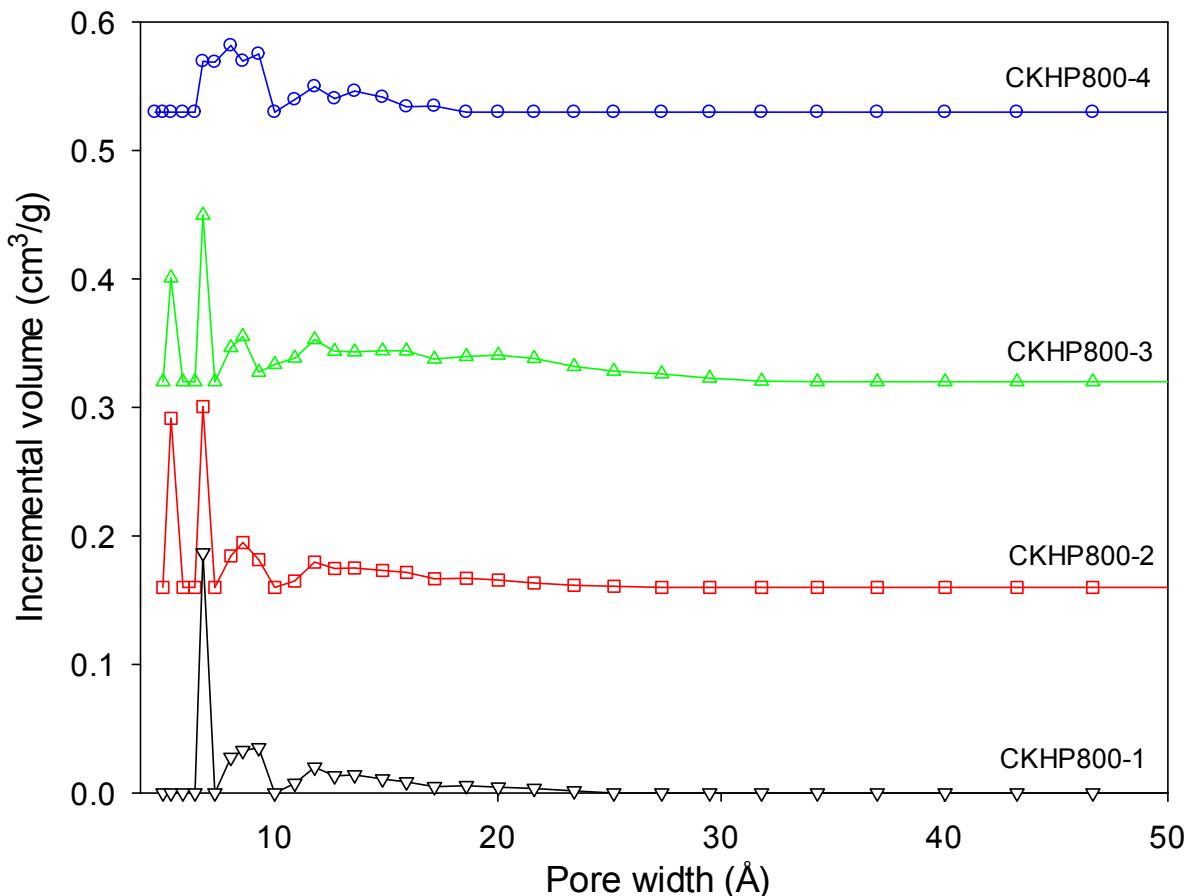
Supporting Figure S1. Differential thermogravimetric (DTG) profiles of CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at various temperatures for 2 h.



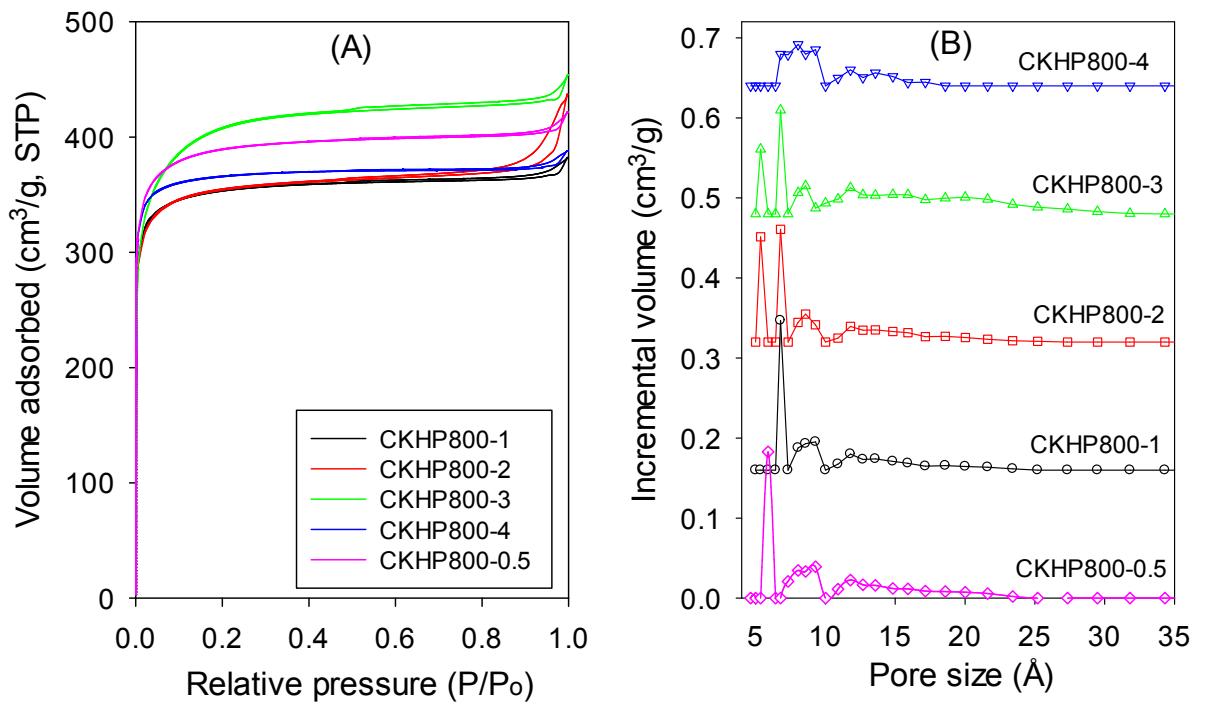
Supporting Figure S2. Powder XRD patterns of CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at various temperatures for 2 h.



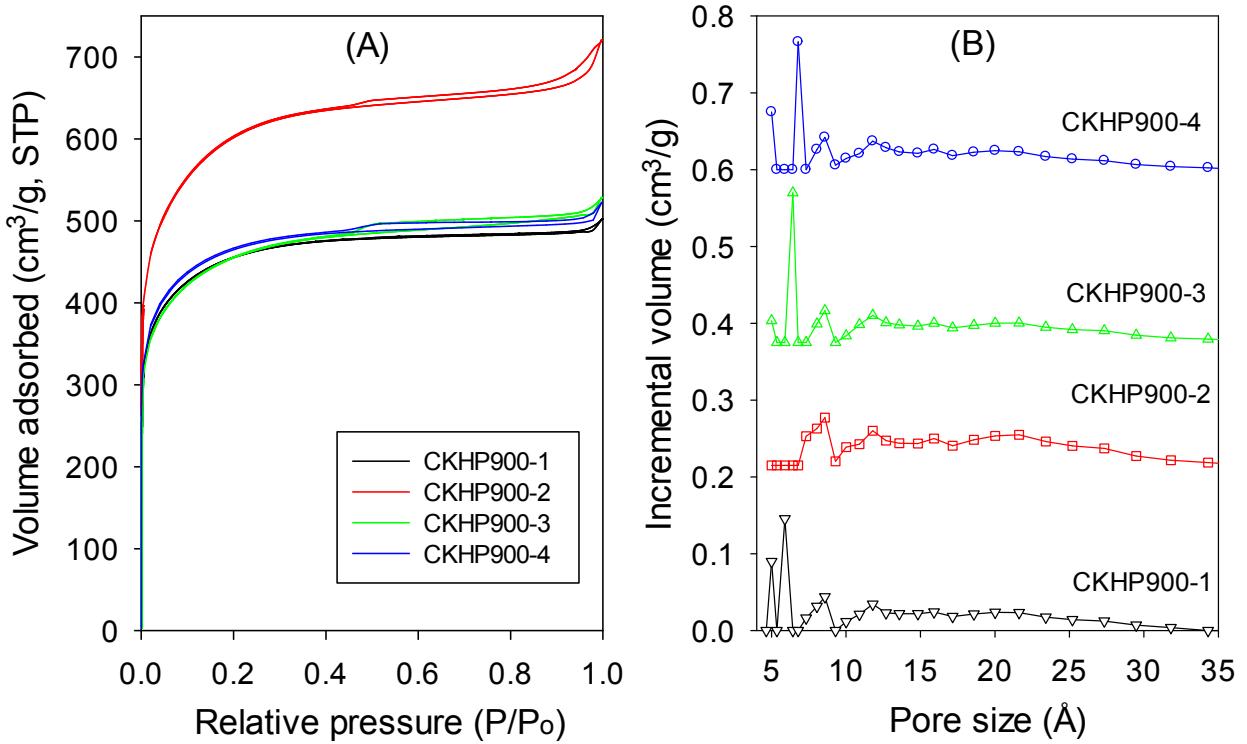
Supporting Figure S3. Pore size distribution curves of CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at various temperatures for 2 h. Samples CKHP600-2, CKHP700-2 and CKHP800-2 have no pores larger than 15 Å. Largest pores for CKHP900-2 and CKHP1000-2 are up to 30 Å and 40 Å, respectively, although even for these samples, only a small proportion of pores are larger than 15 Å.



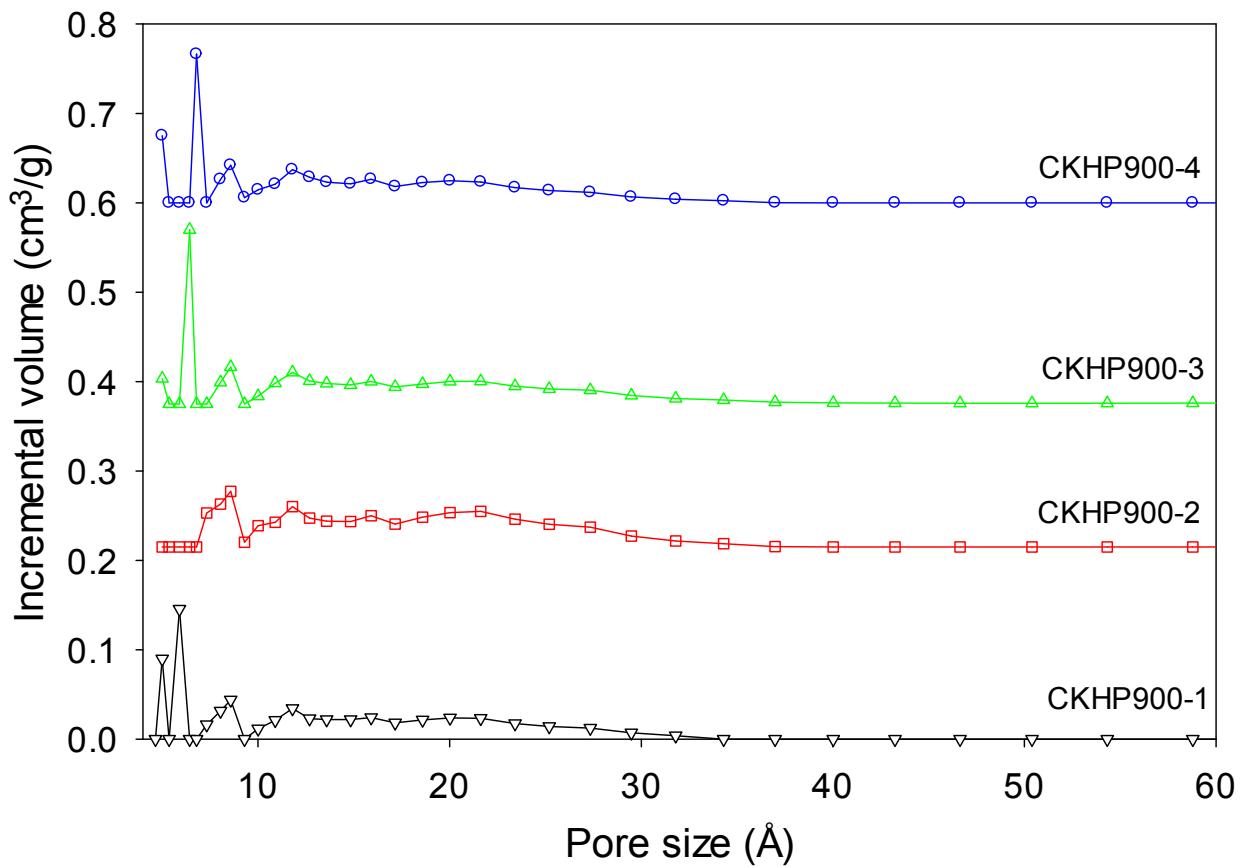
Supporting Figure S4. Pore size distribution curves of CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at 800 °C for between 1 and 4 h. Samples CKHP800-1, CKHP800-2 and CKHP800-4 have no pores larger than ca. 15 Å. For sample CKHP800-3 the largest pores are up to 30 Å, though only a small proportion of pores are larger than 15 Å.



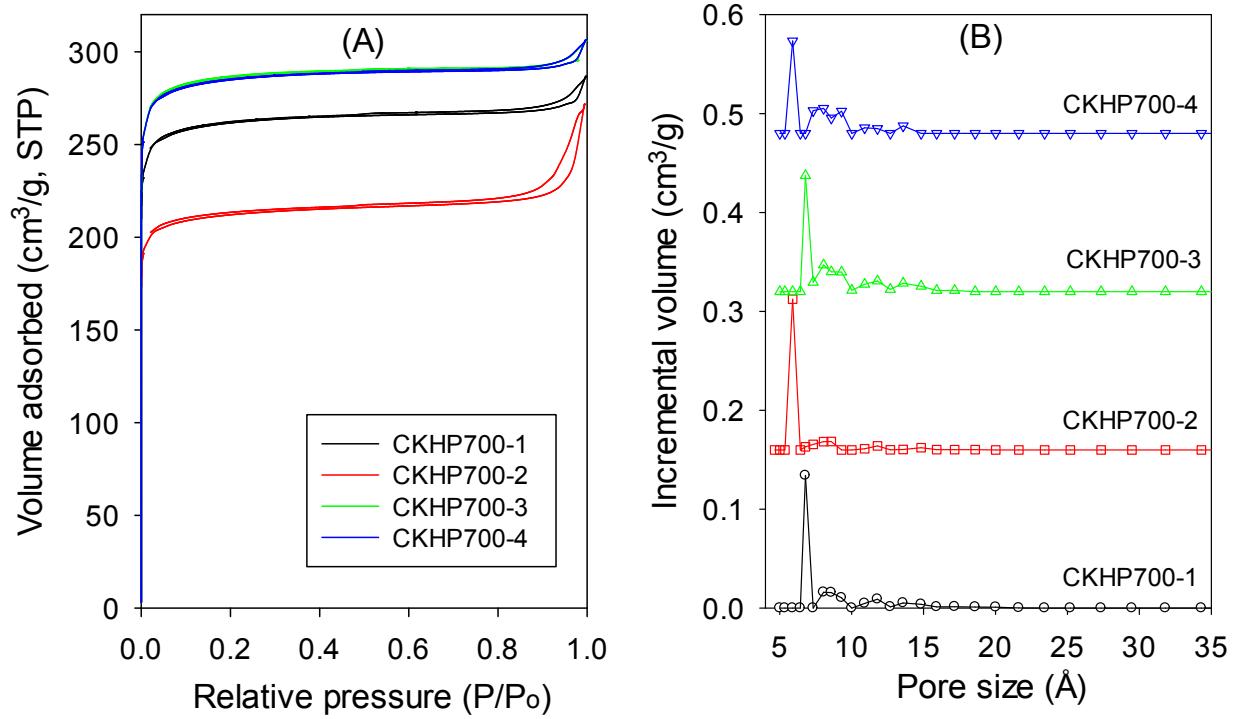
Supporting Figure S5. Nitrogen sorption isotherms (A) and corresponding pore size distribution curves (B) of CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at 800 °C for various periods of time, from as low as 0.5 h to 4 h. The isotherm and pore size distribution of the sample carbonised for 0.5 h is largely similar to the other CKHP800-y samples. The pore size distribution of CKHP800-0.5 exhibits maxima at 6 and 8.5 Å, which is very similar to sample CKHP800-1.



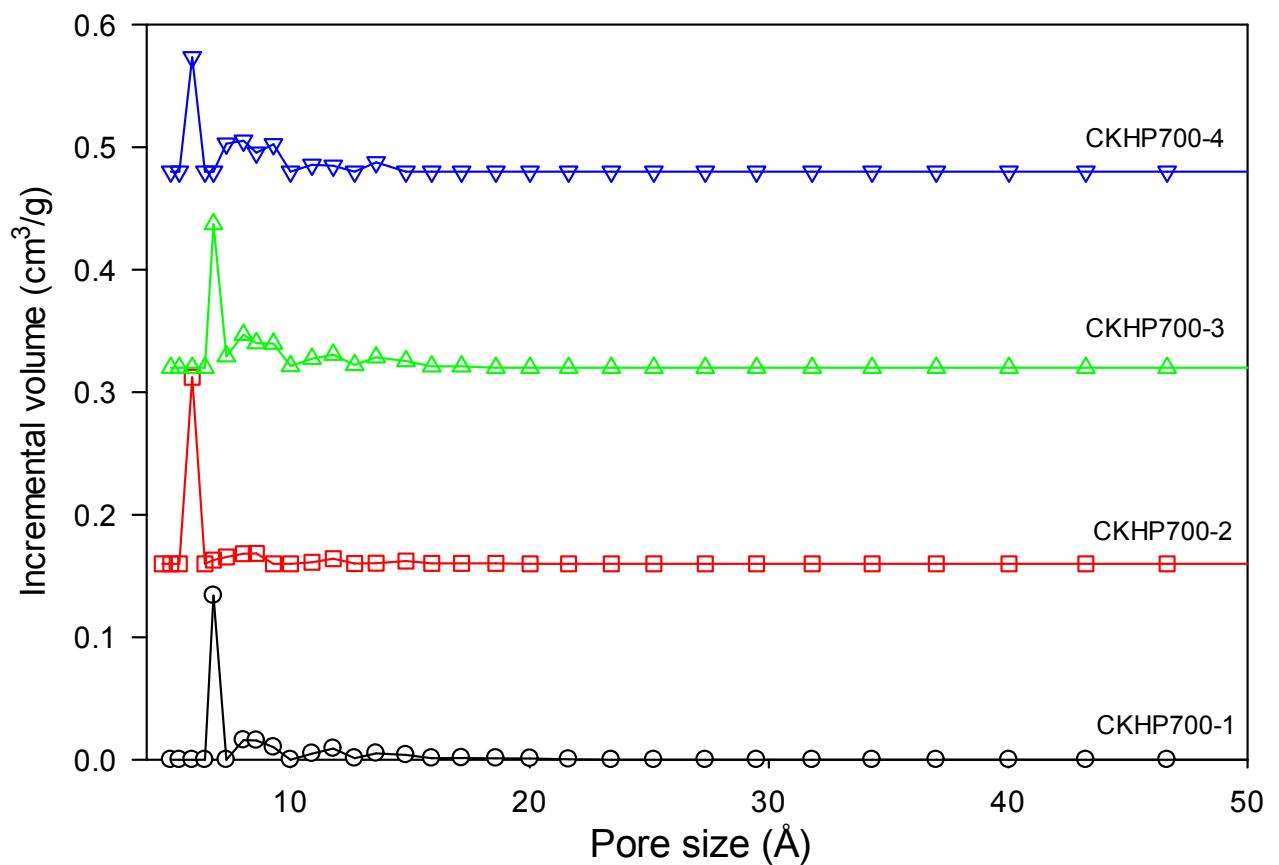
Supporting Figure S6. Nitrogen sorption isotherms (A) and corresponding pore size distribution curves (B) of CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at 900 °C for various periods of time.



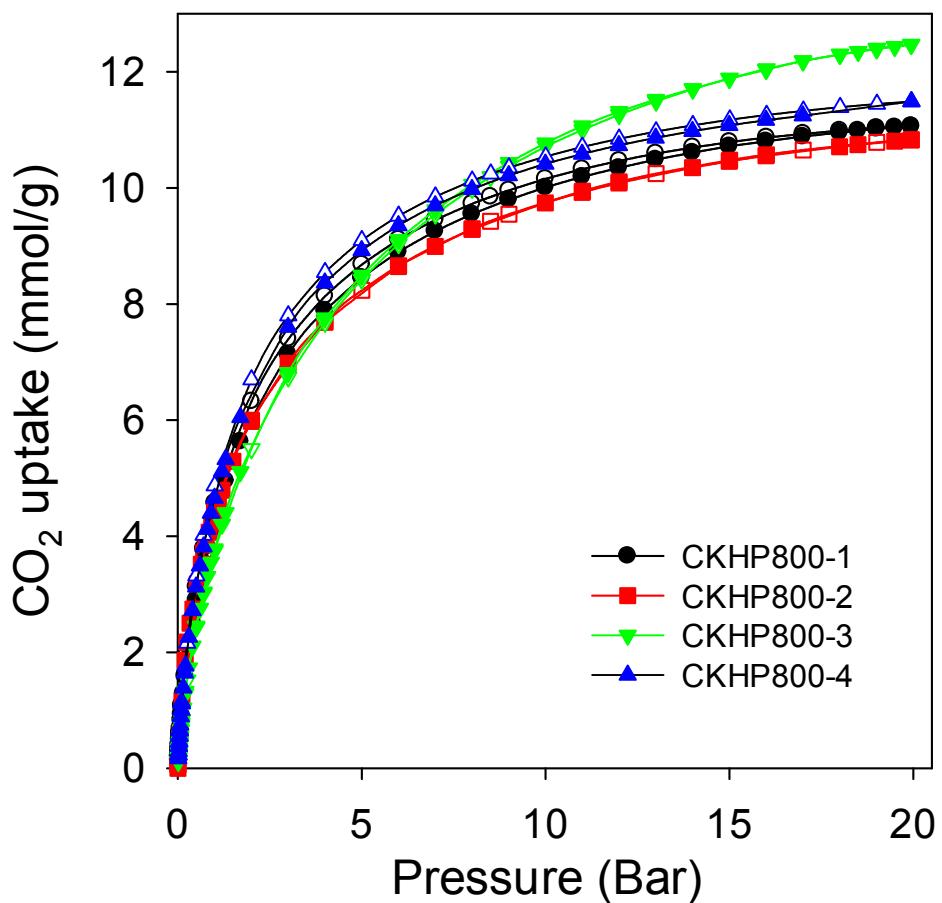
Supporting Figure S7. Pore size distribution curves of CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at 900 °C for between 1 and 4 h. The largest pores in all samples are up to 35 Å, and a significant proportion of pores are larger than 15 Å.



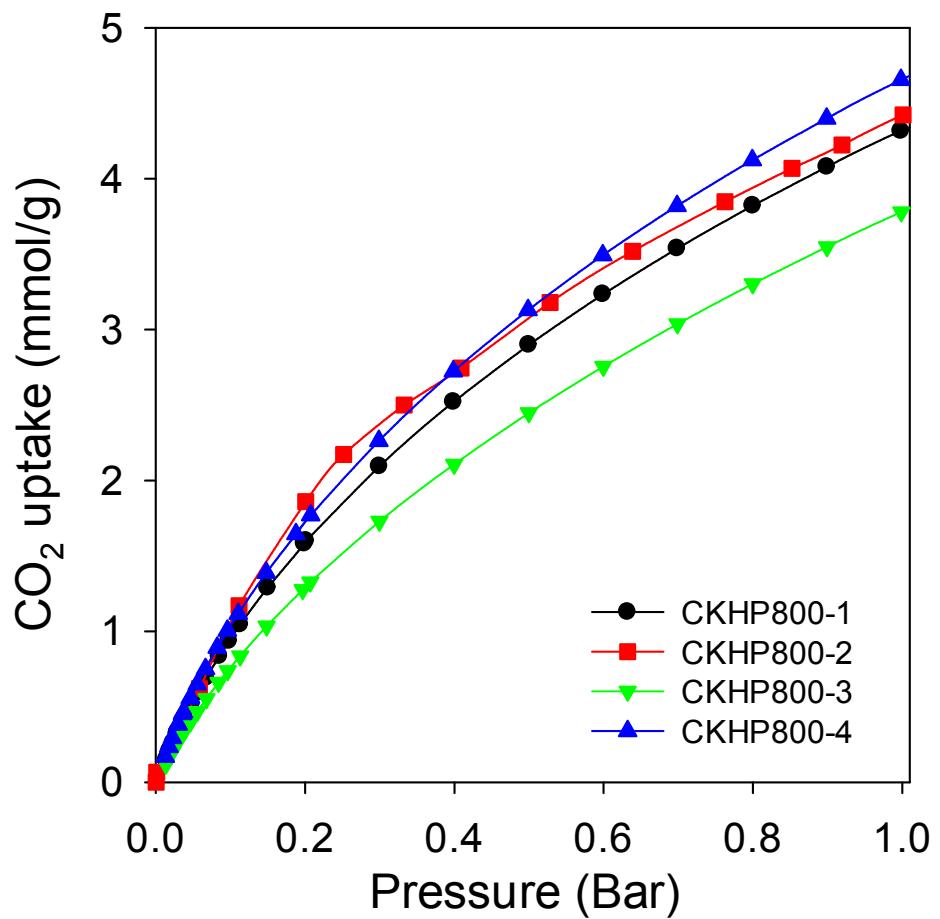
Supporting Figure S8. Nitrogen sorption isotherms (A) and corresponding pore size distribution curves (B) of CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at 700 °C for various periods of time.



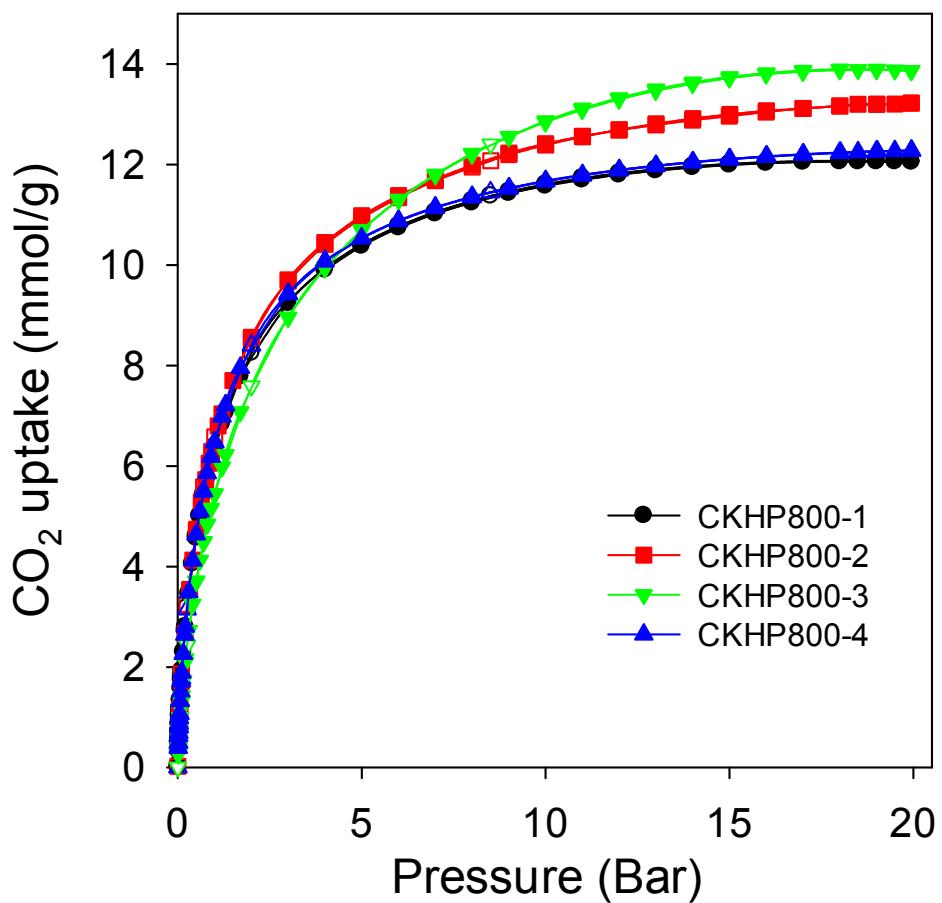
Supporting Figure S9. Pore size distribution curves of CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at 700 °C for between 1 and 4 h. The CKHP700-x samples have virtually no pores larger than 10 \AA .



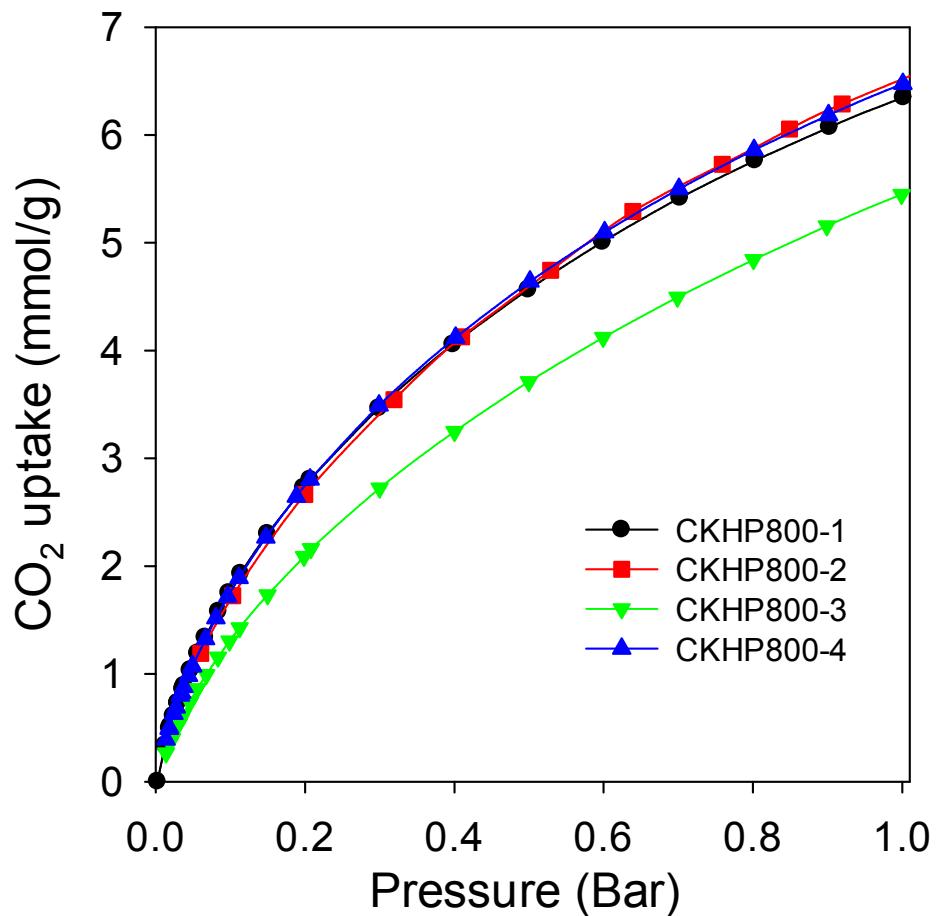
Supporting Figure S10. CO₂ uptake isotherms at 25 °C and 0 – 20 bar for CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at 800 °C for 1 to 4 h.



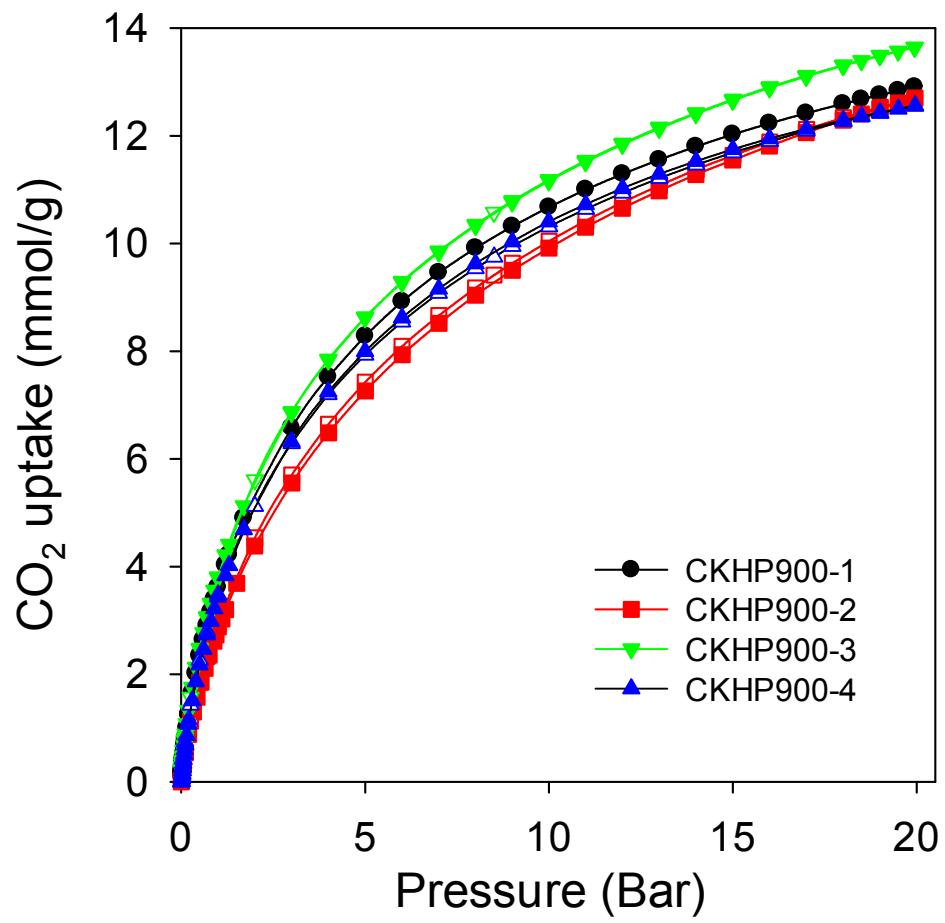
Supporting Figure S11. Low pressure CO₂ uptake isotherms at 25 °C for CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at 800 °C for 1 to 4 h.



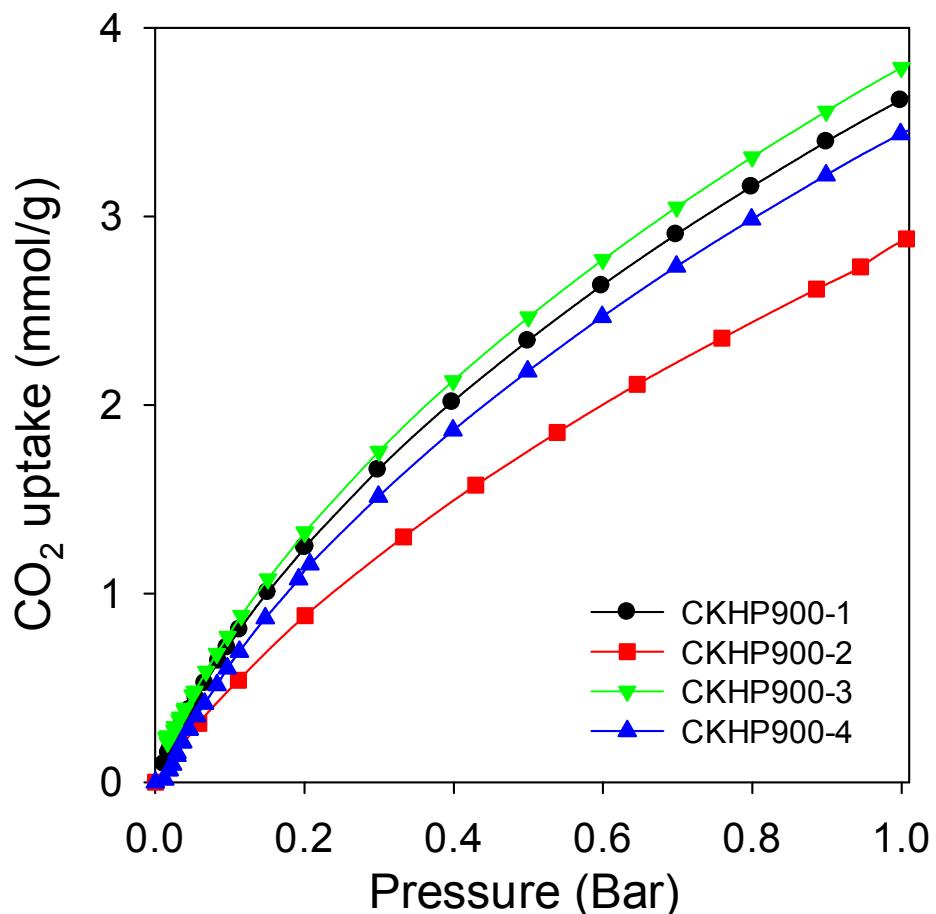
Supporting Figure S12. CO₂ uptake isotherms at 0 °C and 0 – 20 bar for CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at 800 °C for 1 to 4 h.



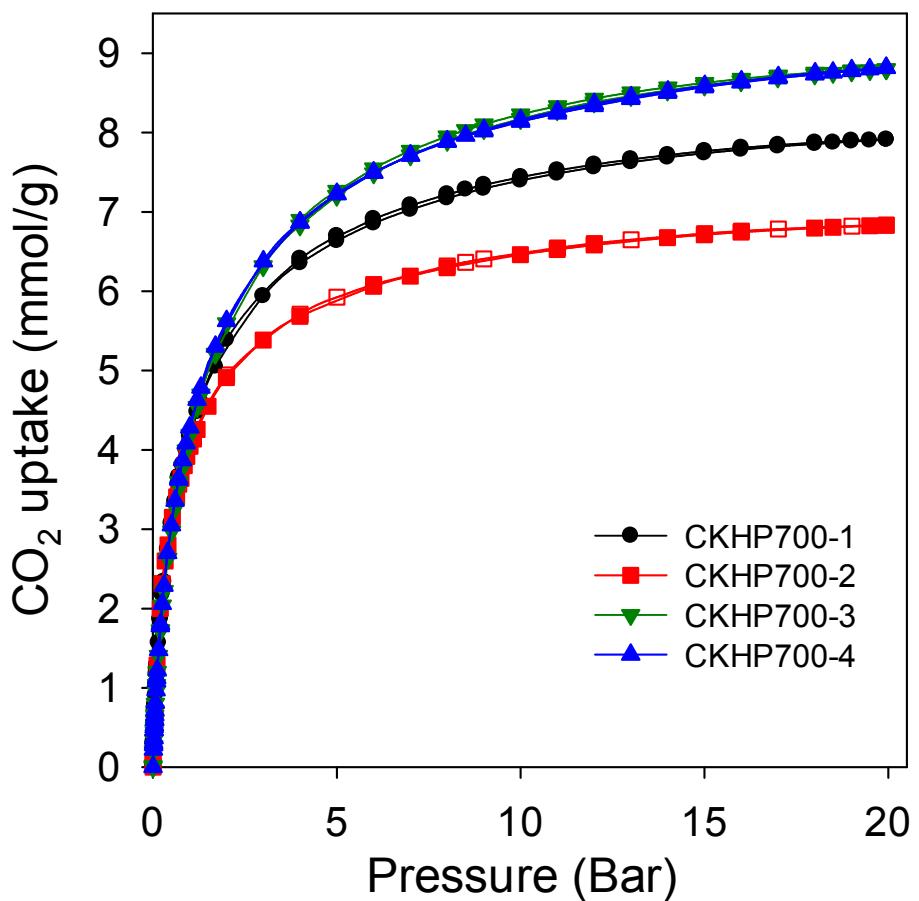
Supporting Figure S13. Low pressure CO₂ uptake isotherms at 0 °C for CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at 800 °C for 1 to 4 h.



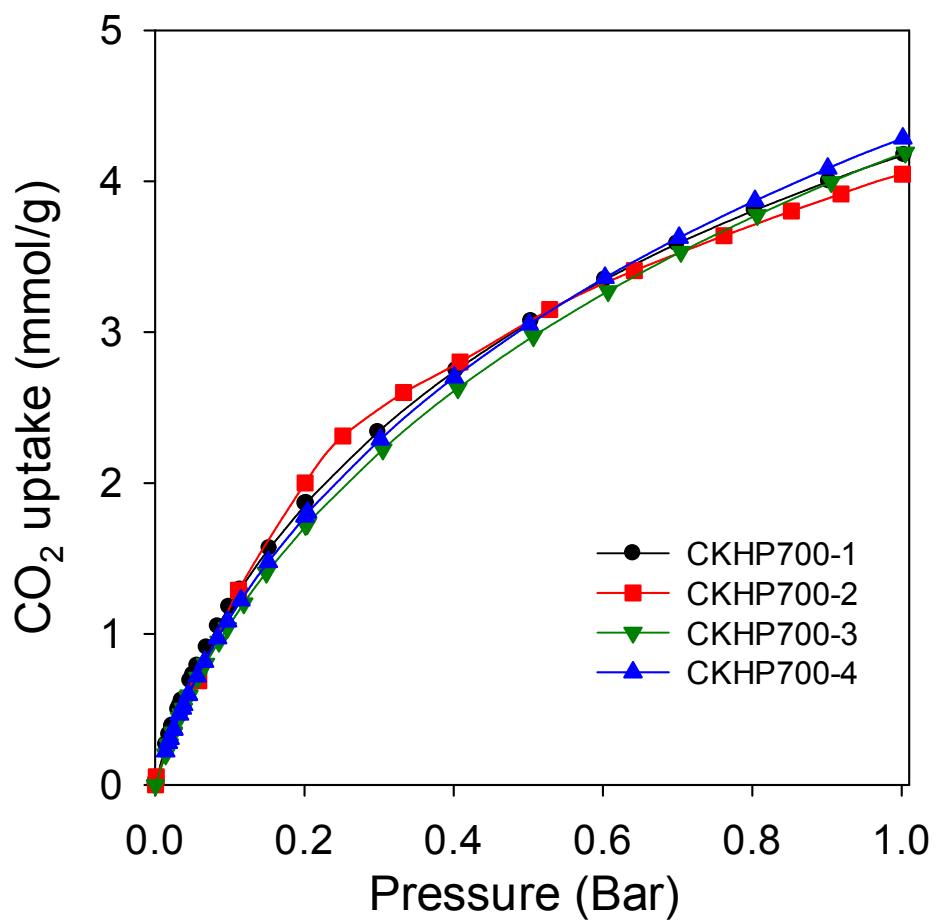
Supporting Figure S14. CO₂ uptake isotherms at 25 °C and 0-20 bar for CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at 900 °C for 1 to 4 h.



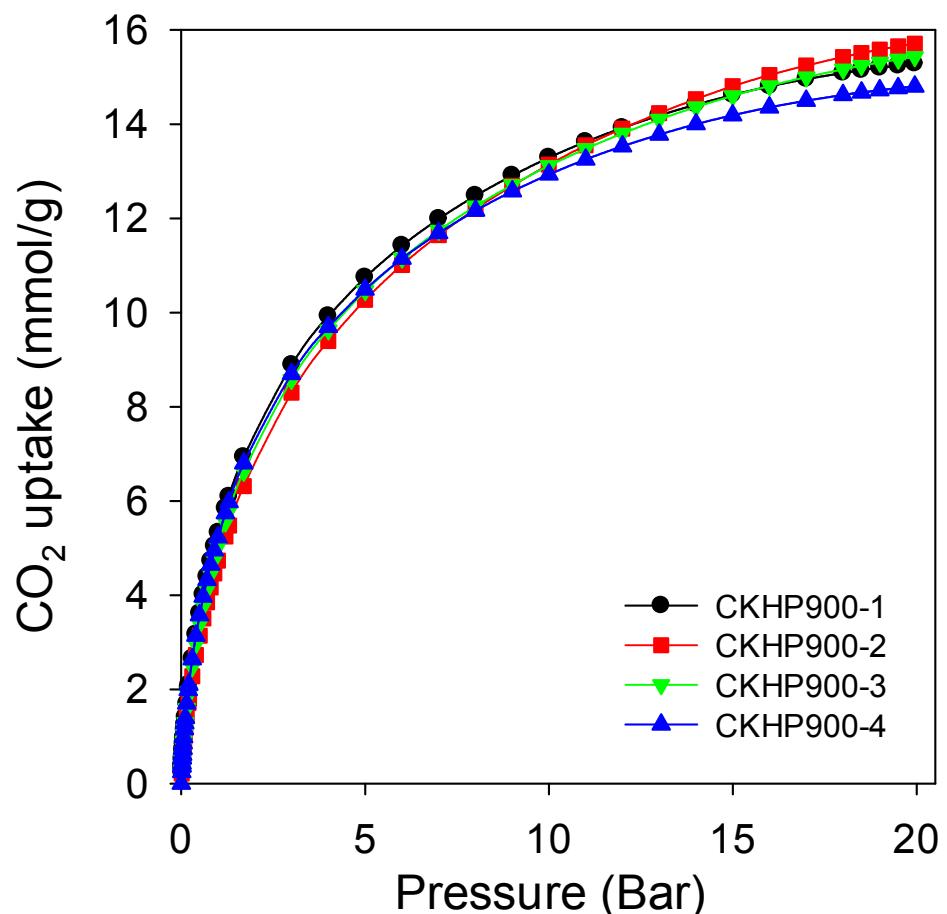
Supporting Figure S15. Low pressure CO₂ uptake isotherms at 25 °C for CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at 900 °C for 1 to 4 h.



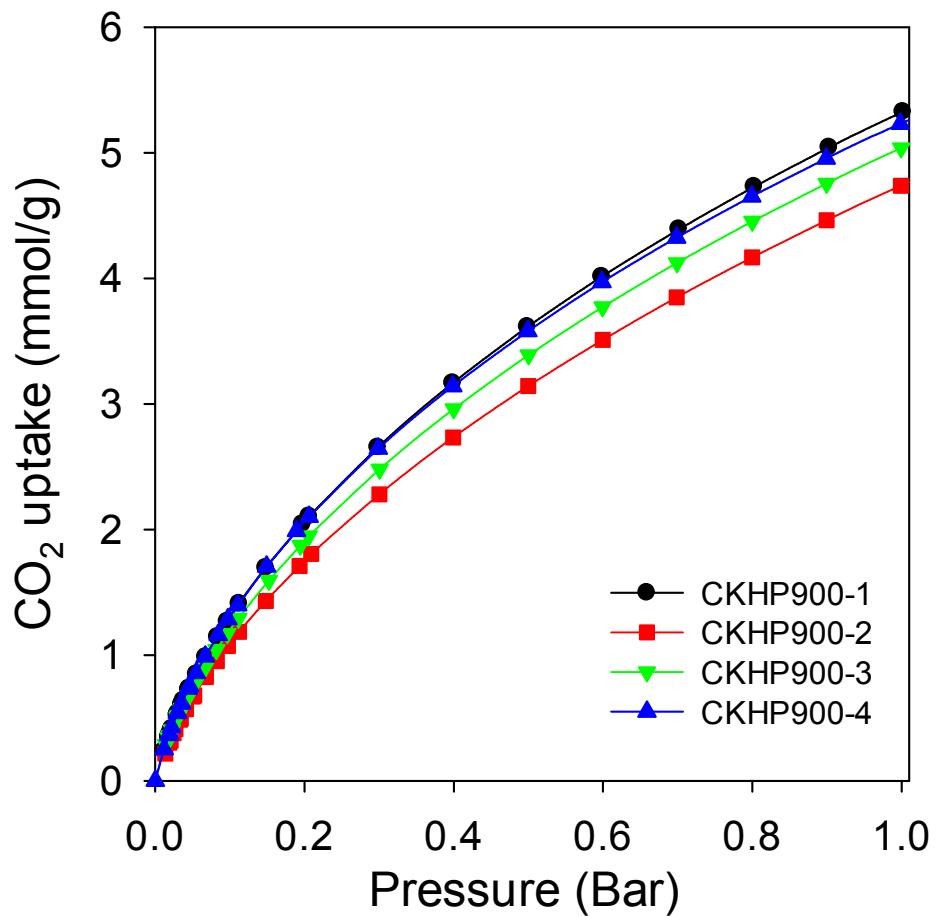
Supporting Figure S16. CO₂ uptake isotherms at 25 °C and 0 – 20 bar for CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at 700 °C for 1 to 4 h.



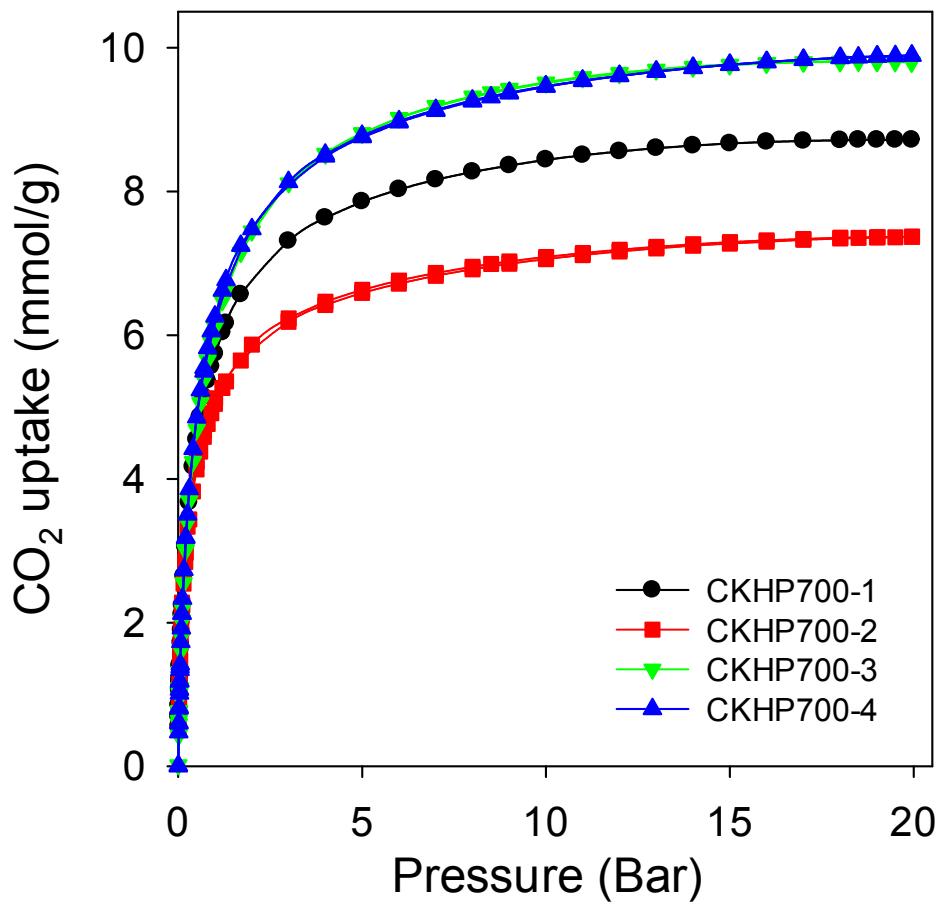
Supporting Figure S17. Low pressure CO₂ uptake isotherms at 25 °C for CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at 700 °C for 1 to 4 h.



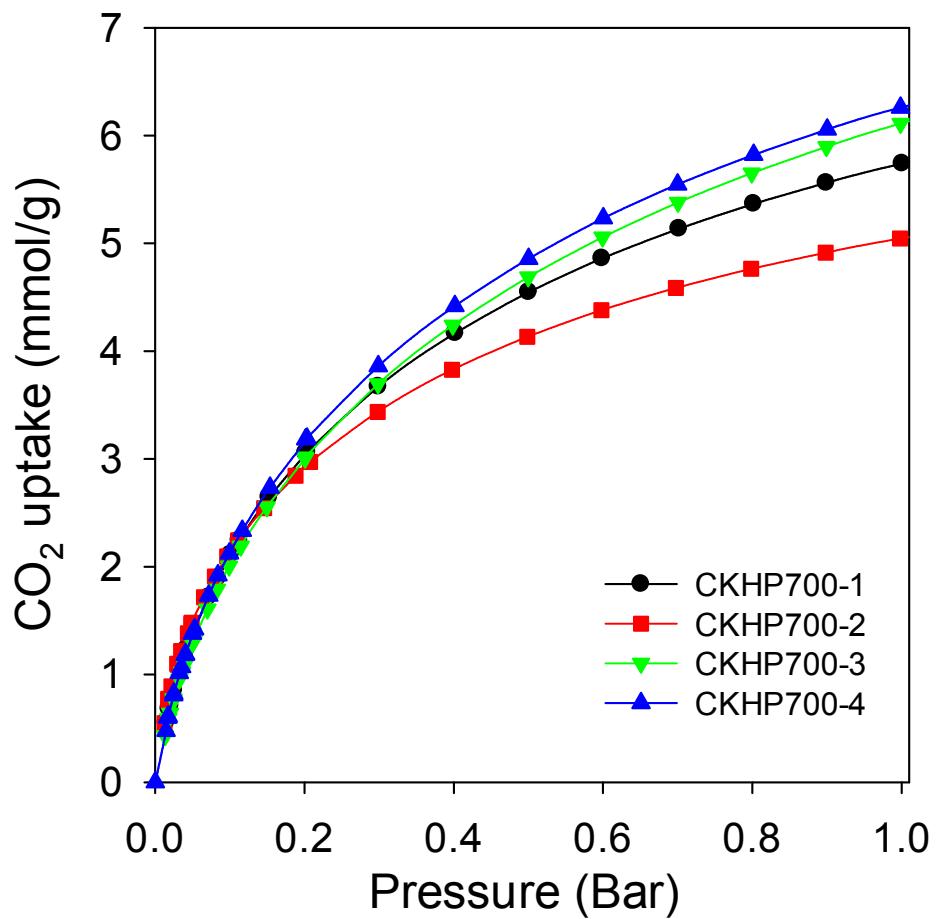
Supporting Figure S18. CO₂ uptake isotherms at 0 °C and 0 – 20 bar for CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at 900 °C for 1 to 4 h.



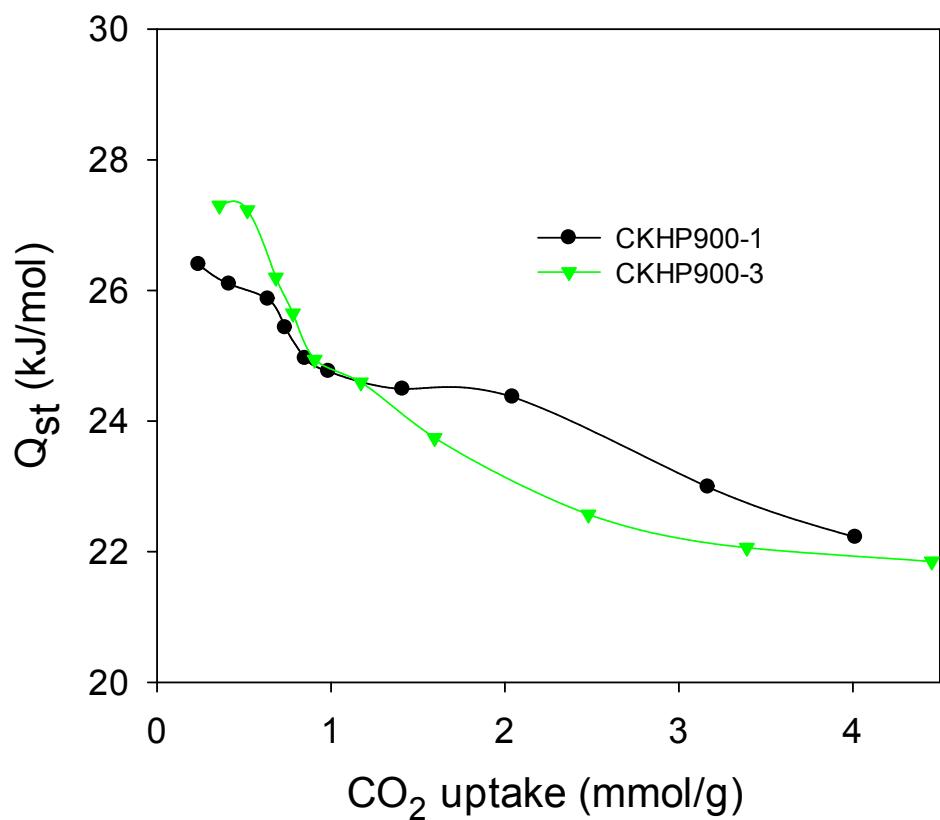
Supporting Figure S19. Low pressure CO₂ uptake isotherms at 0 °C and 0-20 bar for CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at 900 °C for 1 to 4 h.



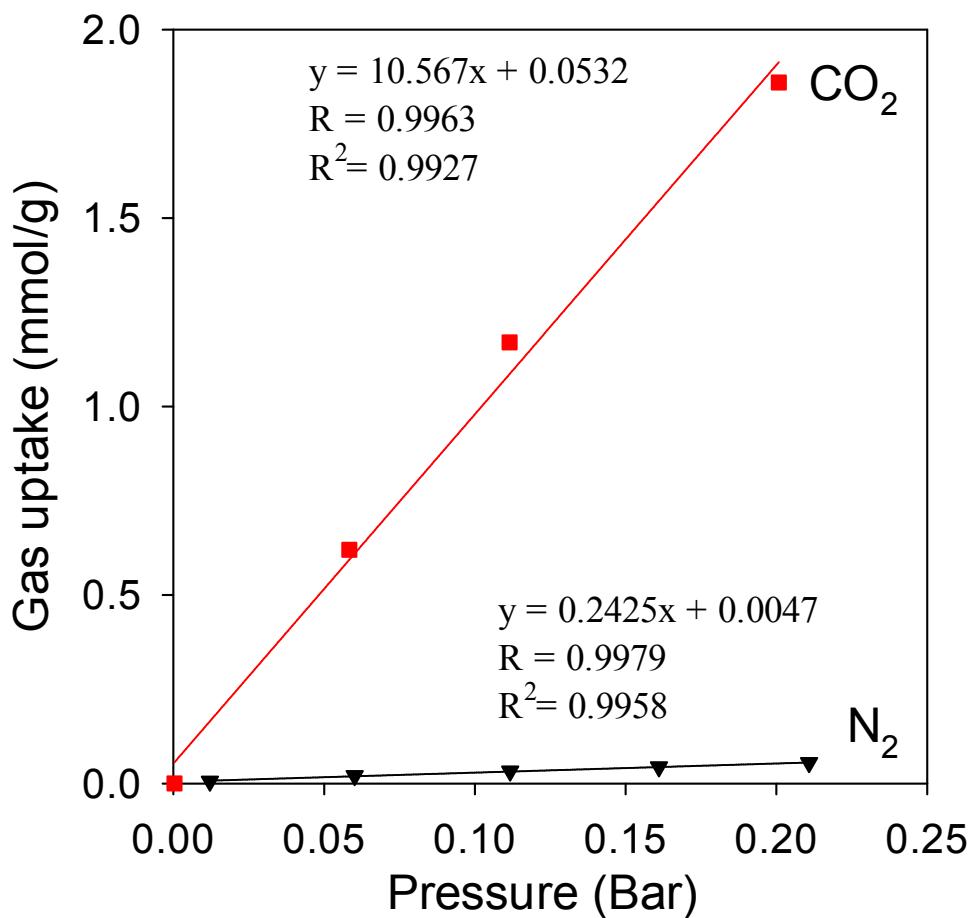
Supporting Figure S20. CO₂ uptake isotherms at 0 °C and 0 – 20 bar for CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at 700 °C for 1 to 4 h.



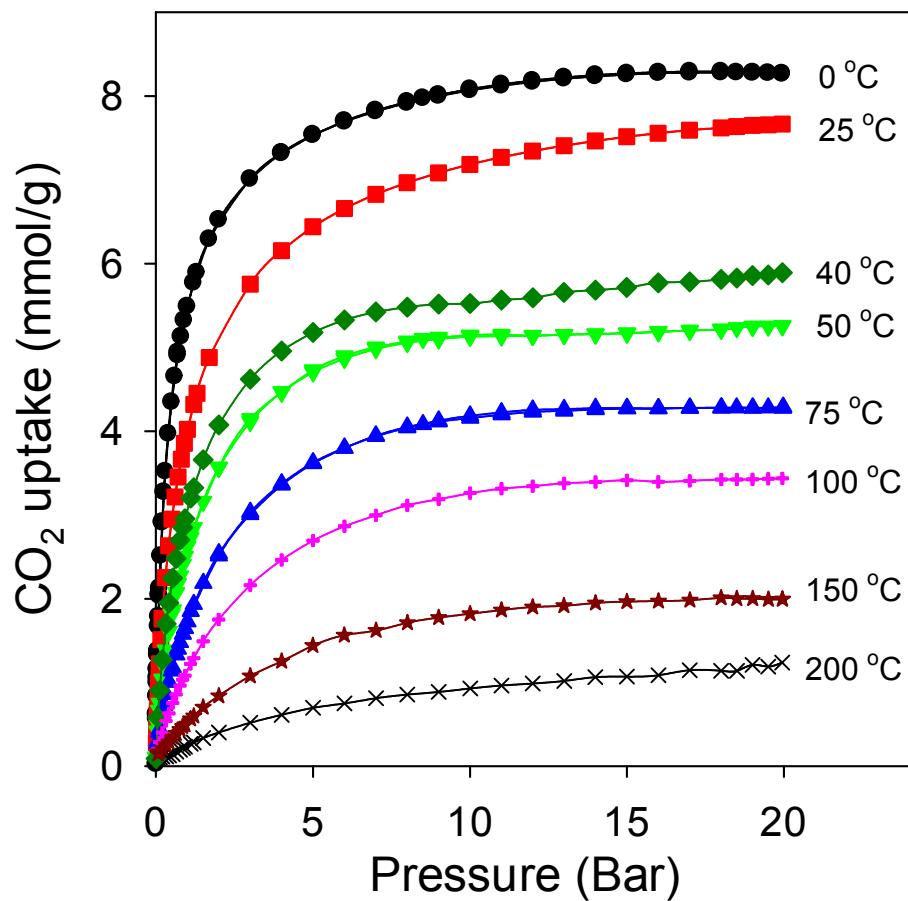
Supporting Figure S21. Low pressure CO₂ uptake isotherms at 0 °C and 0-20 bar for CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at 700 °C for 1 to 4 h.



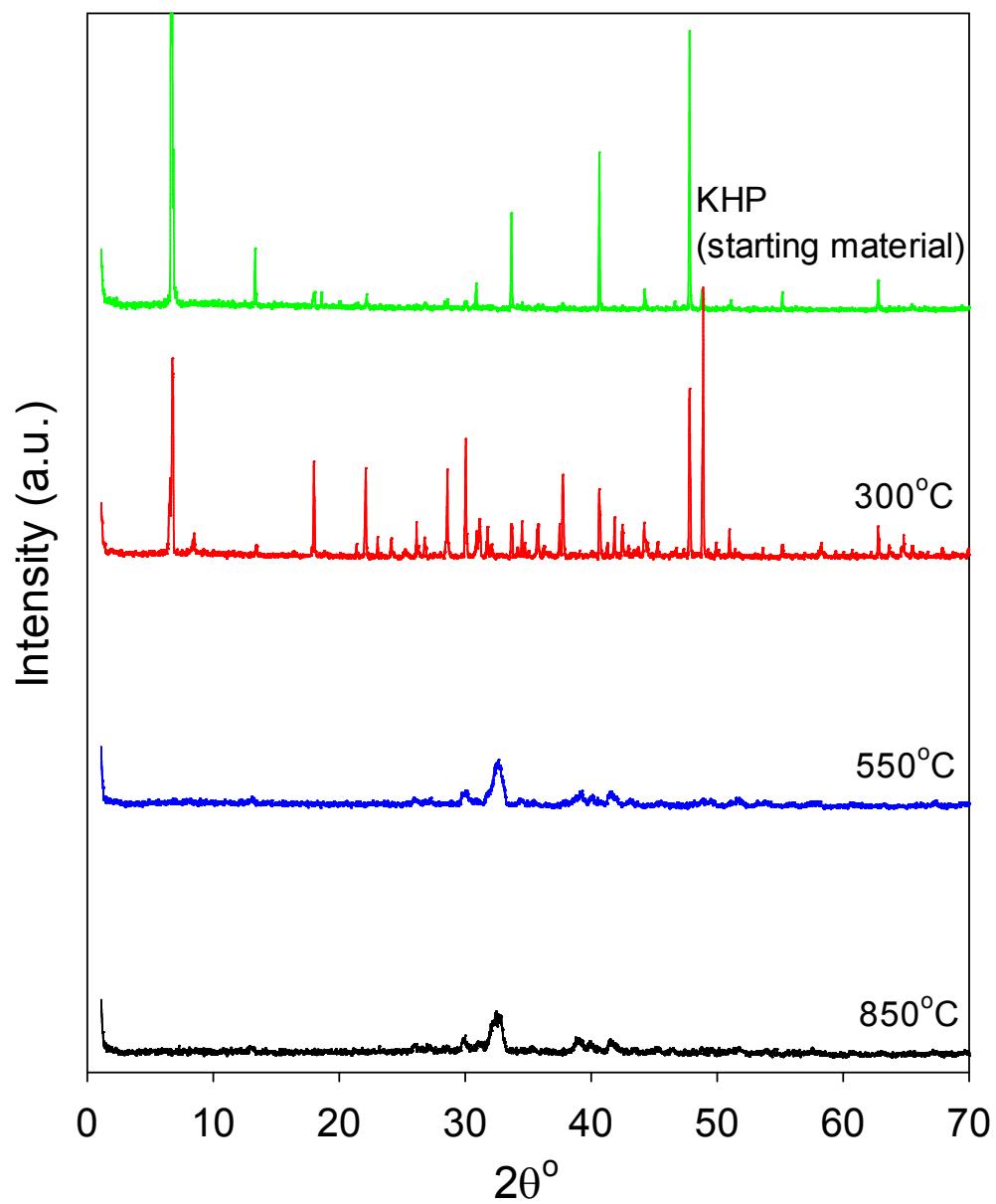
Supporting Figure S22. Isosteric heat of CO_2 adsorption (Q_{st}) as a function of CO_2 uptake for CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at 900°C for 1 or 3 h.



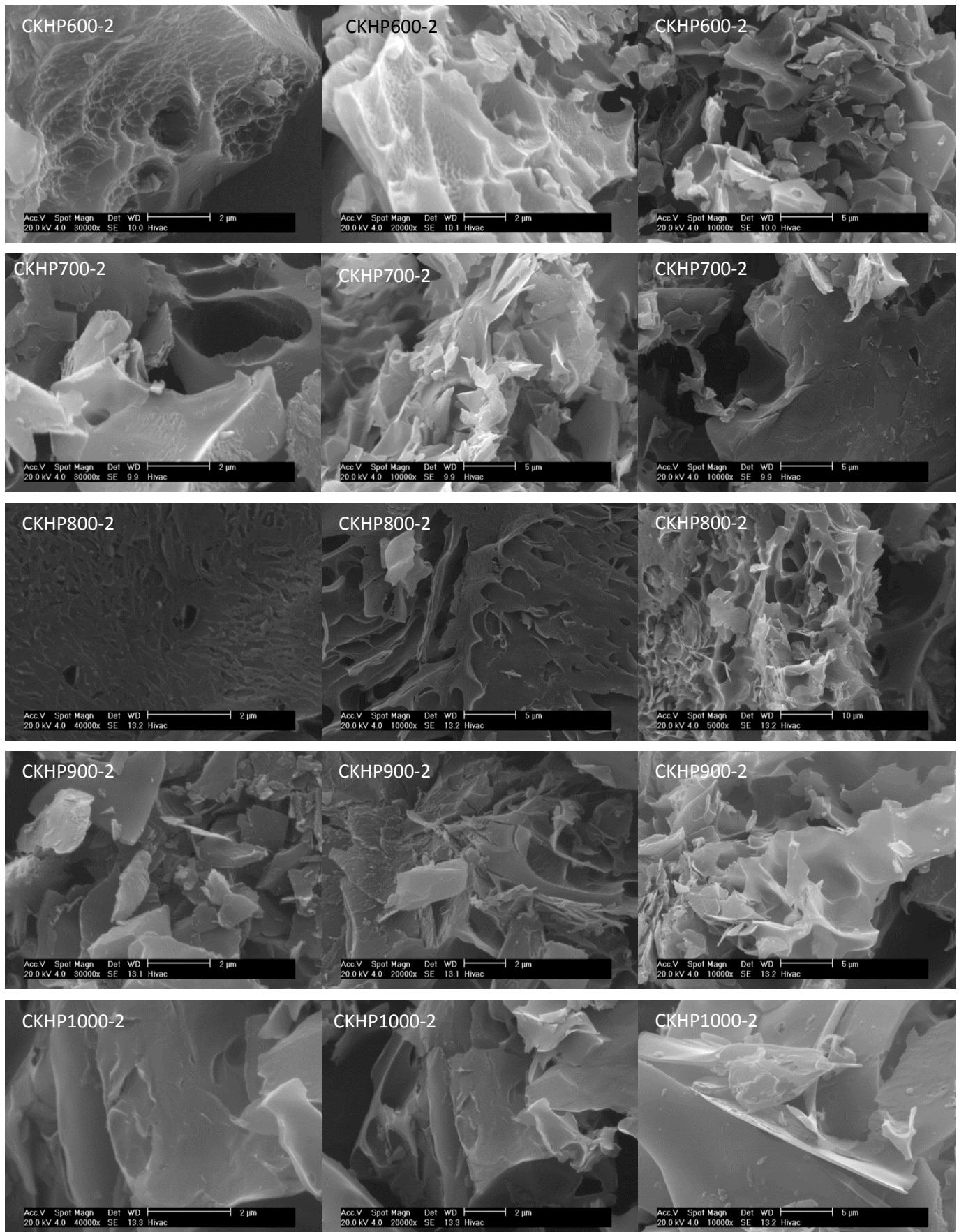
Supporting Figure S23. Comparison of initial rates of CO_2 and N_2 uptake at room temperature for sample CKHP800-2. The comparison gives a CO_2/N_2 selectivity of 43.



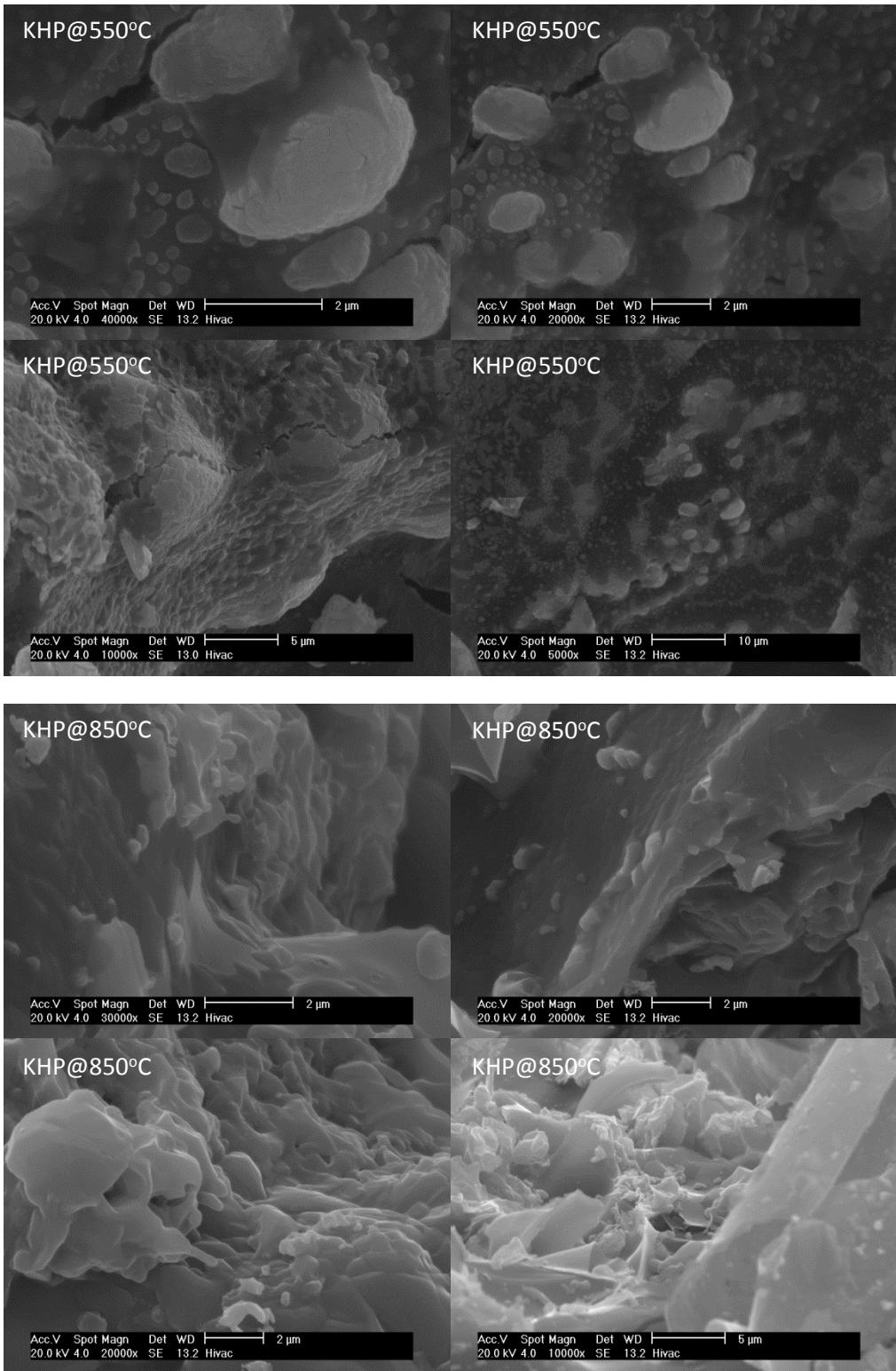
Supporting Figure S24. CO₂ uptake isotherms at various temperatures and 0 – 20 bar for sample CKHP700-1-C5.



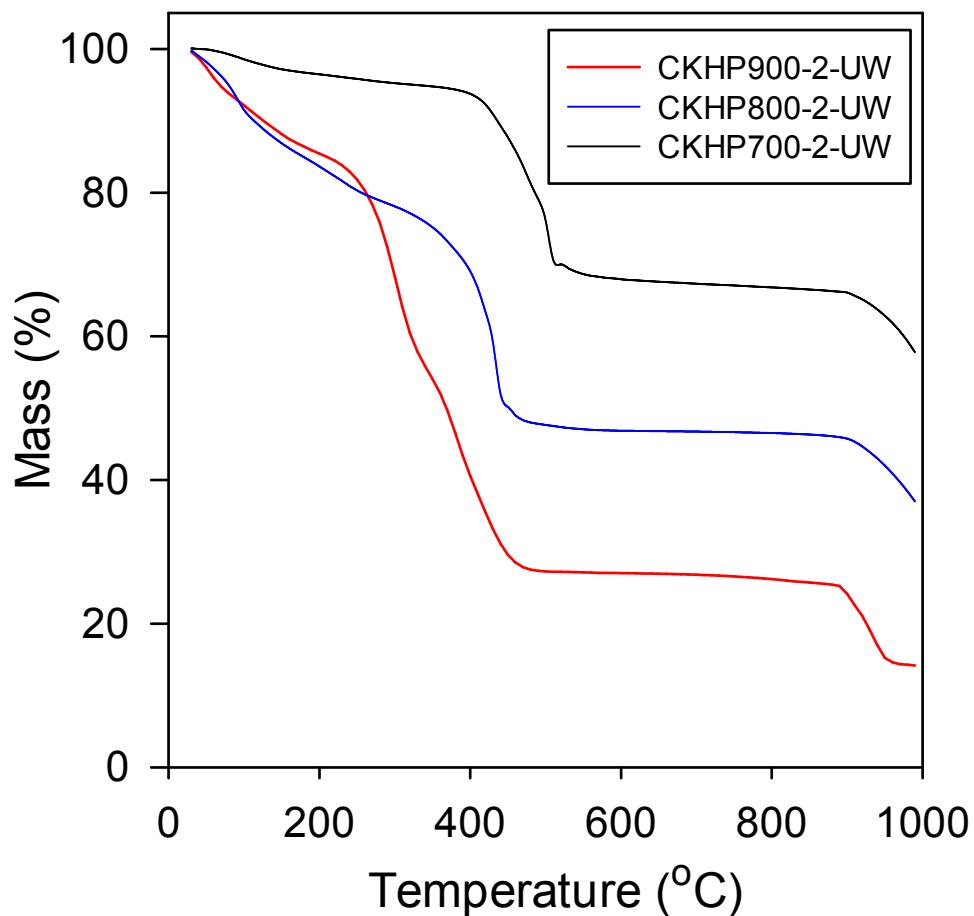
Supporting Figure S25. Powder XRD patterns of potassium hydrogen phthalate (KHP) salt and carbonisation products under nitrogen at various temperatures.



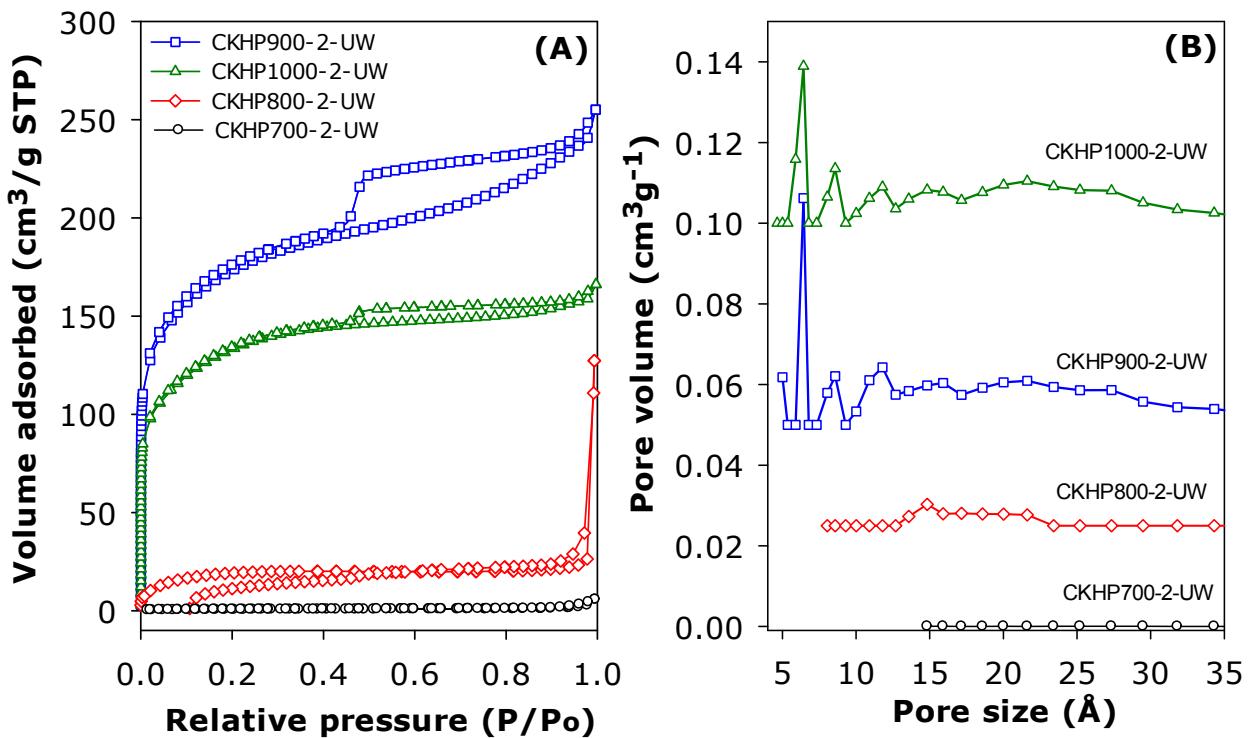
Supporting Figure S26. SEM images of CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at various temperatures for 2 h.



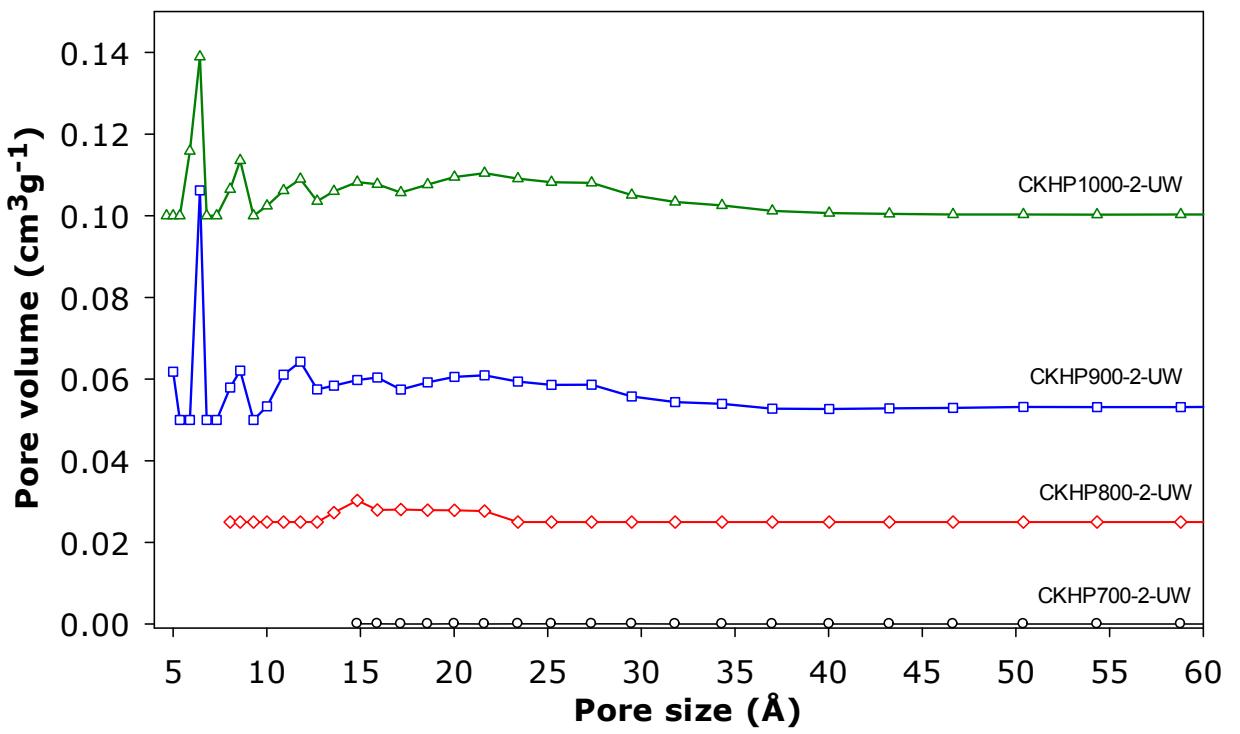
Supporting Figure S27. SEM images of carbonisation products of potassium hydrogen phthalate (KHP) salt under nitrogen at various temperatures. The carbonisation products are not washed.



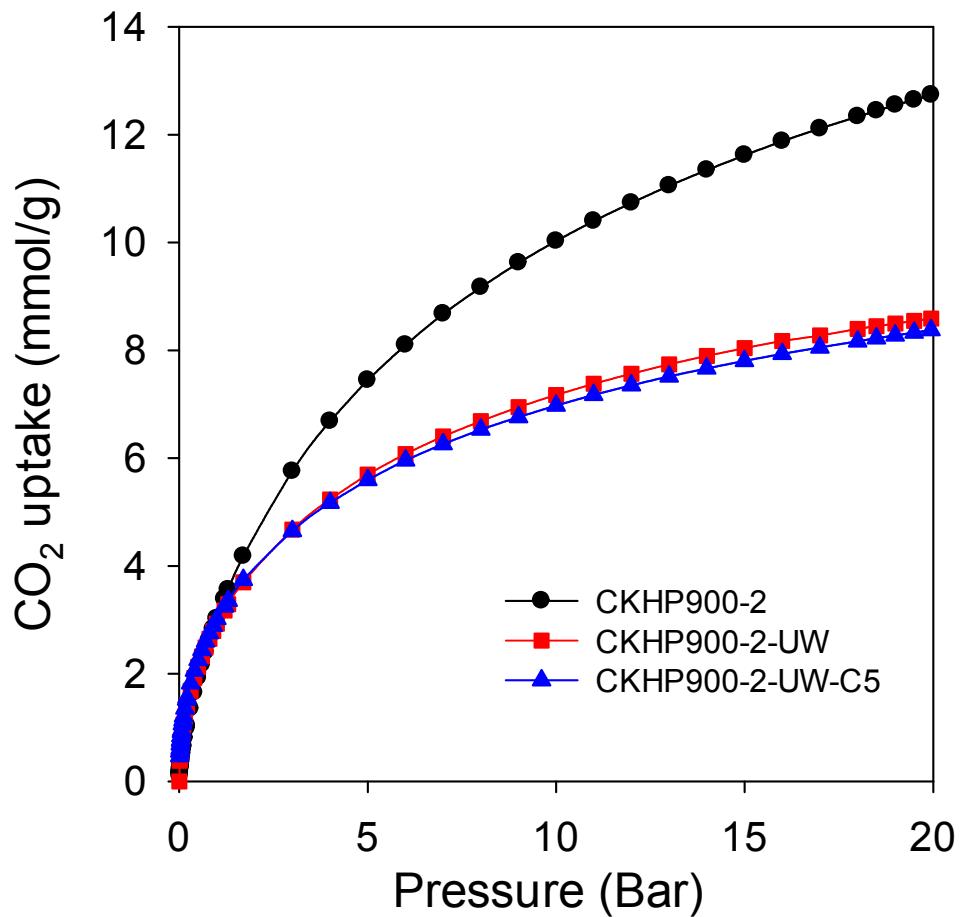
Supporting Figure S28. Thermogravimetric analysis (TGA) curves of unwashed CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at various temperatures for 2 h.



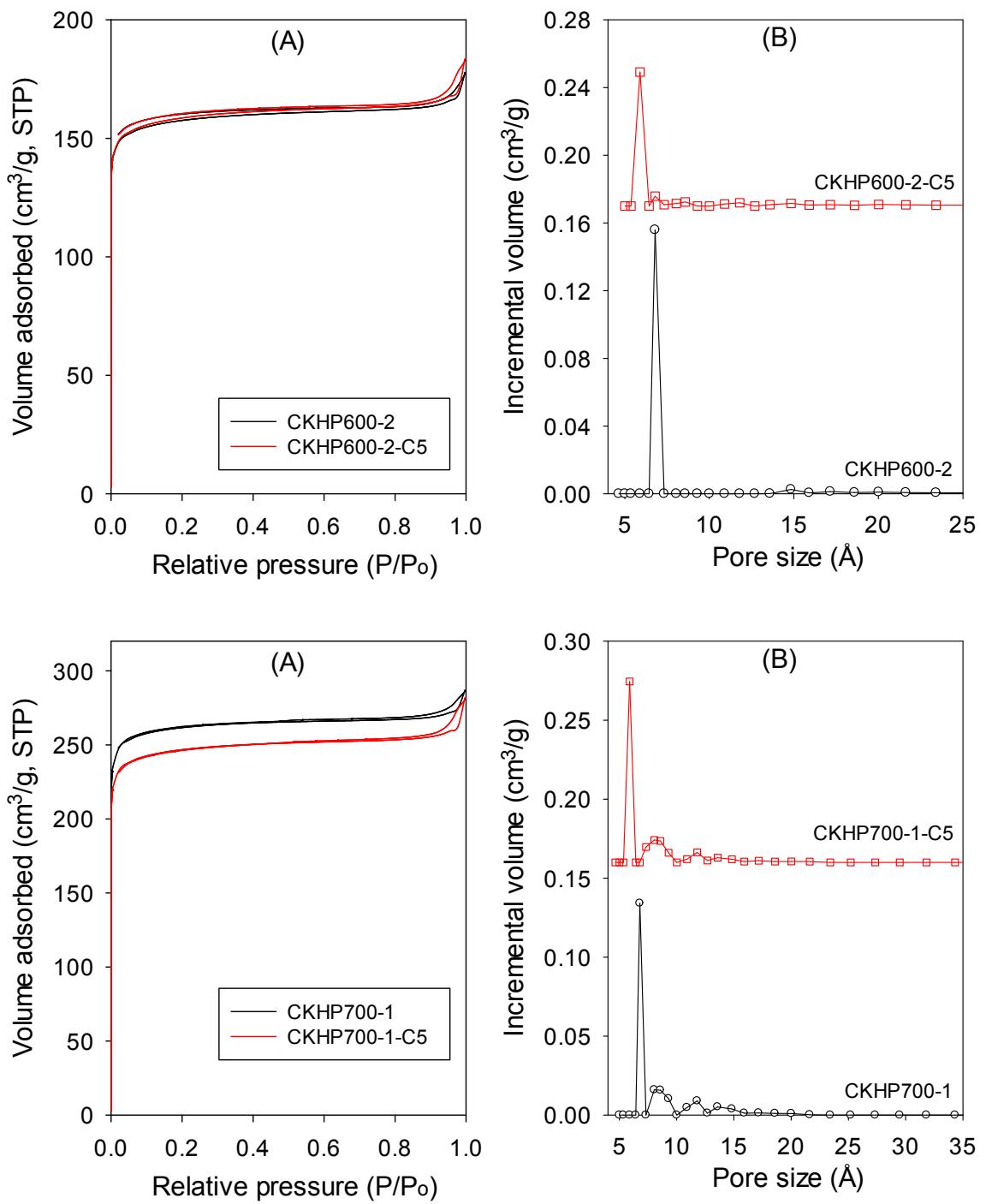
Supporting Figure S29. Nitrogen sorption isotherms (A) and corresponding pore size distribution curves (B) of unwashed CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at various temperatures for 2 h.



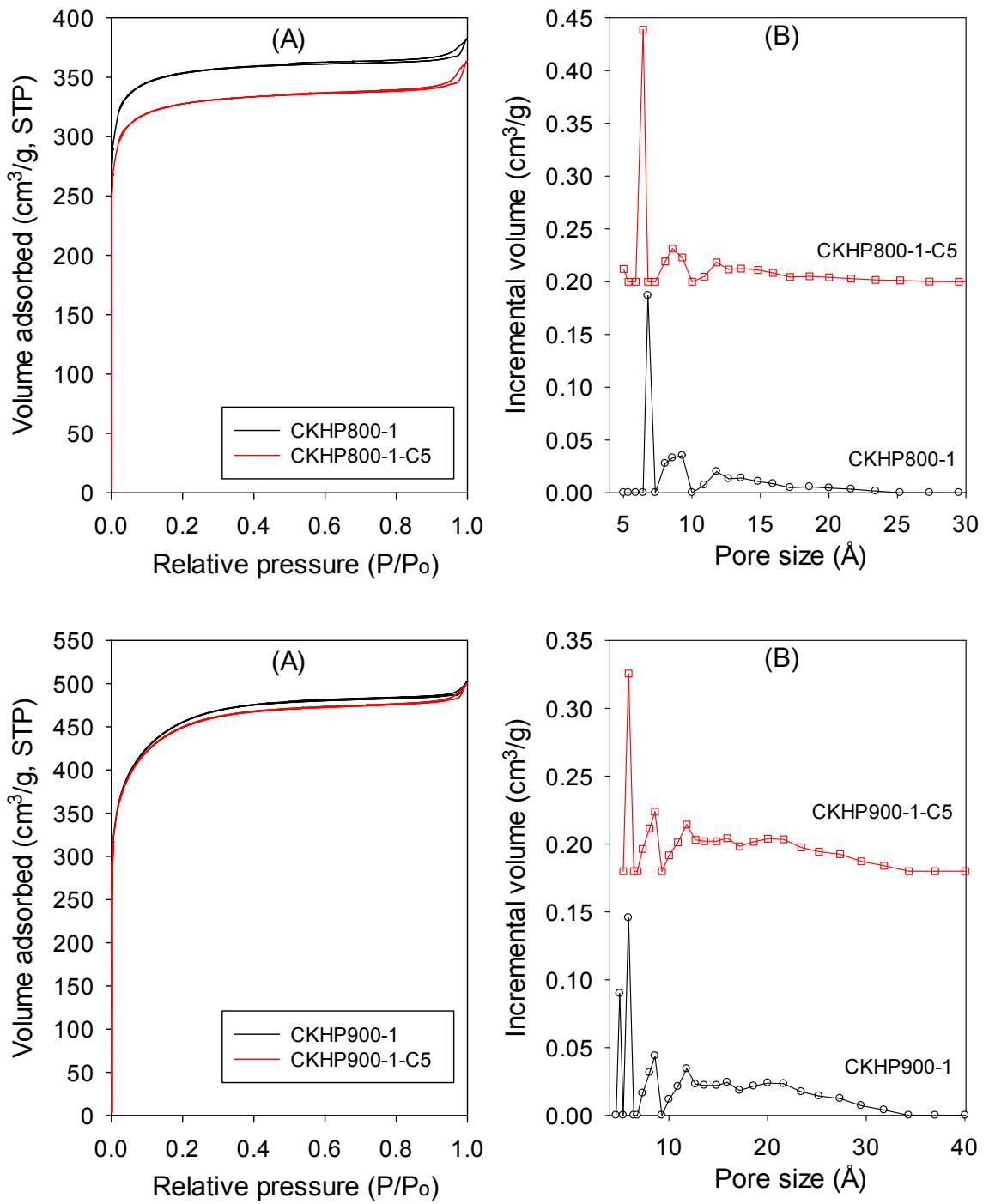
Supporting Figure S30. Pore size distribution curves of unwashed CKHP carbons derived from potassium hydrogen phthalate (KHP) via carbonisation at various temperatures for 2 h. Samples CKHP700-2-UW is non-porous, CKHP800-UW has hardly any pores, while the largest pores in samples CKHP900-2-UW and CKHP1000-UW are up to 35 Å.



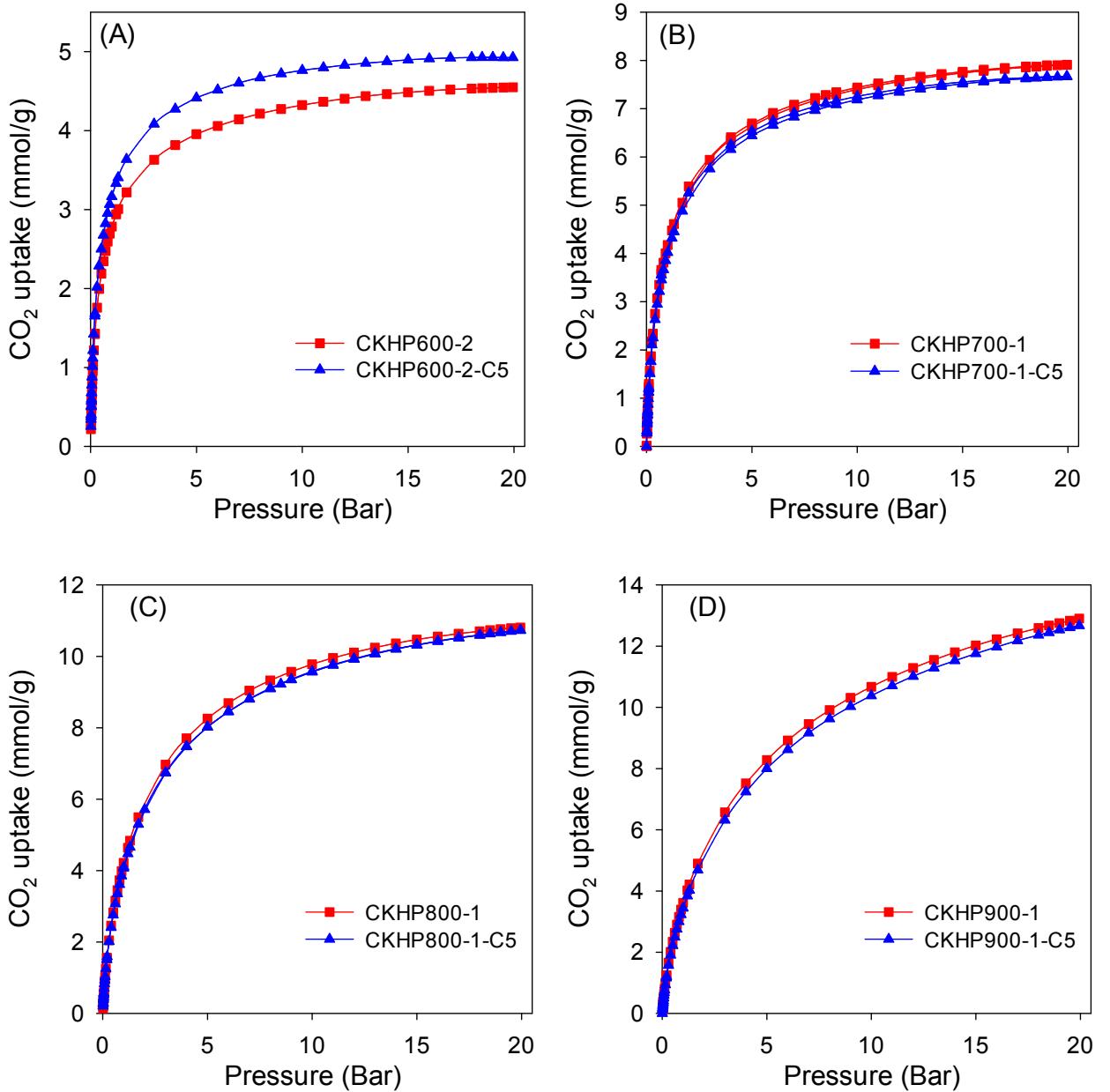
Supporting Figure S31. CO₂ uptake isotherms at 25 °C and 0 – 20 bar for washed (CKHP900-2), unwashed (CKHP900-2-UW) and unwashed compacted (CKHP900-2-UW-C5) samples carbonised at 900 °C for 2 h.



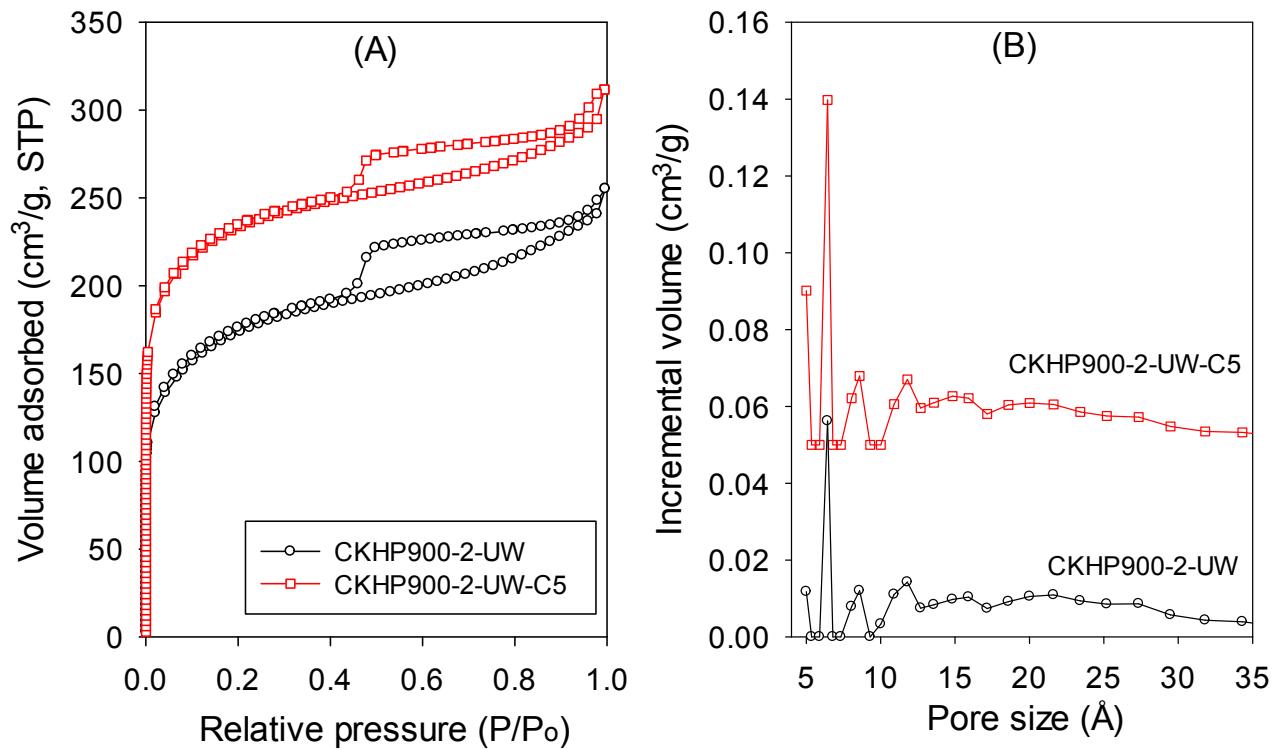
Supporting Figure S32. Nitrogen sorption isotherms (A) and pore size distribution curves (B) of CKHP carbons prepared at 600 °C (top) or 700 °C (bottom), before (CKHP600-2 and CKHP700-1) and after (CKHP600-2-C5 and CKHP700-1-C5) compaction at a load of 5 tons (i.e. 370 MPa).



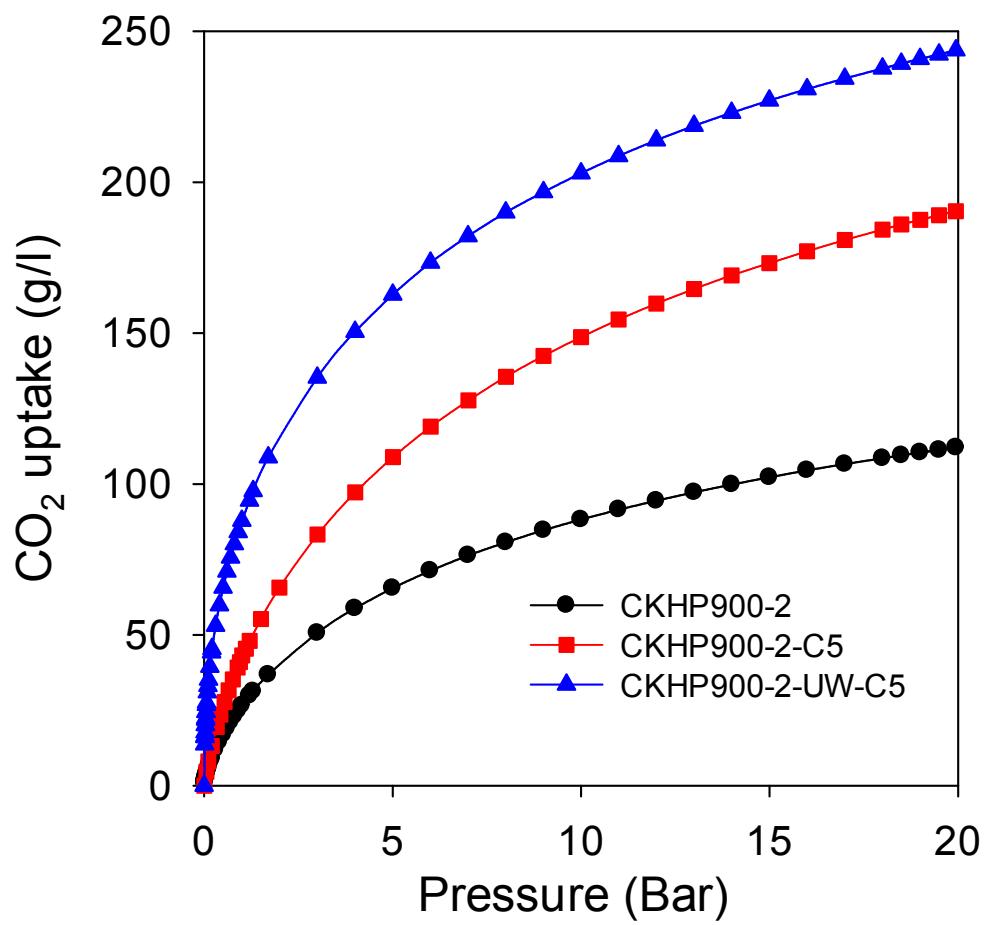
Supporting Figure S33. Nitrogen sorption isotherms (A) and pore size distribution curves (B) of CKHP carbons prepared at 800 °C (top) or 900 °C (bottom), before (CKHP800-1 and CKHP900-1) and after (CKHP800-1-C5 and CKHP900-1-C5) compaction at a load of 5 tons (i.e. 370 MPa).



Supporting Figure S34. CO₂ uptake isotherms at 25 °C and 0 – 20 bar for various CKHP carbons (A) CKHP600-2, (B) CKHP700-1, (C) CKHP800-1 and (D) CKHP900-1, before and after compaction at a load of 5 tons (i.e. 370 MPa).



Supporting Figure S35. Nitrogen sorption isotherms (A) and pore size distribution curves (B) of unwashed CKHP carbon, before (CKHP900-2-UW) and after (CKHP900-2-UW-C5) compaction at a load of 5 tons (i.e. 370 MPa).



Supporting Figure S36. Volumetric CO₂ uptake of washed (CKHP900-2), unwashed (CKHP900-2-UW) and unwashed compacted (CKHP900-2-UW-C5) samples carbonised at 900 °C for 2 h.