

## Supplementary Information

# Effect of buffer/iodide electrolyte on electrochemical capacitor performance

Amelia Klimek, Maciej Tobis, Elzbieta Frackowiak\*

Institute of Chemistry and Technical Electrochemistry, Poznan University of Technology,

Berdychowo 4, Poznan 60-965, Poland

\*Corresponding author

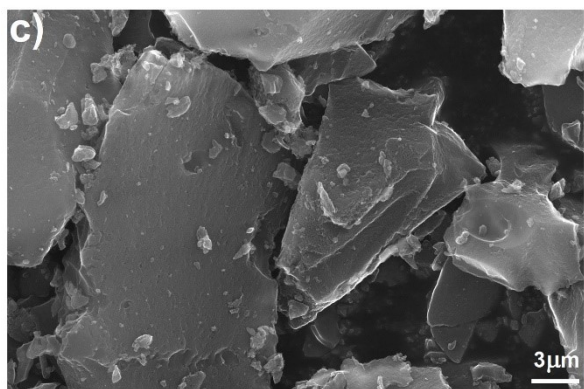
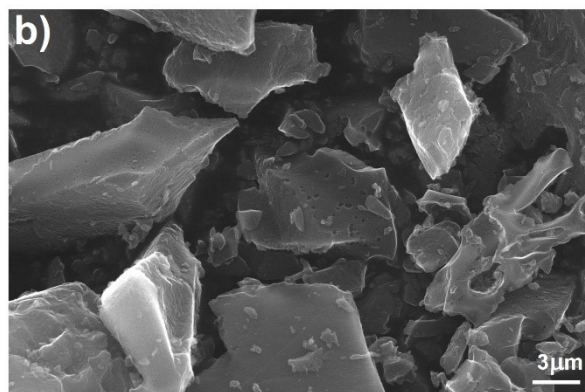
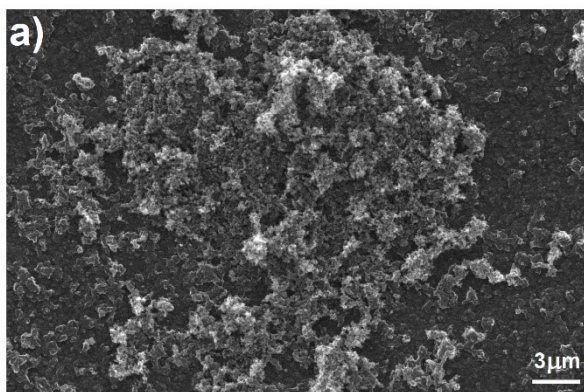
[elzbieta.frackowiak@put.poznan.pl](mailto:elzbieta.frackowiak@put.poznan.pl)

**Table S1** Specific surface area and pore volume of micro/mesoporous carbons: Kynol 5092-20 and salt templated C-ST.

	$S_{\text{BET}}$ ( $\text{m}^2 \text{g}^{-1}$ )	$V_{\text{micro}}$ ( $\text{cm}^3 \text{g}^{-1}$ )	$V_{\text{meso}}$ ( $\text{cm}^3 \text{g}^{-1}$ )
<b>Kynol 5092-20</b>	2100	0.80	0.05
<b>C-ST</b>	2640	0.97	0.16

**Table S2** Diameters of selected solvated ions:  $\text{Na}^+$ ,  $\text{I}^-$ ,  $\text{HPO}_4^{2-}$ ,  $\text{CH}_3\text{COO}^-$ ,  $\text{C}_3\text{H}_4(\text{OH})(\text{COO})_3^{3-}$  [46-49]

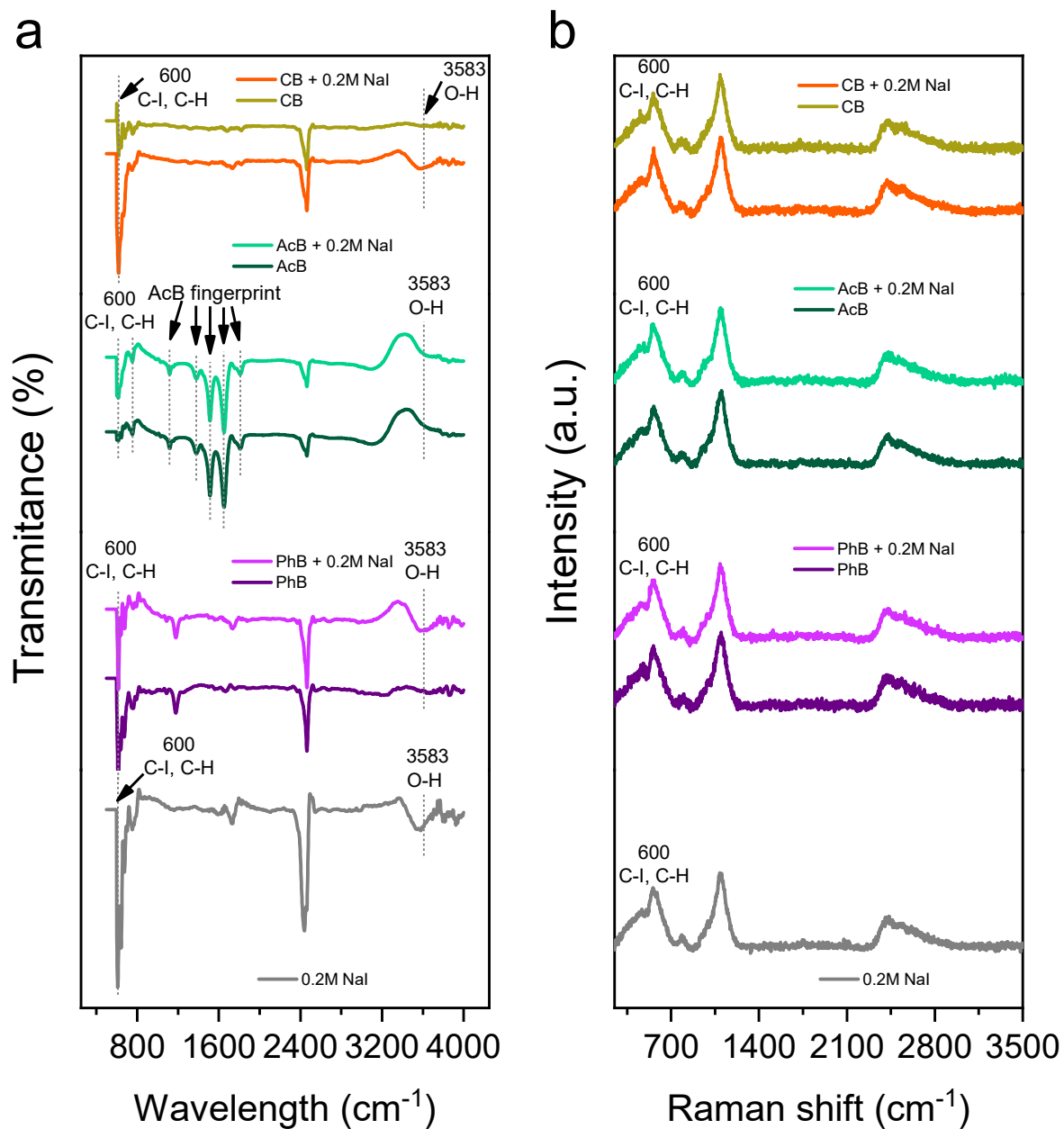
<b>Ion</b>	<b>Solvated ion diameter (nm)</b>
<b><math>\text{Na}^+</math></b>	0.486
<b><math>\text{I}^-</math></b>	0.600
<b><math>\text{HPO}_4^{2-}</math></b>	0.604
<b><math>\text{CH}_3\text{COO}^-</math></b>	0.706
<b><math>\text{C}_3\text{H}_4(\text{OH})(\text{COO})_3^{3-}</math></b>	1.099



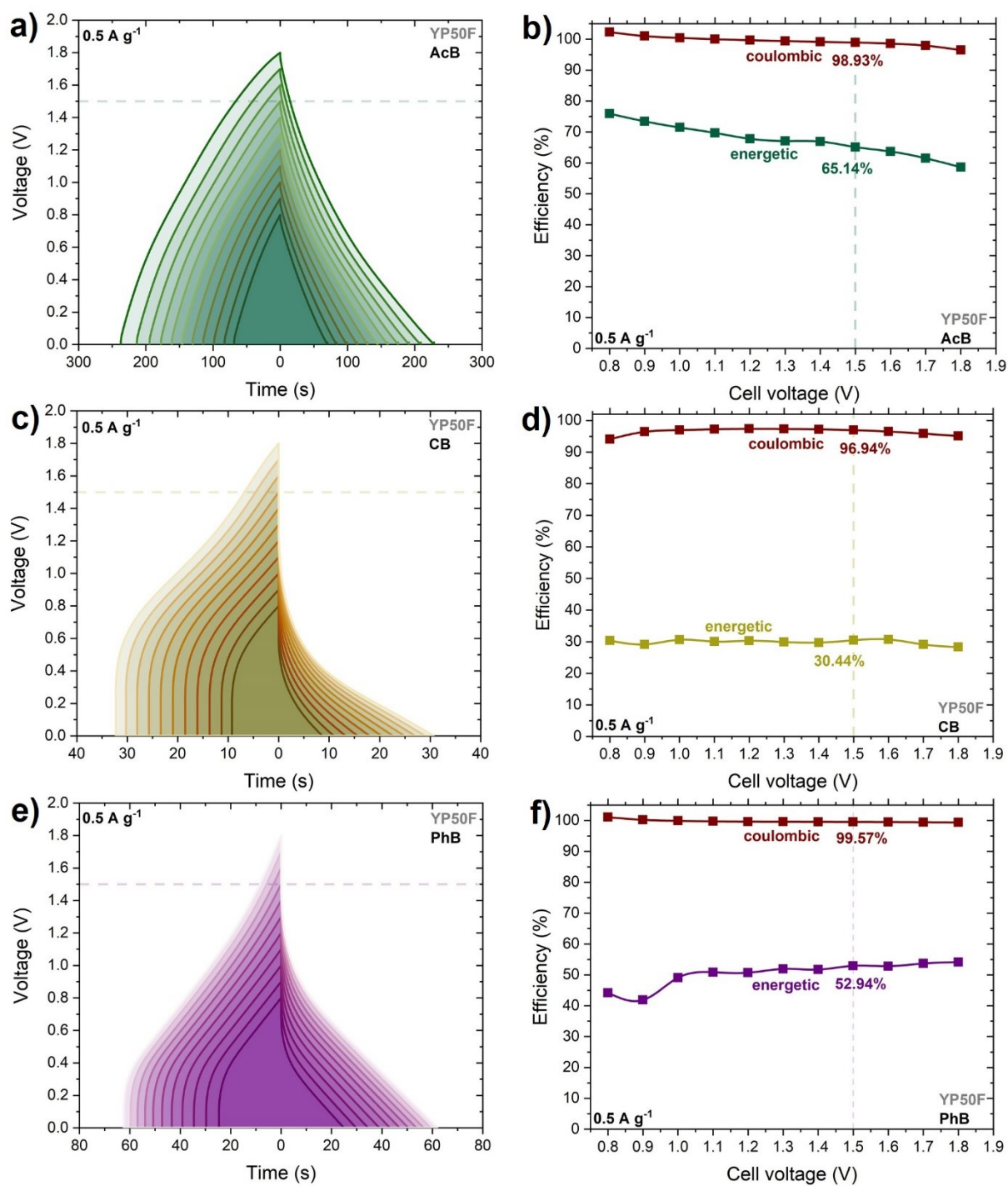
**Fig. S1** SEM images of **a)** BP2000 activated carbon; **b)** YP80F activated carbon; **c)** YP50F activated carbon.

**Table S3** Conductivity ( $\text{mS cm}^{-1}$ ) and pH measurements of sodium iodide ( $0.2 \text{ mol L}^{-1}$ ) and acetate buffer (AcB), citrate buffer (CB), phosphate buffer (PhB) with/without additive of sodium iodide ( $0.2, 0.5, 1, 2 \text{ mol L}^{-1}$ ).

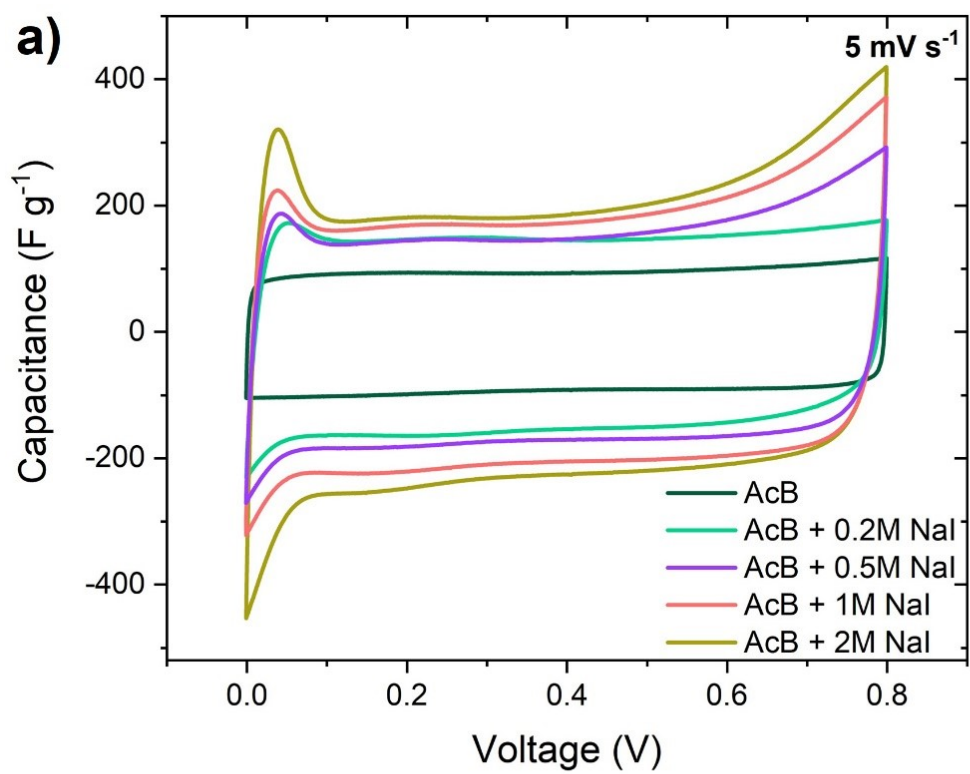
<b>Electrolyte</b>	<b>Conductivity (<math>\text{mS cm}^{-1}</math>)</b>	<b>pH</b>
<b><math>0.2 \text{ mol L}^{-1} \text{ NaI}</math></b>	20.4	8.5
<b>AcB</b>	63.3	5.08
<b>AcB + <math>0.2 \text{ mol L}^{-1} \text{ NaI}</math></b>	72.2	5.05
<b>AcB + <math>0.5 \text{ mol L}^{-1} \text{ NaI}</math></b>	84.5	5.24
<b>AcB + <math>1 \text{ mol L}^{-1} \text{ NaI}</math></b>	102.4	4.94
<b>AcB + <math>2 \text{ mol L}^{-1} \text{ NaI}</math></b>	132.4	4.94
<b>CB</b>	3.0	3.74
<b>CB + <math>0.2 \text{ mol L}^{-1} \text{ NaI}</math></b>	16.5	3.58
<b>CB + <math>0.5 \text{ mol L}^{-1} \text{ NaI}</math></b>	41.2	3.48
<b>CB + <math>1 \text{ mol L}^{-1} \text{ NaI}</math></b>	89.0	3.30
<b>CB + <math>2 \text{ mol L}^{-1} \text{ NaI}</math></b>	76.0	3.25
<b>PhB</b>	6.95	7.18
<b>PhB + <math>0.2 \text{ mol L}^{-1} \text{ NaI}</math></b>	25.5	6.98
<b>PhB + <math>0.5 \text{ mol L}^{-1} \text{ NaI}</math></b>	51.9	6.77
<b>PhB + <math>1 \text{ mol L}^{-1} \text{ NaI}</math></b>	92.1	6.55
<b>PhB + <math>2 \text{ mol L}^{-1} \text{ NaI}</math></b>	154.7	6.24



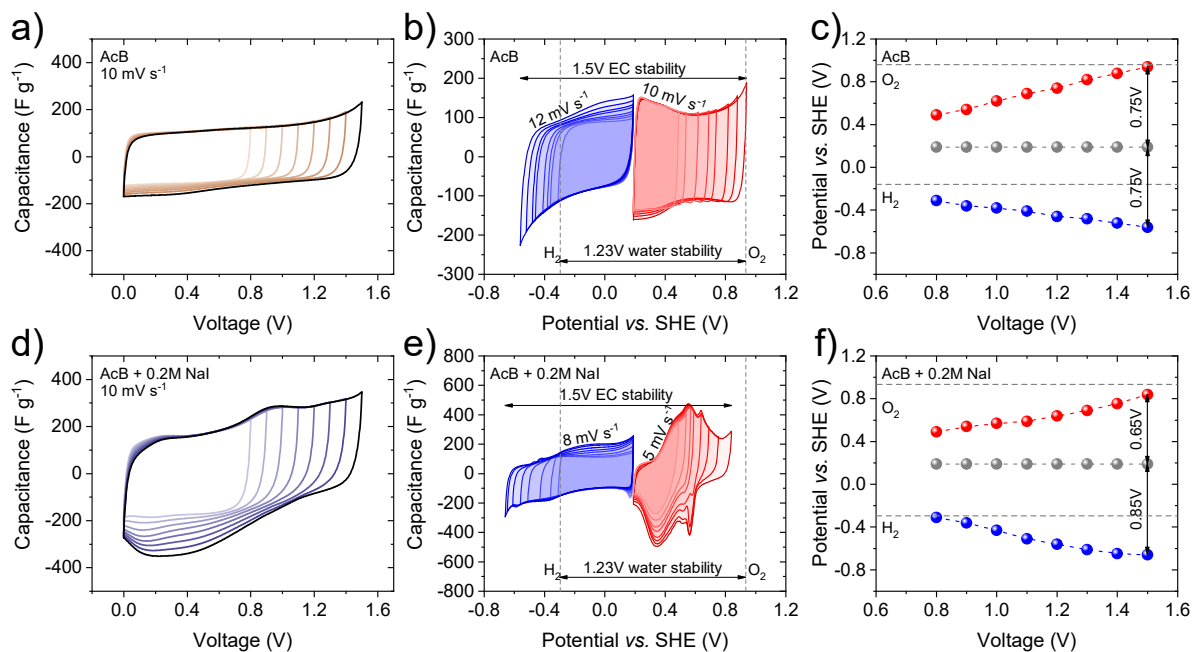
**Fig. S2 a)** FT-IR and **b)** Raman spectra for the electrolyte solutions with indicated stretching and vibrational modes.



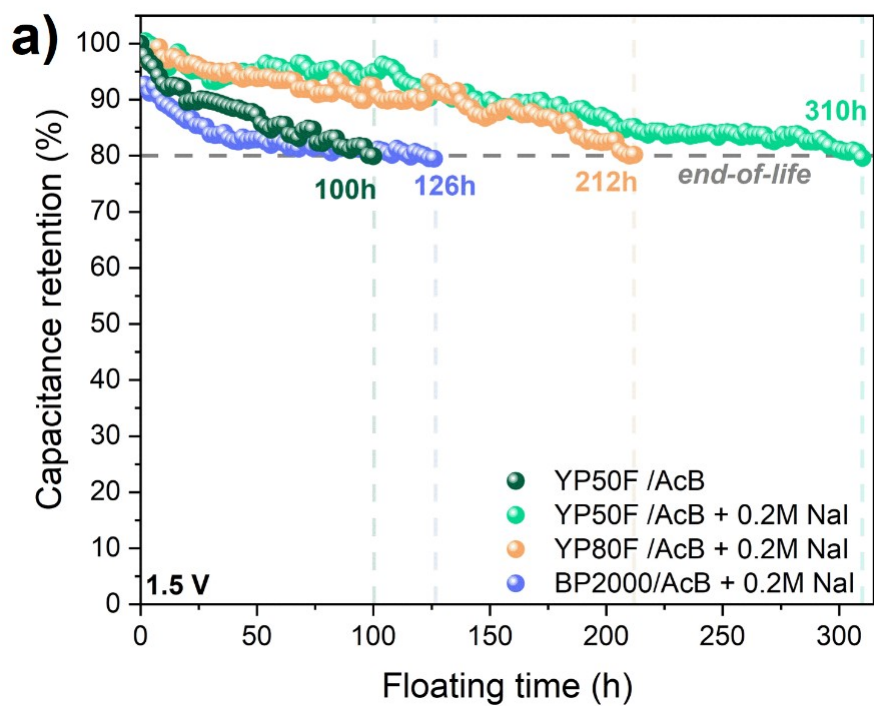
**Fig. S3** Galvanostatic charge/discharge voltage extension at 0.5 A g<sup>-1</sup> for EC operating in **a)** acetate buffer; **c)** citrate buffer; **e)** phosphate buffer for various voltages. Energetic and coulombic efficiency calculated from GCD curves for EC operating in **b)** acetate buffer; **d)** citrate buffer; **f)** phosphate buffer.



**Fig. S4** Cyclic voltammograms recorded at  $5 mV s^{-1}$  for ECs based on YP50F operating in acetate buffer without and with NaI additive of several concentrations (0.2, 0.5, 1, 2 mol  $L^{-1}$ ).

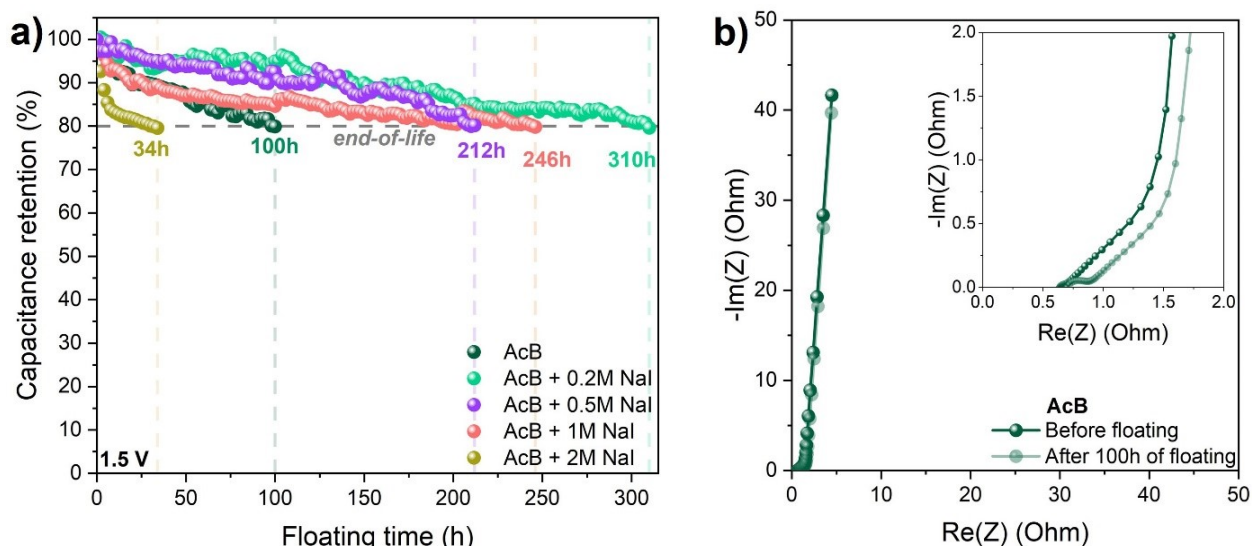


**Fig. S5** Electrochemical studies of three-electrode Swagelok cell with reference while monitoring the potential of positive and negative electrode operating in different electrolytes. Cyclic voltammograms of cells operating in AcB and AcB + 0.2 mol L<sup>-1</sup> NaI **a), d)**. Electrochemical behavior of electrodes during CV measurements **b), e)**. Cut-off potential range for positive and negative electrodes **c), f)**.

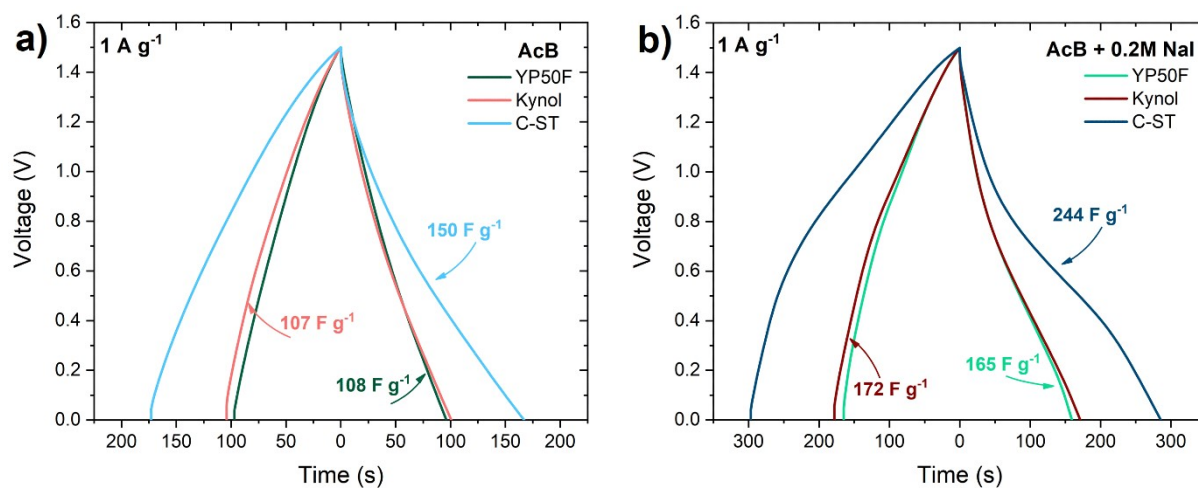


**Fig. S6 a)** Capacitance retention of different carbons micro/mesoporous carbons (YP50F, YP80F, BP2000) based ECs operating in acetate buffer with/without 0.2 mol L<sup>-1</sup> NaI additive during floating (at 1.5 V).

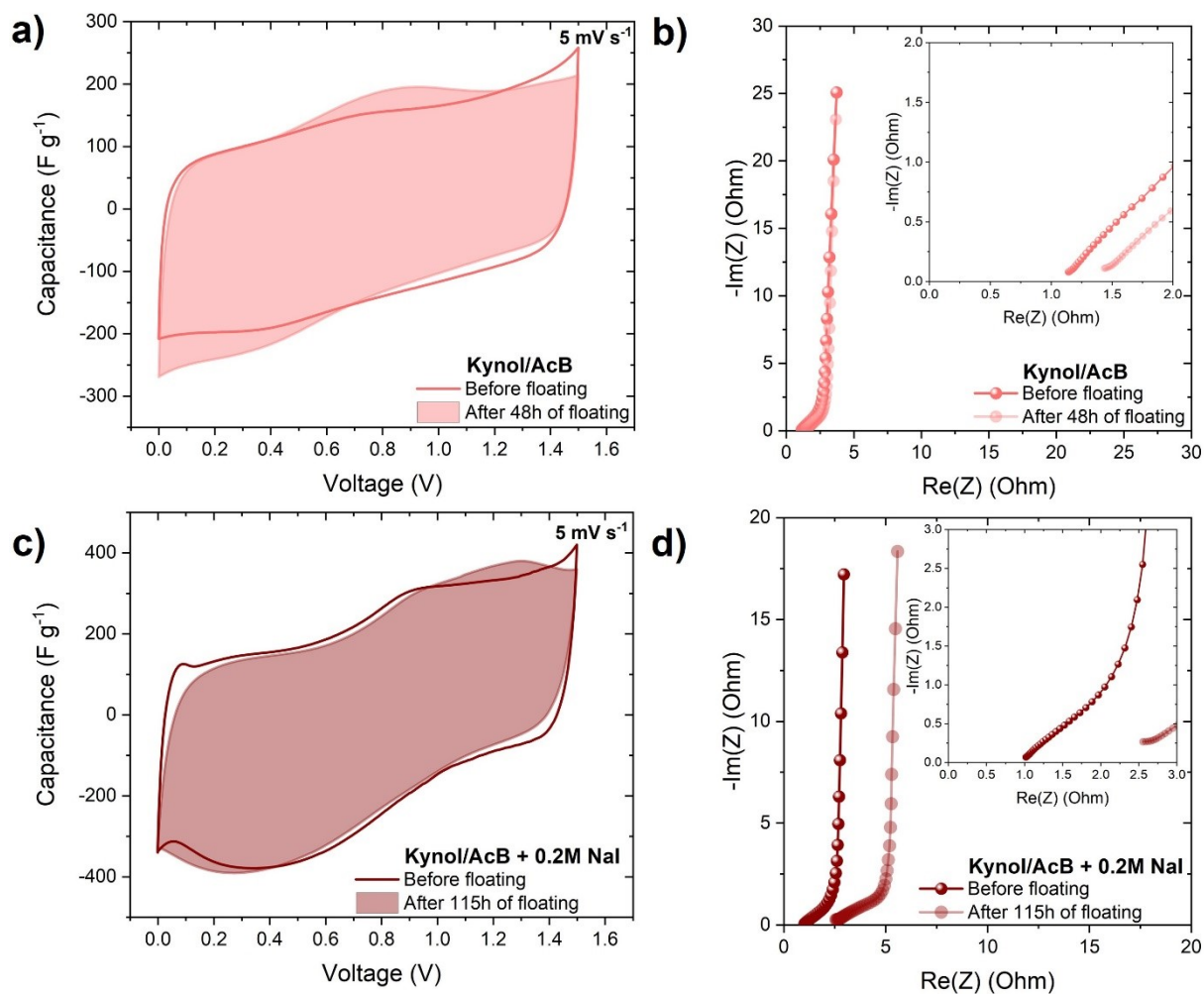




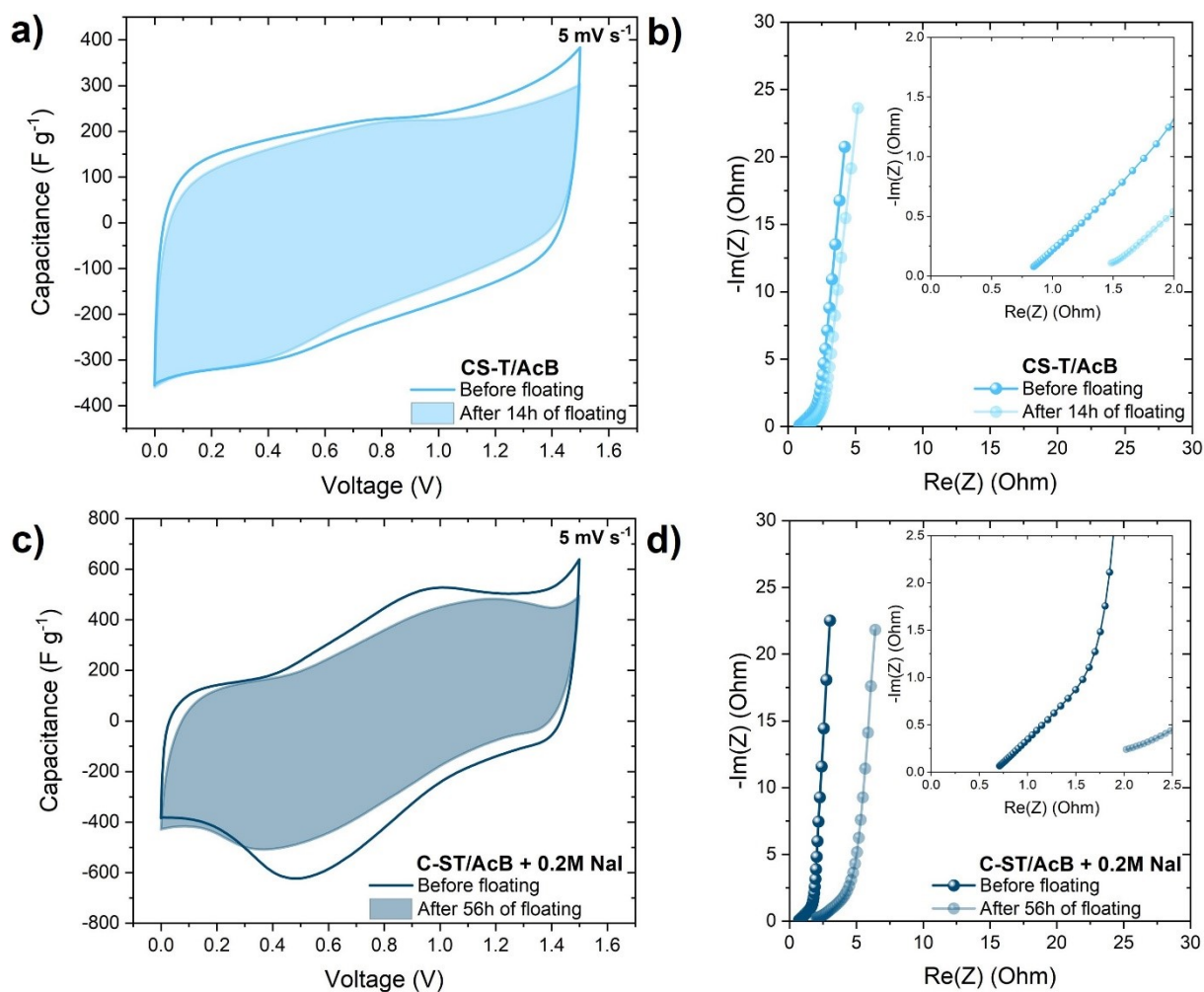
**Fig. S7 a)** Capacitance retention of YP50F based ECs operating in acetate buffer with/without 0.2, 0.5, 1, and 2 mol L<sup>-1</sup> NaI additive during floating (at 1.5 V); **b)** Nyquist plots obtained for YP50F based ECs operating in acetate buffer before and after 100 h of floating.



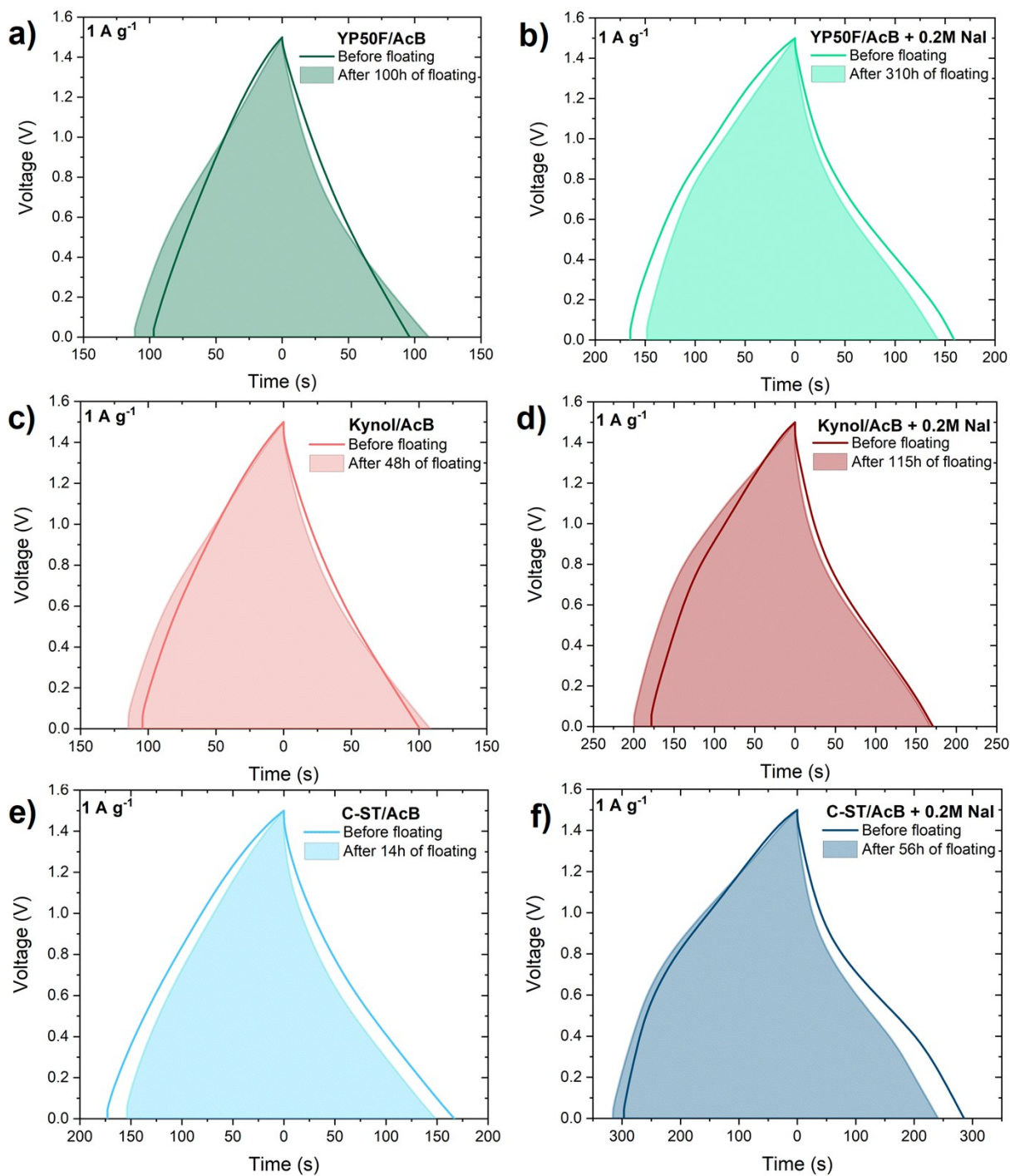
**Fig. S8** Galvanostatic charge/discharge profiles recorded at 1 A g<sup>-1</sup> for ECs based on YP50F, Kynol and C-ST carbons operating in **a)** acetate buffer; **b)** acetate buffer + 0.2 mol L<sup>-1</sup> NaI.



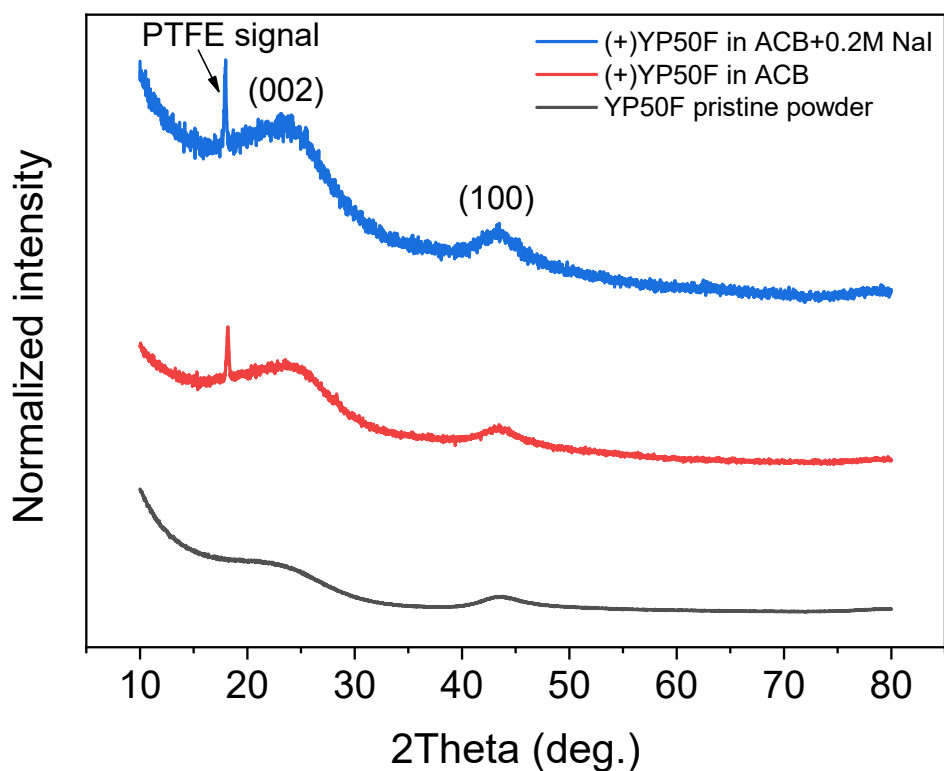
**Fig. S9** Cyclic voltammogram at  $5 mV s^{-1}$  recorded for EC based on Kynol operating in **a)** acetate buffer before and after 48 h floating; **c)** acetate buffer +  $0.2 mol L^{-1}$  NaI before and after 115 h floating. Nyquist plots obtained for Kynol based ECs operating in **b)** acetate buffer before and after 48 h of floating and **d)** acetate buffer +  $0.2 mol L^{-1}$  NaI before and after 115 h floating.



**Fig. S10** Cyclic voltammogram at  $5 \text{ mV s}^{-1}$  recorded for EC based on C-ST operating in **a)** acetate buffer before and after 48 h floating; **c)** acetate buffer +  $0.2 \text{ mol L}^{-1}$  NaI before and after 115 h floating; Nyquist plots obtained for C-ST based ECs operating in **b)** acetate buffer before and after 48 h of floating and **d)** acetate buffer +  $0.2 \text{ mol L}^{-1}$  NaI before and after 115 h floating.



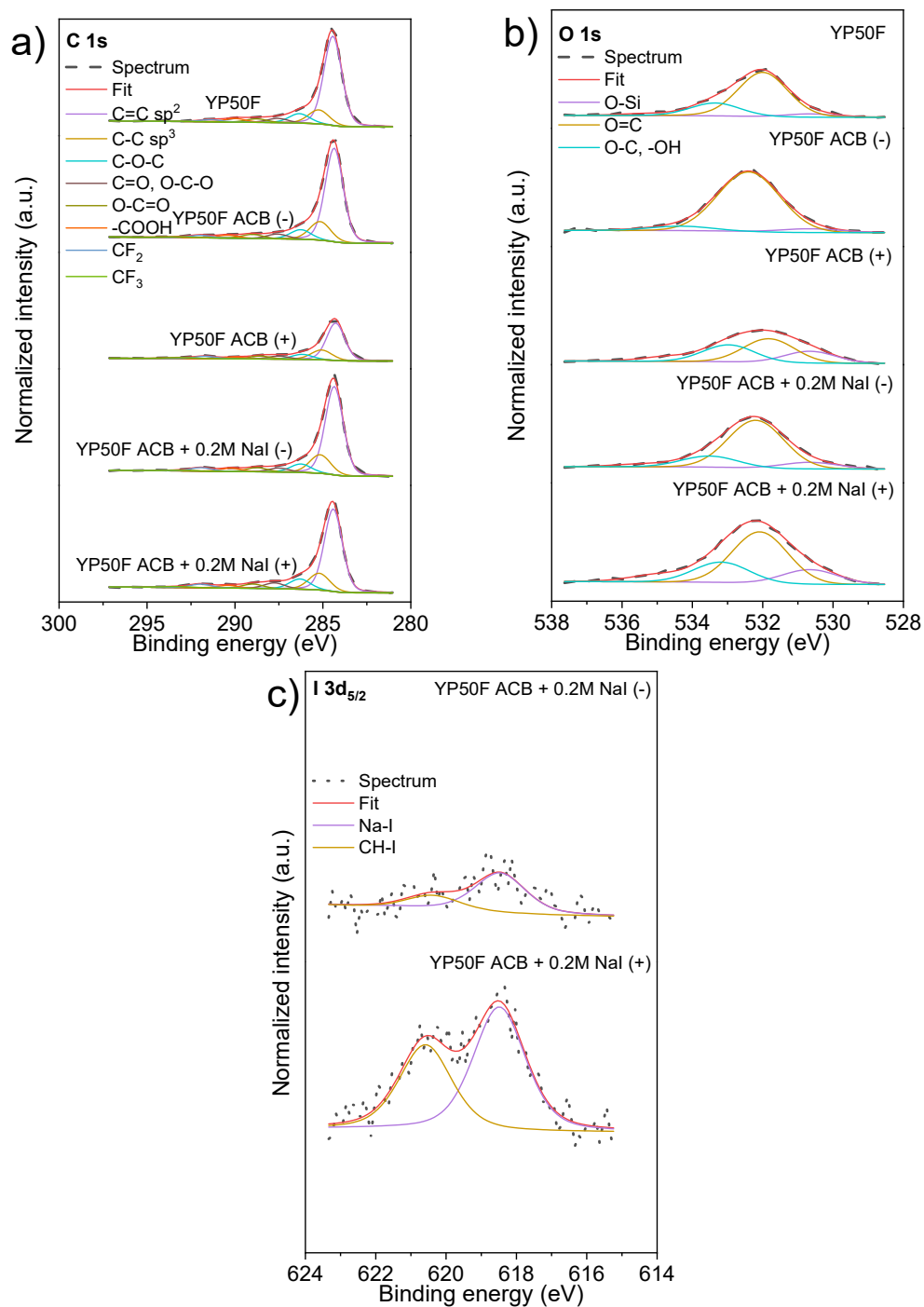
**Fig. S11** Galvanostatic charge/discharge profiles before and after floating for ECs based on **a)** YP50F activated carbon operating in acetate buffer; **b)** YP50F activated carbon operating in acetate buffer + 0.2 mol L<sup>-1</sup> NaI; **c)** Kynol carbon cloth operating in acetate buffer; **d)** Kynol carbon cloth operating in acetate buffer + 0.2 mol L<sup>-1</sup> NaI; **e)** C-ST activated carbon operating in acetate buffer; **f)** C-ST activated carbon operating in acetate buffer + 0.2 mol L<sup>-1</sup> NaI.



**Fig. S12** X-ray diffraction patterns (XRD) of pristine YP50F powder and positive electrodes after floating tests conducted in AcB and AcB + 0.2 mol L<sup>-1</sup> NaI.

**Table S4.** Surface composition (atomic %) determined by fitting XPS spectra for electrodes after aging in different electrolytes.

Binding energy [eV]	C						O		I	
	284.4	285.0	286.3	287.6	288.9	290.3	532.2	533.8	618.4	620.6
Compound / Oxidation State	C=C sp <sup>2</sup>	C-C sp <sup>3</sup>	C-O-C, C-OH	C=O, O-C-O	O-C=O	-COOH	O-Si, O=C, C-O-C	C-O, -OH	NaI	CH-I
Pristine YP50F	56.9	10.4	7.0	3.2	2.6	2.7	7.7	2.2	0.0	0.0
AcB (-)	51.6	11.8	6.1	3.1	2.6	1.9	10.6	0.8	0.0	0.0
AcB (+)	41.4	12.2	6.3	3.8	3.7	2.2	6.8	4.9	0.0	0.0
AcB+0.2M NaI (-)	51.2	12.5	5.7	2.7	2.7	2.2	8.0	1.9	0.05	0.02
AcB+0.2M NaI (+)	47.0	10.8	6.6	3.8	2.8	1.7	8.3	3.3	0.16	0.11



**Fig. S13.** X-ray photoelectron spectroscopy of a) C 1s region, b) O 1s region, and c) I 3d<sub>5/2</sub> region for pristine, positive and negative YP50F electrodes after floating in different electrolytes with and without addition of iodides.