

Supporting Information

Enhanced Toughness of Hydroxyapatite-poly(ethylene terephthalate) Composites by Immersion in Water

Yui Okuda, Ken Hirota, Tadashi Mizutani,* and Yusuke Numamoto

^a Department of Applied Chemistry, Graduate School of Science and Engineering, Doshisha University, Kyotanabe, Kyoto 610-0321 Japan

^b Goo Chemical Corporation, Limited., 58 Ijiri, Iseda-cho, Uji, Kyoto 611-0043 Japan

*E-mail address of corresponding author

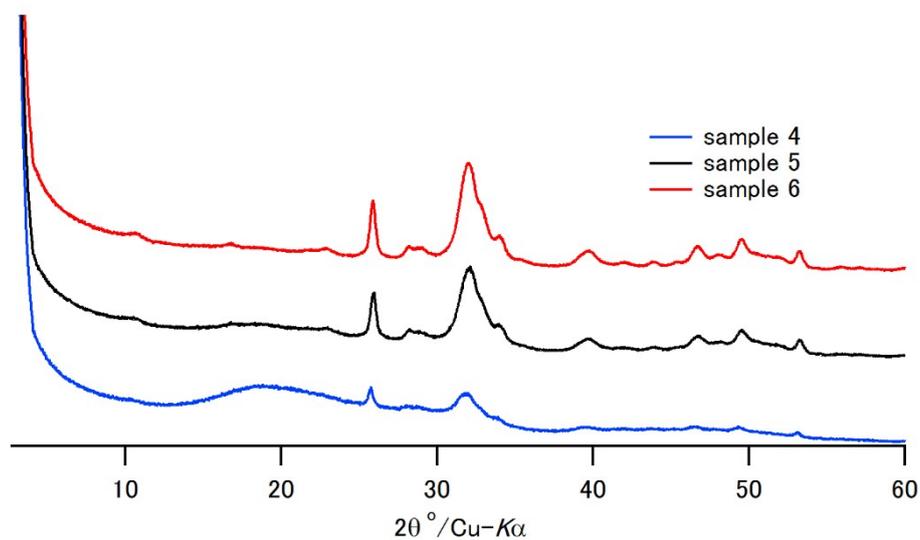
tmizutan@mail.doshisha.ac.jp

Number of pages: 15

Number of figures: 8

Number of tables 3.

1. Characterization of the composite powder



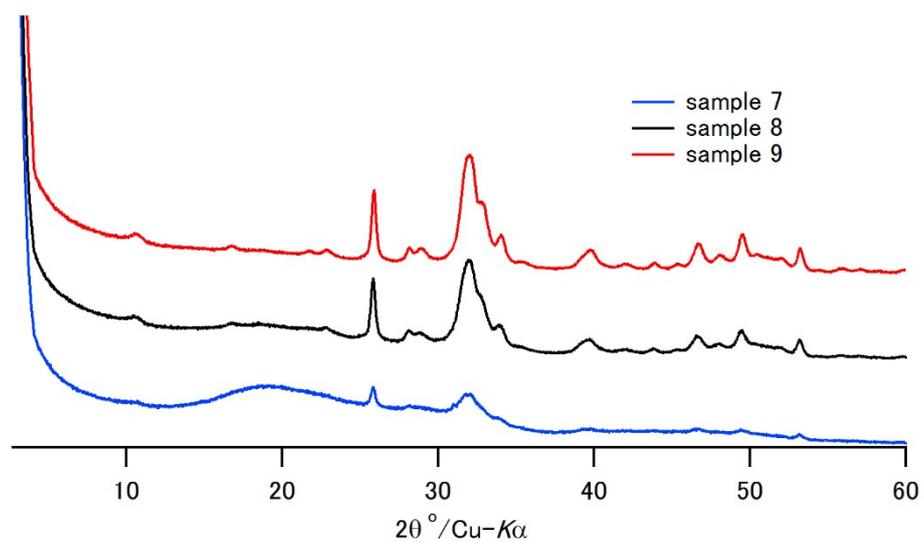
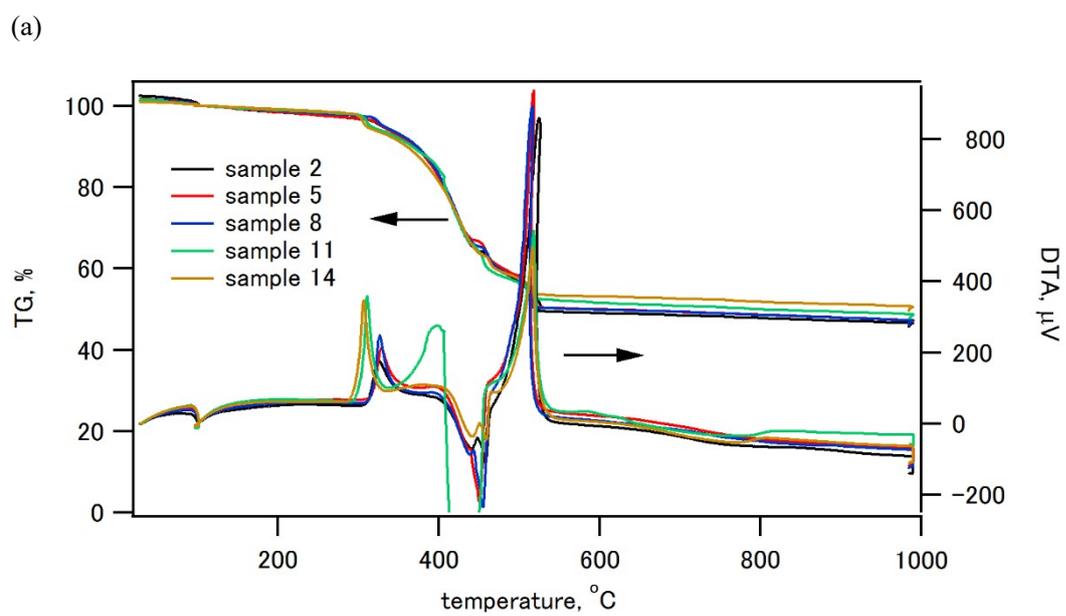


Figure S1. XRD patterns of samples 4-9.



(b)

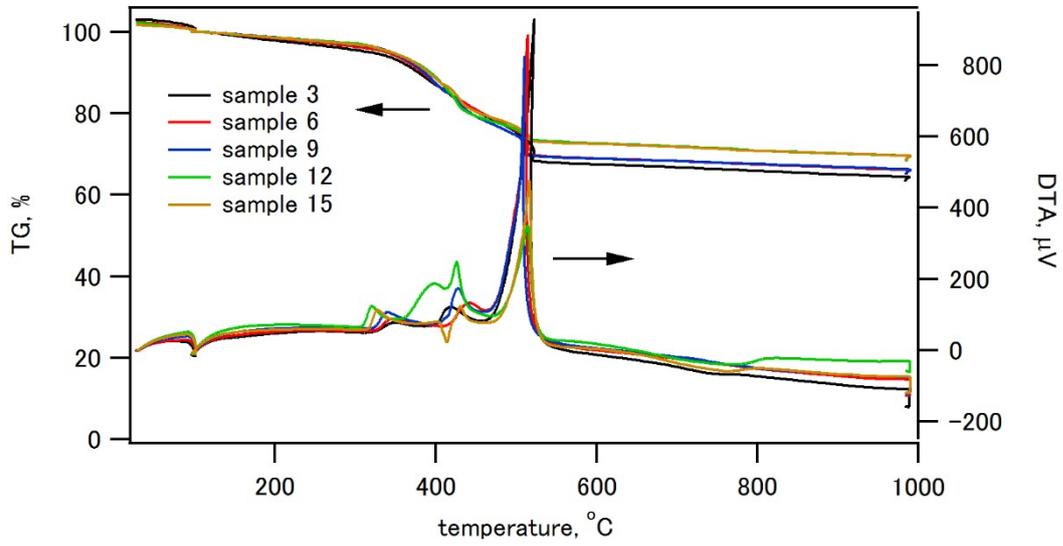


Figure S2. TG/DTA traces of the composites containing 50 wt% (a) and 70 wt% (b) of HAP.

2. Stress-strain curves of the composite compacts.

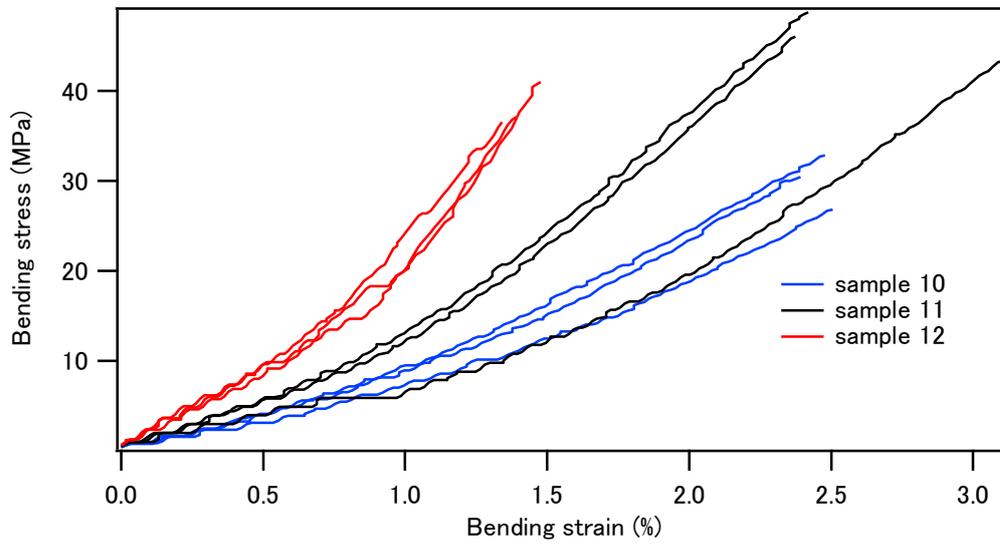
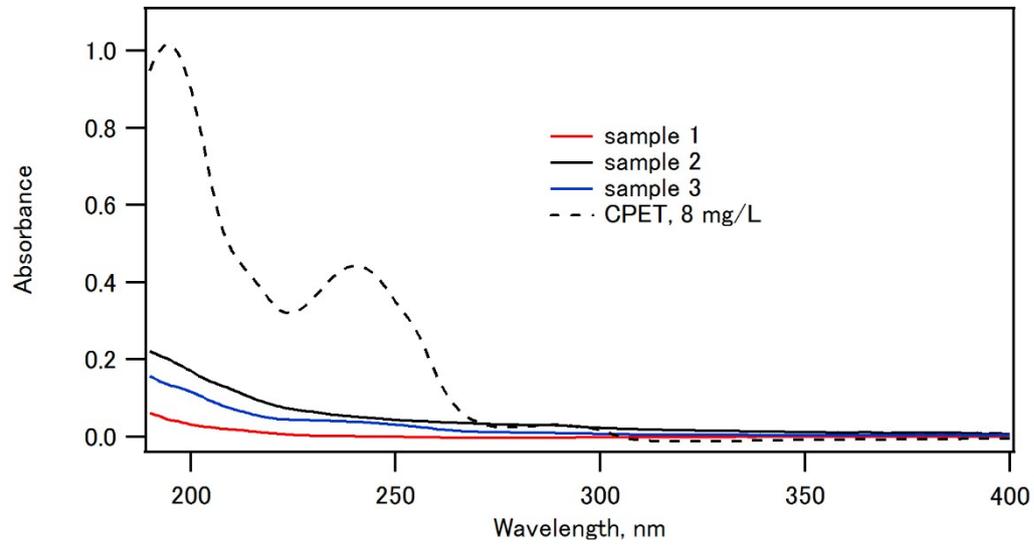


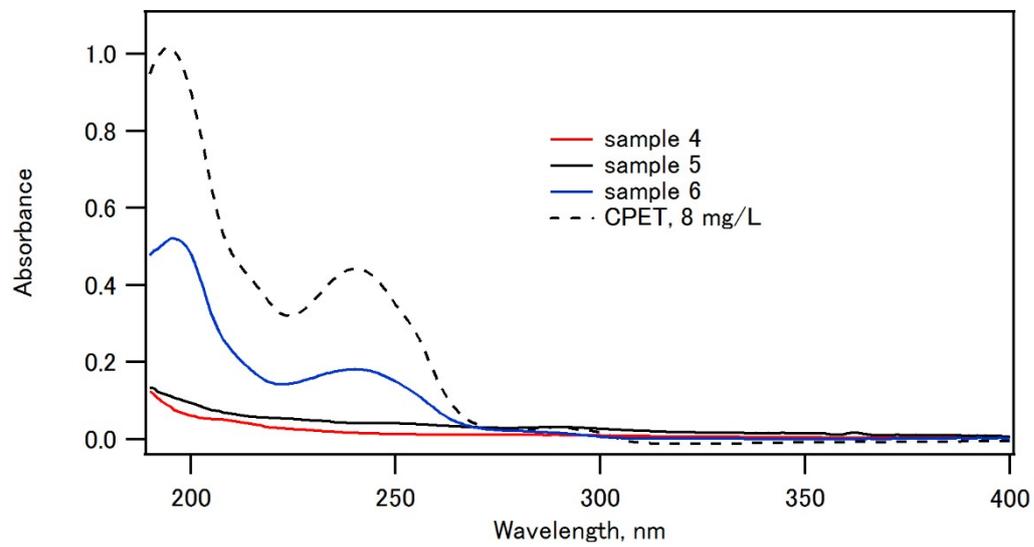
Figure S3. Representative stress-strain curves of samples 10-12 ($N = 3$).

3. Water resistant test: UV-visible spectra of the aqueous phase after immersion of the composite compacts.

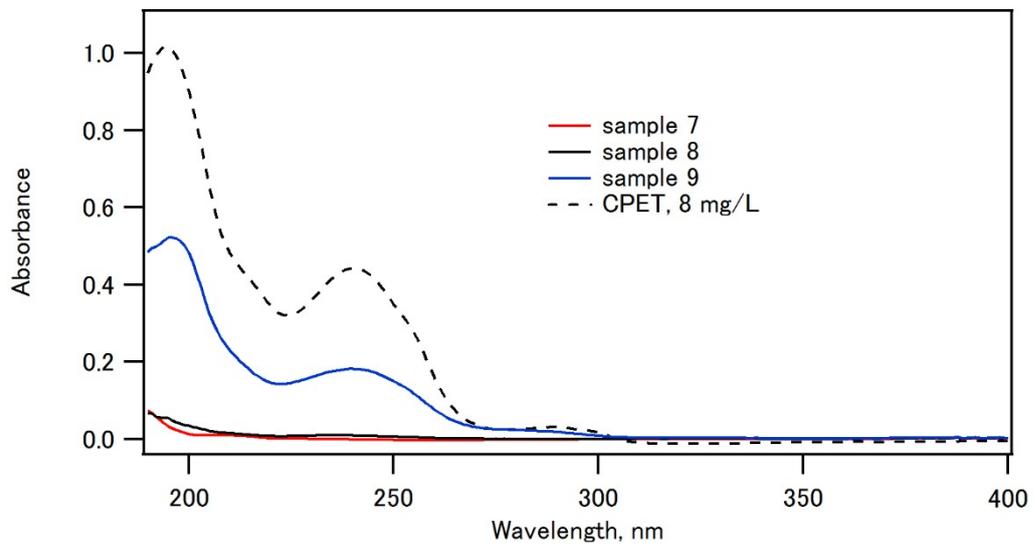
(a)



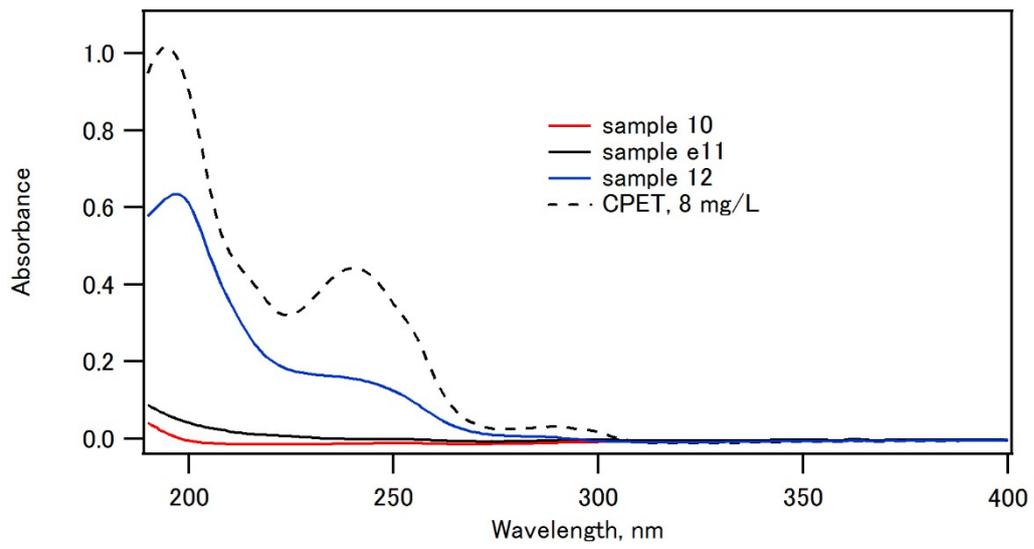
(b)



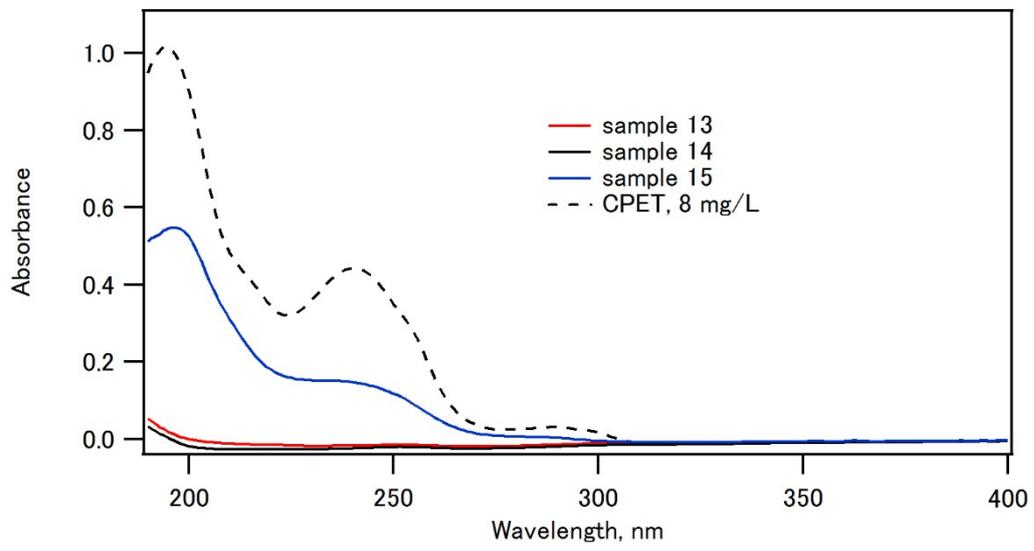
(c)



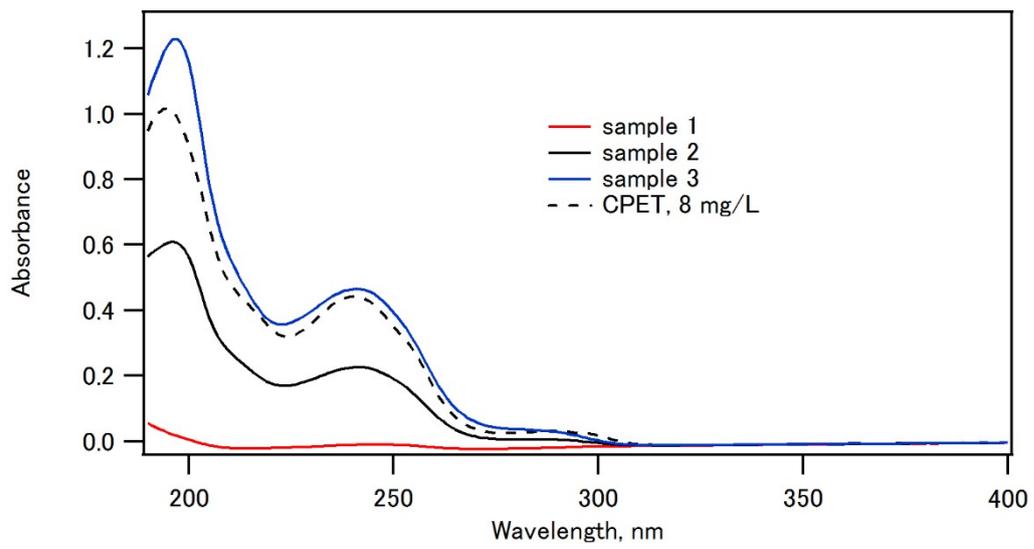
(d)



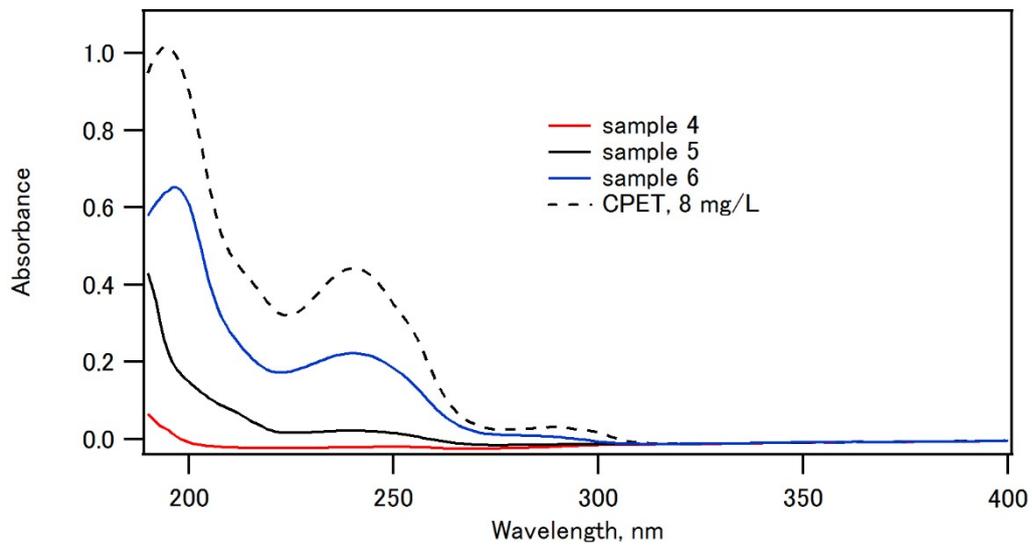
(e)



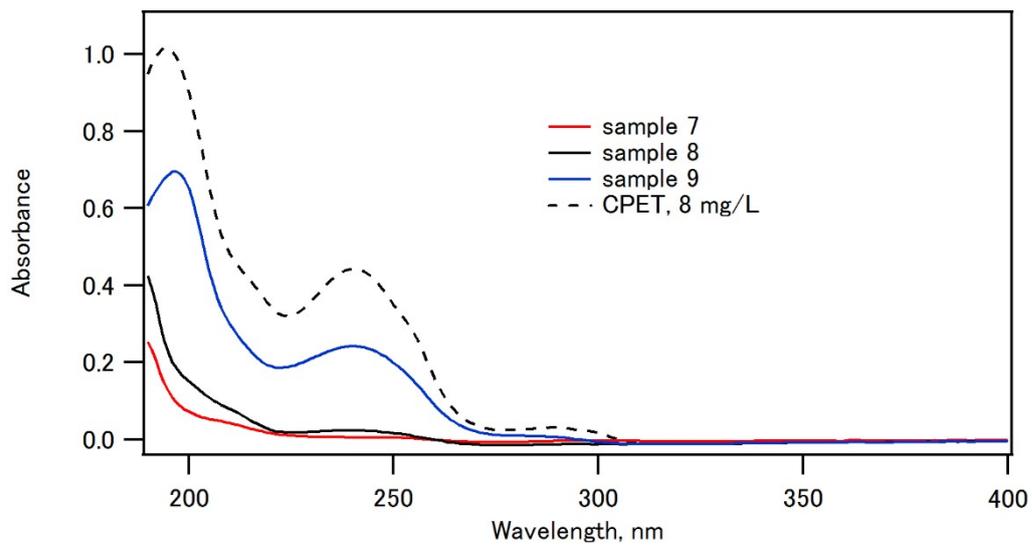
(f)



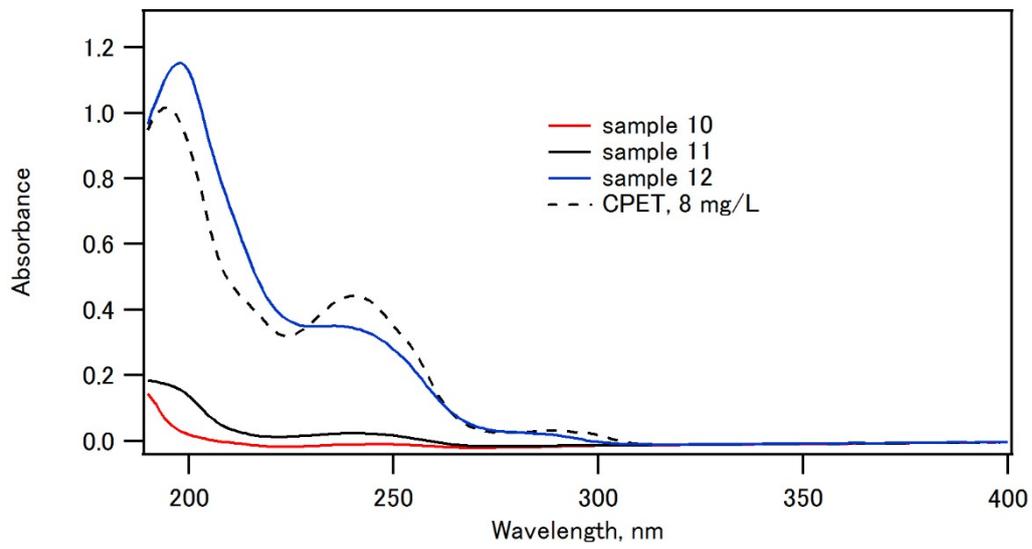
(g)



(h)



(i)



(j)

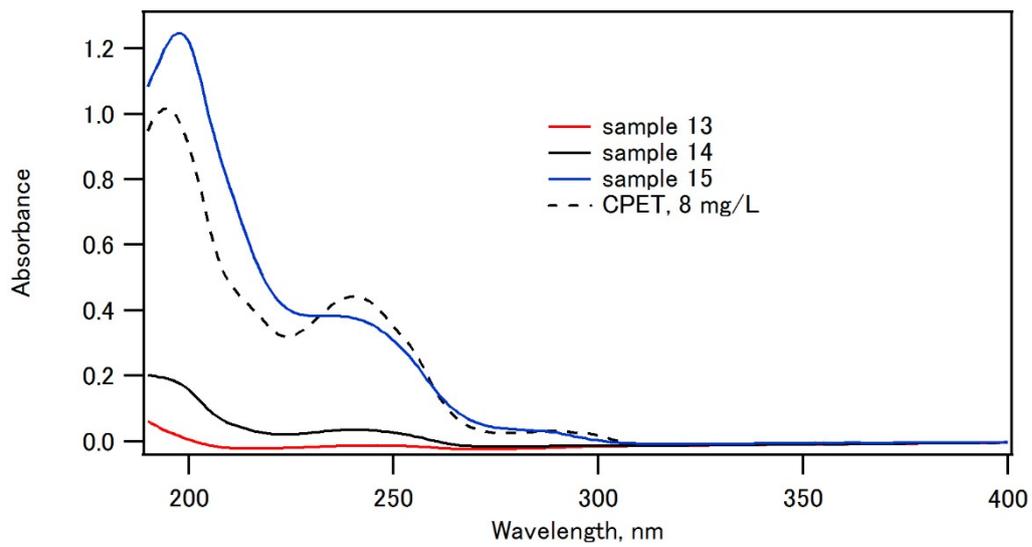


Figure S4. UV-visible spectra of the aqueous phase after immersion of the composite compacts. (a)--(e): immersion at room temperature for 24 h, (f)-(j): immersion at 37 °C for 1 week.

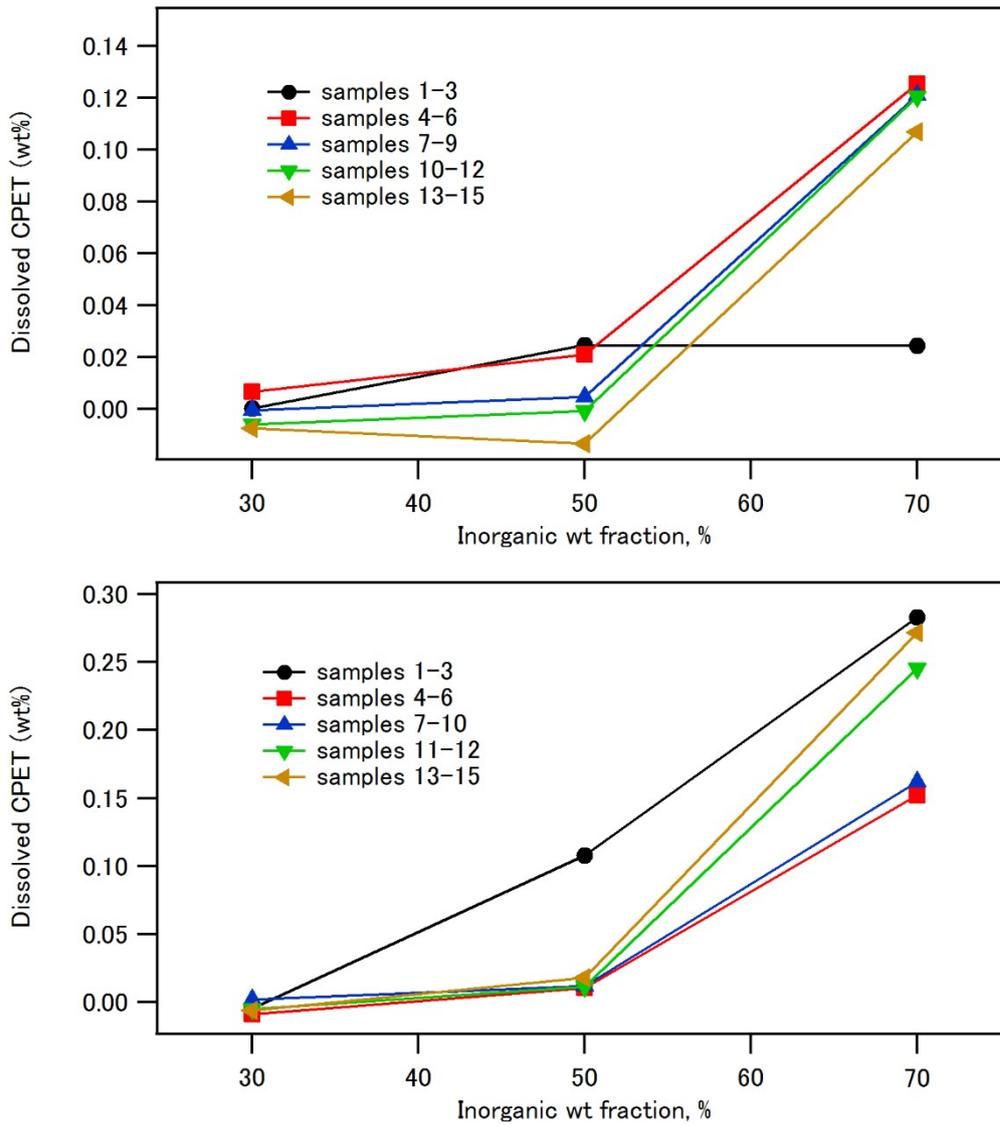
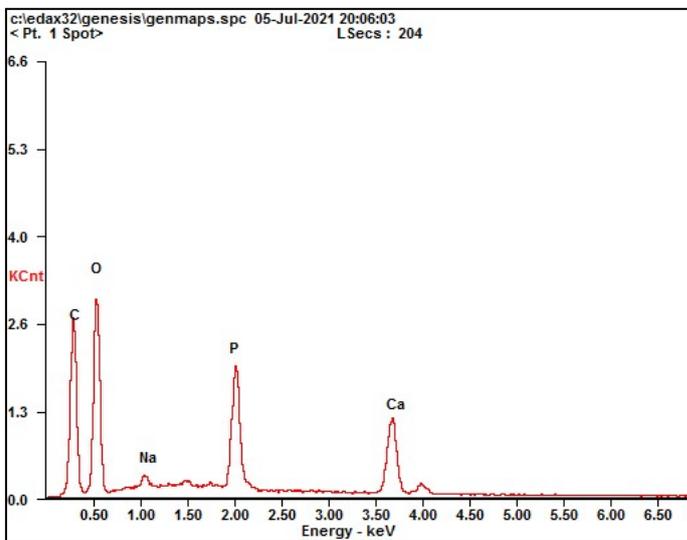


Figure S5. The weight % of CPET dissolved in water after immersion in water at room temperature for 24 h (a), and at 37 °C for 1 week, determined by UV-visible spectra of the water phase.

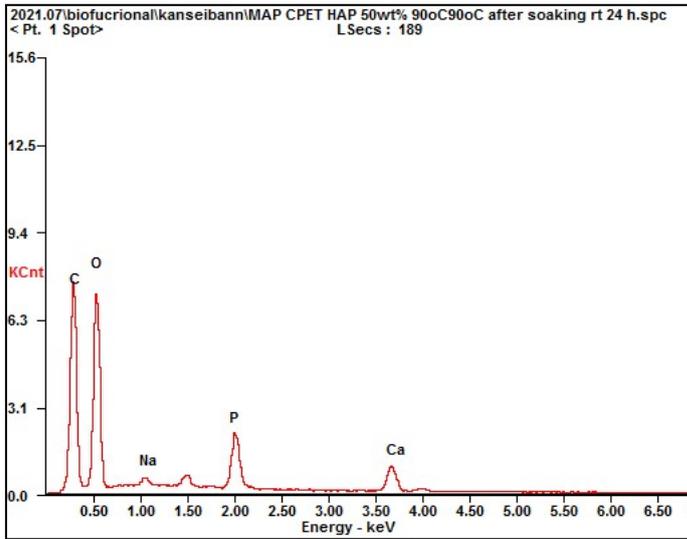
Table S1. Concentrations of CPET (mg/L) in the aqueous phase after immersion of the compacts in 20 mL water at room temperature for 24 h or at 37 °C for 1 week.

Sample	Room temperature, 24 h	37 °C, 1 week
1	0.01	0
2	0.9	4.1
3	0.7	8.4
4	0.3	0
5	0.8	0.4
6	3.3	4.0
7	0	0.1
8	0.2	0.4
9	3.3	4.4
10	0	0
11	0	0.4
12	2.8	6.2
13	0	0
14	0	0.6
15	2.7	6.8



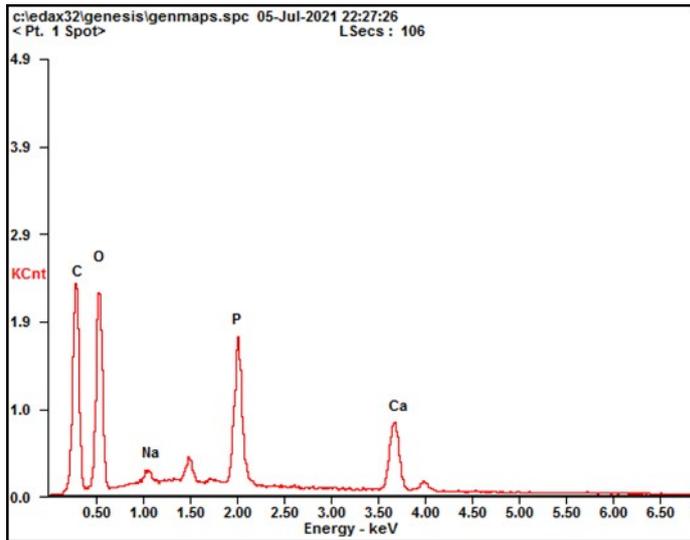
<i>Element</i>	<i>Wt%</i>	<i>At%</i>
<i>CK</i>	23.82	37.41
<i>OK</i>	34.16	40.27
<i>NaK</i>	01.21	00.99
<i>PK</i>	15.28	09.31
<i>CaK</i>	25.53	12.02
<i>Matrix</i>	Correction	ZAF

Figure S6. Energy dispersion spectra of the fracture surface of pristine sample 14.



<i>Element</i>	<i>Wt%</i>	<i>At%</i>
<i>CK</i>	35.49	47.25
<i>OK</i>	42.58	42.56
<i>NaK</i>	01.03	00.72
<i>PK</i>	09.75	05.03
<i>CaK</i>	11.15	04.45
<i>Matrix</i>	Correction	ZAF

Figure S7. Energy dispersion spectra of the fracture surface of sample 14 after immersed in water at room temperature for 24 h.



<i>Element</i>	<i>Wt%</i>	<i>At%</i>
<i>CK</i>	27.50	42.10
<i>OK</i>	31.74	36.48
<i>NaK</i>	01.21	00.97
<i>PK</i>	17.09	10.15
<i>CaK</i>	22.45	10.30
<i>Matrix</i>	Correction	ZAF

Figure S8. Energy dispersion spectra of the fracture surface of sample 14 after immersed in water at 37 °C for 1 week.

4. Mechanical properties of compacts after immersion in water.

Table S2. Mechanical properties and densities of the compacts after immersion in water at room temperature for 24 h.

Sample	density (g/cm ³)	Bending strain (%)	Bending strength (MPa)	Elastic modulus 1 (GPa)	Elastic modulus 2 (GPa)	Fracture energy (MJ/m ³)	<i>N</i> ^b
1	1.331 ± 0.004	1.3 ± 0.1	15 ± 2	0.70 ± 0.03	1.6 ± 0.1	0.083 ± 0.008	3
2	1.54 ± 0.01	1.7 ± 0.1	14 ± 2	0.7 ± 0.3	1.0 ± 0.1	0.12 ± 0.03	3
3	1.82 ± 0.03	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	3
4	1.382 ± 0.008	1.8 ± 0.1	21.0 ± 0.2	1.0 ± 0.1	1.8 ± 0.2	0.19 ± 0.02	3
5	1.53 ± 0.03	1.6 ± 0.3	17 ± 1	0.8 ± 0.1	1.5 ± 0.1	0.12 ± 0.02	3
6	1.608 ± 0.009	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	3
7	1.376 ± 0.006	1.8 ± 0.3	22 ± 3	0.80 ± 0.04	1.7 ± 0.2	0.17 ± 0.05	3
8	1.597 ± 0.004	2.2 ± 0.1	25 ± 2	0.9 ± 0.2	1.8 ± 0.1	0.23 ± 0.03	3
9	1.73 ± 0.03	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	3
10	1.39 ± 0.01	1.4 ± 0.2	21 ± 1	1.1 ± 0.2	1.8 ± 0.3	0.130 ± 0.009	3
11	1.62 ± 0.02	2.0 ± 0.1	30 ± 1	1.1 ± 0.2	2.5 ± 0.1	0.26 ± 0.01	3
12	1.87 ± 0.02	1.0 ± 0.3	13.1 ± 0.5	1.2 ± 0.4	1.5 ± 0.2	0.06 ± 0.01	3
13	1.369 ± 0.007	2.4 ± 0.2	26 ± 2	0.46 ± 0.06	2.1 ± 0.2	0.232 ± 0.009	3
14	1.64 ± 0.02	2.8 ± 0.3	38 ± 2	0.55 ± 0.03	1.9 ± 0.1	0.45 ± 0.05	3
15	1.81 ± 0.01	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	3

a The compacts were collapsed after immersion in water, and a three point bending test cannot be carried out. b The number of determinations.

Table S3. Mechanical properties and densities of the compacts after immersion in water at 37 °C for 1 week.

Sample	density (g/cm ³)	Bending strain (%)	Bending strength (MPa)	Elastic modulus 1 (GPa)	Elastic modulus 2 (GPa)	Fracture energy (MJ/m ³)	<i>N</i> ^b
1	1.38 ± 0.02	1.5 ± 0.1	13 ± 2	0.85 ± 0.08	1.20 ± 0.07	0.08 ± 0.02	3
2	1.56 ± 0.02	0.6 ± 0.2	5 ± 1	1.0 ± 0.1	1.0 ± 0.2	0.017 ± 0.009	3
3	1.82 ± 0.03	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	3
4	1.39 ± 0.02	1.6 ± 0.4	15 ± 3	0.8 ± 0.1	1.14 ± 0.09	0.11 ± 0.04	3
5	1.542 ± 0.001	0.13 ± 0.08	2 ± 1	0.6 ± 0.3	0.6 ± 0.3	0.002 ± 0.002	3
6	1.79 ± 0.03	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	3
7	1.378 ± 0.004	2.1 ± 0.2	15.4 ± 0.9	0.50 ± 0.05	1.03 ± 0.08	0.14 ± 0.01	3
8	1.587 ± 0.008	0.45 ± 0.05	4.6 ± 0.7	0.9 ± 0.1	0.9 ± 0.1	0.011 ± 0.003	3
9	1.830 ± 0.008	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	3
10	1.39 ± 0.01	1.54 ± 0.05	19 ± 1	0.68 ± 0.02	1.7 ± 0.2	0.123 ± 0.008	5
11	1.61 ± 0.02	1.2 ± 0.1	18 ± 1	1.1 ± 0.1	1.5 ± 0.1	0.10 ± 0.01	5
12	1.74 ± 0.03	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	5
13	1.371 ± 0.009	2.2 ± 0.2	22 ± 1	0.68 ± 0.06	1.50 ± 0.02	0.21 ± 0.03	3
14	1.60 ± 0.03	7.43 ± 0.05	22.0 ± 0.9	0.53 ± 0.04	0.18 ± 0.01	0.96 ± 0.01	3
15	1.79 ± 0.04	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	<i>a</i>	3

a The compacts were collapsed after immersion in water, and a three point bending test cannot be carried out. b The number of determinations.