## Supplementary Information for

## Treatment of Electrochemical Plating Wastewater by Heterogeneous Photocatalysis: The Simultaneous Removal of 6:2 Fluorotelomer Sulfonate and Hexavalent Chromium

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Photocatalyst	BET Surface area (m <sup>2</sup> /g)			
Ga <sub>2</sub> O <sub>3</sub>	19.5			
In <sub>2</sub> O <sub>3</sub>	7.1			
TiO <sub>2</sub>	51.9			

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 Table S1. BET surface area of the as received photocatalysts.

**Table S2.** Formula and molecular weight of investigated PFASs.

Chemical	Abbreviation	Formula	Molecular weight (g/mol)	
6:2 Fluorotelomer sulfonate	6:2 FtS	$C_6F_{13}CH_2CH_2SO_3H$	428.16	
6:2 Fluorotelomer carboxylate	6:2 FtCA	C <sub>6</sub> F <sub>13</sub> CH <sub>2</sub> COOH	378.09	
Perfluoroheptanoate	PFHpA	C <sub>6</sub> F <sub>13</sub> COOH	364.06	
Perfluorohexanoate	PFHxA	C <sub>5</sub> F <sub>11</sub> COOH	314.05	
Perfluoropentanoate	PFPeA	C <sub>5</sub> F <sub>9</sub> COOH	264.05	
Perfluorobutanoate	PFBA	C <sub>4</sub> F <sub>7</sub> COOH	214.04	

Instrument	Shimadzu LCMS-8030 Triple Quadrupole Mass Spectrometer						
Ionization	Negative electrospray						
Precolumn	Guard Column Thermo Scientific <sup>™</sup> Acclaim 120 Å C18, 4.6 x 10 mm, 5 µm						
Column	Agilent Infinity Lab Poroshell 120 Å EC-C18, 3.0 x 50 mm, 2.7 µm						
Column oven temperature	30°C						
Injection	30 µL						
Mobile phases	A: 5 mM ammonium acetate in LCMS grade water B: LCMS grade Methanol						
Flow rate	0.5 mL/min						
	Time (min)		Eluent A Conc. (%)	Eluent B Conc. (%)			
Gradient	0		60		40		
	0.5		60	40			
	4.5		20	80			
prome	10		20		80		
	10.5		60	40			
	13.5		Stop				
	Analytes	Ion transitions	Internal standards	Ion transitions	Calibration range (µg/L)		
Monitored	6:2 FtS	427>407	[ <sup>13</sup> C <sub>2</sub> ]M6:2FtS	429 > 81	0.05 - 20		
ion	6:2 FtCA	377 > 293	[ <sup>13</sup> C <sub>2</sub> ]M6:2FtCA	379>294	0.05 - 20		
transitions	PFHpA	363 > 319	[ <sup>13</sup> C <sub>4</sub> ]MPFHpA	367 > 322	0.05 - 20		
	PFHxA	313>629	[ <sup>13</sup> C <sub>5</sub> ]MPFHxA	315>270	0.05 - 20		
	PFPeA	263 > 219	[ <sup>13</sup> C <sub>5</sub> ]MPFPeA	268 > 223	0.25 - 20		
	PFBA	213>169	[ <sup>13</sup> C <sub>4</sub> ]MPFBA	217>172	0.25 - 20		

 Table S3. Quantitative analytical method.



Figure S1. SEM images and EDS analyses of the as received photocatalysts.



**Figure S2.** (a) photograph of the 4W UV-C lamp-equipped photoreactor box and (b) intensity of UV-C light (6 units) versus wavelength.



Figure S3. The removals of 6:2 FtS (a) and Cr(VI) (b) by Ga<sub>2</sub>O<sub>3</sub>, In<sub>2</sub>O<sub>3</sub>, and TiO<sub>2</sub> photocatalysts in the absence of light. [catalyst] = 0.5 g/L, [6:2 FtS]<sub>0</sub> = 100  $\mu$ g/L, [Cr(VI)]<sub>0</sub> = 1 mg/L, pH = 3.0 ± 1.



Figure S4. The removals of 6:2 FtS and Cr(VI) by UV-C/Ga<sub>2</sub>O<sub>3</sub> system.  $[Ga_2O_3] = 0.5$  g/L, [6:2 FtS]<sub>0</sub> = 2 mg/L,  $[Cr(VI)]_0 = 20$  mg/L, pH =  $3.0 \pm 1$ .





Figure S6. Effect of HCOOH doses on the degradation of 6:2 FtS by the UV-C/Ga<sub>2</sub>O<sub>3</sub> in the presence of Cr(VI). ([Ga<sub>2</sub>O<sub>3</sub>]<sub>0</sub> = 0.5 g/L (surface area-based dose: 9.76 m<sup>2</sup>/L), [6:2 FtS]<sub>0</sub> = 100  $\mu$ g/L, [Cr(VI)] = 1 mg/L, [MeOH]  $_0$  = 0.3 mM, pH<sub>i</sub> = 3, Pre-sorption time = 60 min).



Figure S7. Effects of bromate or N<sub>2</sub> sparging on the degradation of 6:2 FtS by the UV-C/Ga<sub>2</sub>O<sub>3</sub> system. ([Ga<sub>2</sub>O<sub>3</sub>]<sub>0</sub>=0.5 g/L (surface area-based dose: 9.76 m<sup>2</sup>/L), [6:2 FtS]<sub>0</sub>=100  $\mu$ g/L, [MeOH]  $_0 = 0.3 \text{ mM}$ , [BrO<sub>3</sub>-]<sub>0</sub> = 10 mM, pH<sub>i</sub> = 3, Pre-sorption time = 60 min)



Figure S8. Photographs of Ga<sub>2</sub>O<sub>3</sub>/Cr, TiO<sub>2</sub>/Cr, and In<sub>2</sub>O<sub>3</sub>/Cr suspensions after 8 h UVC irradiation, respectively.  $([Ga_2O_3]_0 = [TiO_2]_0 = [In_2O_3]_0 = 0.5 \text{ g/L}$  (surface area-based doses:  $[Ga_2O_3] = 9.76 \text{ m}^2/\text{L}$ ,  $[In_2O_3] = 3.57 \text{ m}^2/\text{L}$ ,  $[TiO_2] = 25.94 \text{ m}^2/\text{L}$ ),  $[6:2 \text{ FtS}]_0 = 100 \text{ µg/L}$ , [Cr(VI)] = 1 mg/L,  $[MeOH]_0 = 0.3 \text{ mM}$ ,  $pH_i = 3$ , Pre-sorption time = 60 min).

( 2 μm	a) Ga <sub>2</sub> O <sub>3</sub> 2 μm Ο K	Ga K	2 µm	(b) In <sub>2</sub> O <sub>3</sub> 2μm Ο Κ	In L Cr K	2 µm	(c) TiO <sub>2</sub> 2 µm О К	Ti K Cr K
<u>2 μm</u>	2 µm		2 µm	2 μm		2 µm	2 µm	
Element	Weight %	Atomic %	Element	Weight %	Atomic %	Element	Weight %	Atomic %
ОК	27.95	62.80	ОК	22.50	67.42	ОК	19.00	41.35
Ga K	71.83	37.04	ln L	77.06	32.17	In L	77.28	56.15
Cr K	0.22	0.15	Cr K	0.44	0.40	Cr K	3.72	2.50

Figure S9. SEM/EDS images of the spent catalysts.