The Efficacy of Chlorine Photolysis as an Advanced Oxidation Process for Drinking Water Treatment: Supplemental Information

Christina K. Remucal^{a,b*} and Devon Manley^b

^aDepartment of Civil and Environmental Engineering

University of Wisconsin – Madison

Madison, Wisconsin

^bEnvironmental Chemistry and Technology Program

University of Wisconsin – Madison

Madison, Wisconsin

Submitted to: Environmental Science: Water Research & Technology, Jan. 28, 2016.

Contents (9 pages): Tables S1-S4

^{*} Corresponding author e-mail: remucal@wisc.edu; telephone: 608-262-1820; fax: 608-262-0454.

Tables S1-S4 summarize the experimental conditions for studies that quantified the formation of disinfection by-products (DBPs) by chlorine in the presence and absence of light. The type and fluence of the light source, the experimental duration (i.e., time between chlorine addition and DBP measurement), the free available chlorine (FAC) concentration, pH, and dissolved organic carbon (DOC) concentration are indicated for each data point. The difference between the concentration of DBP measured in the irradiated solutions and in dark control experiments is presented graphically in Figure 3 in the manuscript. Note that full-scale studies that prechlorinated water before any treatment are not included due to the elevated background DBP concentrations detected using that approach.

Table S1: Summary of experimental conditions for studies quantifying the formation of trihalomethanes (THMs) produced by chlorine in the presence and absence of light. The studies quantified either chloroform alone (denoted CHCl₃) or total trihalomethanes (denoted TTHM; the sum of bromodichloromethane, bromoform, chloroform, and dibromochloromethane).

DBPs	Light	Fluence	Duration	Ouration pH	[FAC] (mg/L	[DOC]	[DBP] in Irradiated	[DBP] in Dark	Reference
Quantified	Source	(mJ/cm²)	(min)	рп	as Cl ₂)	(mg/L)	Experiment (µg/L)	Experiment (µg/L)	Reference
				8.1	2		4.2	9.5	1
	Hg	N/A	60	6.5		4	2.9	3.1	
				7		5	122	87	
	LP UV			7		5	78	55	
CHCl ₃	LPOV			6.8	7	1.8	35	24	3
		60	4,320	7		5	80	47	
	MP UV		4,320	7		5	92	87	
				7		5	77	55	
				6.8		1.8	40	24	
				7		5	86	47	
	LP UV	3,900	120	7.5	10	2.6	14	13	
	UV-A	7,000					24	13	
					2		7	4	
TTHM				6.5	6		8	6	
	MP UV	1,820	1		10	3.5	5	4	4
					2		8	7	
				7.5	6	ŗ	8	10	
					10		6	8	

Table S1: Continued.

DBPs	Light	Fluence	Duration	ъU	[FAC] (mg/L	[DOC]	[DBP] in Irradiated	[DBP] in Dark	Reference
Quantified	Source	(mJ/cm²)	(min)	рН	as Cl ₂)	(mg/L)	Experiment (µg/L)	Experiment (µg/L)	Reference
					2		10	9	4
	MP UV	1,820	1	8.5	6	3.5	10	11	
					10		7	9	
				6.5	7.5 6.5		100	50	
TTHM			1,440	7.5			100	65	
				8.5			120	85	
	LP UV	/ 800	120		7.2 1.5	3.2	15	31	5
			800 1,440 4,320	7.2			17	40	
							19	50	

Table S2: Summary of experimental conditions for studies quantifying the formation of haloacetic acids (HAAs) produced by chlorine in the presence and absence of light. The studies quantified the sum of dichloroacetic acid (DCAA) and trichloroacetic acid (TCAA; denoted DCAA + TCAA), the five regulated HAAs (denoted HAA5; the sum of bromoacetic acid, dibromoacetic acid, dichloroacetic acid, monochloroacetic acid, and trichloroacetic acid), or nine HAAs (denoted HAA9; the sum of HAA5, bromochloroacetic acid, bromodichloroacetic acid, chlorodibromoacetic acid, and tribromoacetic acid).

DBPs	Light	Fluence	Duration	LJ	[FAC] (mg/L	[DOC]	[DBP] in Irradiated	[DBP] in Dark	Deference
Quantified	Source	(mJ/cm²)	(min)	pН	as Cl ₂)	(mg/L)	Experiment (µg/L)	Experiment (µg/L)	Reference
				7		5	57	76	
	LP UV			7		5	118	108	
	LFOV			7		5	80	68	
DCAA +		60	5	6.8	7	1.8	48	37	2
TCAA		00	ິ	7	,	5	66	76	
	MP UV			7		5	98	108	
				6.8		1.8	50	37	
				7		5	95	68	
HAA5	LP UV	3,900	. 120	7.5	10		37	17	3
IIAAS	UV-A	7,000				2.6	31	17	
	LP UV	3,900					42	27	
	UV-A	7,000					47	27	
				6.5	2		8	3	4
HAA9		1,820			6		7	3	
IIAAS	MP UV		1		10	3.5	4	2	
	IVIF UV				2	3.5	13	3	
				7.5	6		9	3	
					10		5	3	

Table S2: Continued.

DBPs	Light	Fluence	Duration	рН	[FAC] (mg/L	[DOC]	[DBP] in Irradiated	[DBP] in Dark	Reference													
Quantified	Source	(mJ/cm ²)	(min)	рп	as Cl₂)	(mg/L)	Experiment (µg/L)	Experiment (µg/L)	Reference													
					2		15	3														
			1	8.5	6		8	3														
HAA9	MP UV	1,820			10		5	2	4													
ПААЭ			1,440	6.5	6.5	-	140	70	4													
				7.5			100	75														
				8.5		Ī	70	60														
	LP UV	UV 800														30				10	22	
HAA5			120 1,440	7.2	1.5	3.2	12	25	5													
HAAS				1.2		5.2	18	34														
			4,	4,320	-			22	39													

Table S3: Summary of experimental conditions for studies quantifying the formation of haloacetonitriles (HANs) produced by chlorine in the presence and absence of light. Both studies quantified the sum of bromochloroacetonitrile, dibromoacetonitrile, dichloroacetonitrile, and trichloroacetonitrile.

DBPs	Light	Fluence	Duration (min) pH	[FAC] (mg/L	[DOC]	[DBP] in Irradiated	[DBP] in Dark	Deference							
Quantified	Source	(mJ/cm²)		рп	as Cl ₂)	(mg/L)	Experiment (µg/L)	Experiment (µg/L)	Reference						
					2		2.5	0							
				6.5	6		2.5	0							
					10		1.5	0							
					2		4	0.5	4						
			1	7.5	6	3.5	3	0.5							
	MP UV	1,820			10		1.5	0							
				8.5	2		5	0.5							
HAN					6		3	0.5							
LIAN					10		1.5	0.5							
			1,440	6.5			30	9							
				7.5	6.5		16	8							
				8.5			4	3							
	LP UV	800	30				17	8	- 5						
			120	7.2	1.5	3.2	22	11							
			1,440		1.5	3.2	26	14							
										4,320				40	20

Table S4: Summary of experimental conditions for studies quantifying the formation of total organic halides (TOX) or adsorbable organic halides (AOX).

DBPs Quantified	Light Source	Fluence (mJ/cm ²)	Duration (min)	рН	[FAC] (mg/L as Cl₂)	[DOC] (mg/L)	[DBP] in Irradiated Experiment (µg/L)	[DBP] in Dark Experiment (µg/L)	Reference
	LP UV			6.5	0.28	5	479.25	443.75	6
		60	10	8.5			390.5	319.5	
	MP UV		10	6.5	0.20	3	461.5	443.75	
	IVII OV			8.5			284	319.5	
	MP UV	220	10	6.5	0.28	5	301.75	337.25	7
TOX	IVII OV	220	10	0.0	0.20	3	461.5	443.75	,
	LP UV		4,320		1.7	1	111	113	
	MP UV	300		8.6	2.2		147	113	8
	LP UV	3,900	400	7.5	10	2.6	65	97	3
	UV-A	7,000	120				97	97	
		1,820		6.5	2		60	35	
					6		40	40	
					10		25	35	
				7.5	2		70	35	4
			0.67		6		50	35	
AOX	MP UV				10	3.5	30	35	
AOX	IVII OV	1,020			2	0.0	90	42	
				8.5	6		70	35	
					10		40	35	
				6.5	6.5		300	200	
			1,440	7.5	6.5		225	200	
				8.5	6.5		210	200	

References

- 1. L. H. Nowell and J. Hoigné, Water Res., 1992, 26, 599-605.
- 2. W. Liu, L.-M. Cheung, X. Yang and C. Shang, *Water Res.*, 2006, **40**, 2033-2043.
- 3. A. N. Pisarenko, B. D. Stanford, S. A. Snyder, S. B. Rivera and A. K. Boal, *J. Adv. Oxid. Technol.*, 2013, **16**, 137-150.
- D. Wang, J. R. Bolton, S. A. Andrews and R. Hofmann, *Sci. Total Environ.*, 2015, 518-519, 49-57.
- 5. X. Zhang, W. Li, I. Blatchley, Ernest R, X. Wang and P. Ren, *Water Res.*, 2015, **68**, 804-811.
- 6. Q. Zhao, C. Shang and X. Zhang, Water Sci. Tech., 2009, **9**, 627-634.
- 7. Q. Zhao, C. Shang, X. Zhang, G. Ding and X. Yang, *Water Res.*, 2011, **45**, 6545-6554.
- 8. M. J. Plewa, E. D. Wagner, D. H. Metz, R. Kashinkunti, K. J. Jamriska and M. Meyer, *Environ. Sci. Technol.*, 2012, **46**, 7811-7817.