

表1 管理値のまとめ

水温	0~10	10~20	20 ~
紫外線吸光度 (250 nm、50mm)	0.083	0.067	0.056

表2 H28年度ハロ酢酸低減化対策実施状況

	塩素注入点 (前または中間)			粉末活性炭 (mg/L)
	着水井	フロック形成池	沈澱池末端	
4/1~5/15				-
5/16~6/5				-
6/6~7/28				3
7/29~9/4				4
9/5~10/11				4
10/12~				4

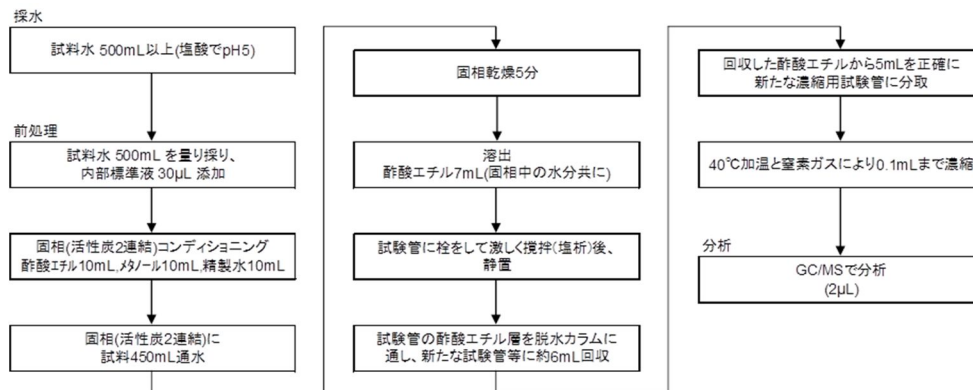


図1 ハロアセトアミドの分析フロー

表3 ハロアセトアミド GC/MS 分析条件

測定装置	GC: Agilent製 7890A MS: Agilent製 5975C	モニターイオン(m/z)	定量	確認
分離カラム	GL Sciences製 InertCap 5MS (30m × 0.25mm、1.00 µm)	1,2-ジブromoロハシ	121	123
注入方法	スプリットレス	クロロアセトアミド*	93	44
カラム温度	35°C (3分) → 10°C / 分 → 190°C → 30°C / 分 → 290°C (10分)	フロモアセトアミド*	137	139
キャリアーガス	He 1.5mL/min	ジクロロアセトアミド*	127	44
注入口温度	250°C	ブromoクロロアセトアミド*	173	44
イオン源温度	230°C	トリクロロアセトアミド*	82	98
		ジブromoアセトアミド*	217	174

表4 対象論文リスト

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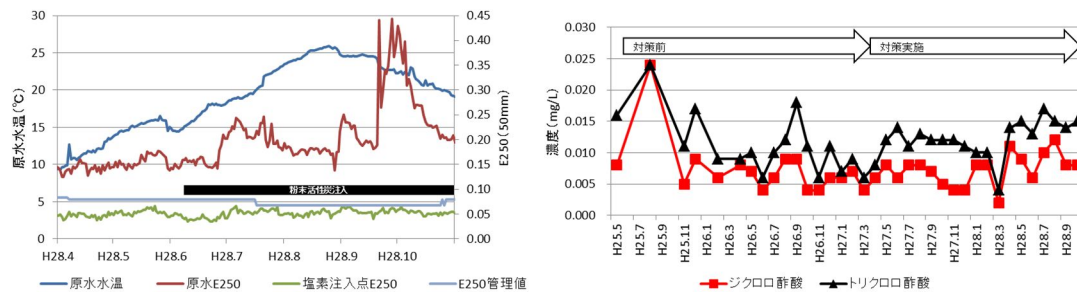


図2 塩素注入点と原水吸光度に基づいたハロ酢酸制御 (左: E250 管理状況, 右: ハロ酢酸月最大値のまとめ)

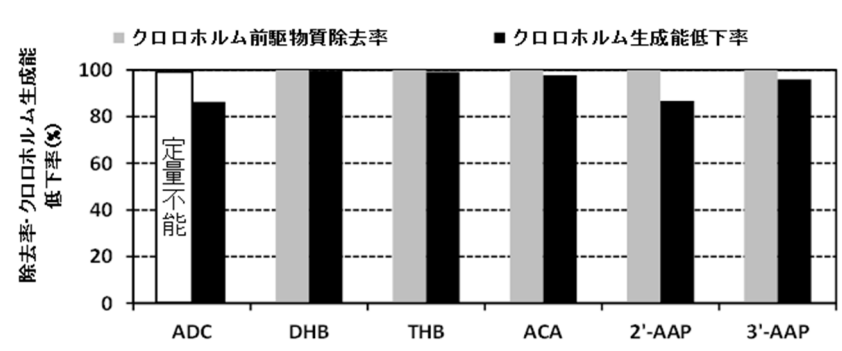


図3 クロロホルム前駆物質の中オゾン処理性調査結果

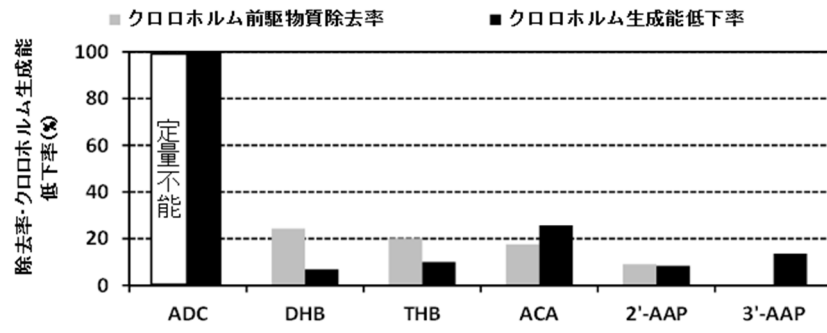


図4 クロロホルム前駆物質の急速砂ろ過処理性調査結果

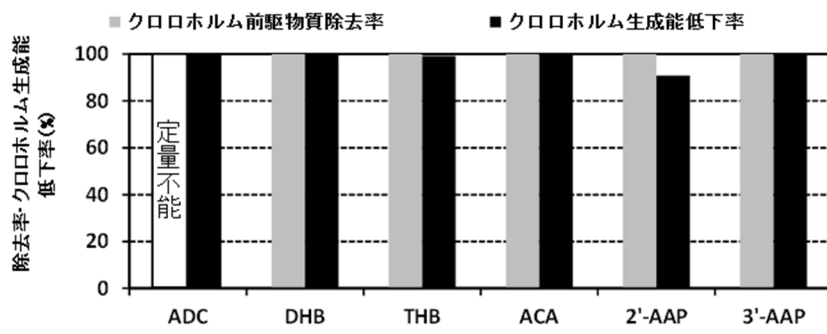


図5 クロロホルム前駆物質のGAC処理性調査結果

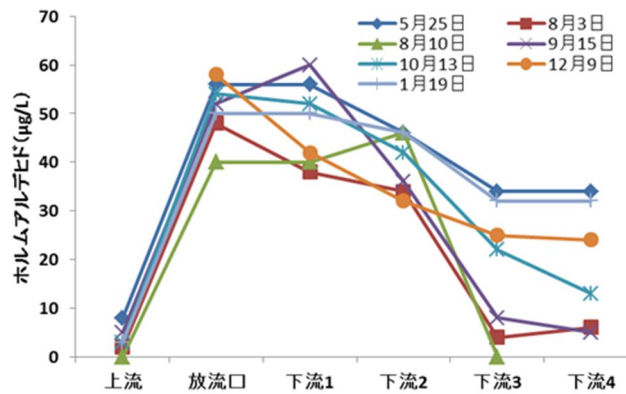


図6 西高瀬川におけるホルムアルデヒドの挙動

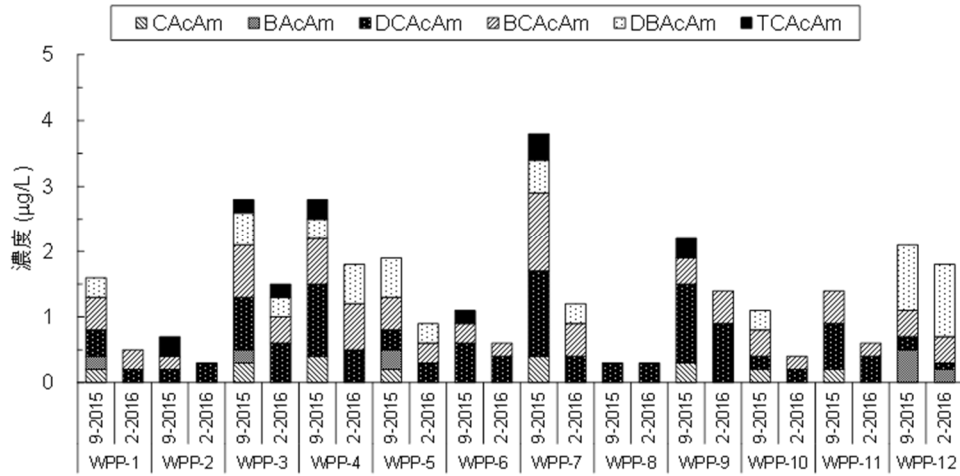


図7 全国 12 浄水場の水道水中の HACams 濃度

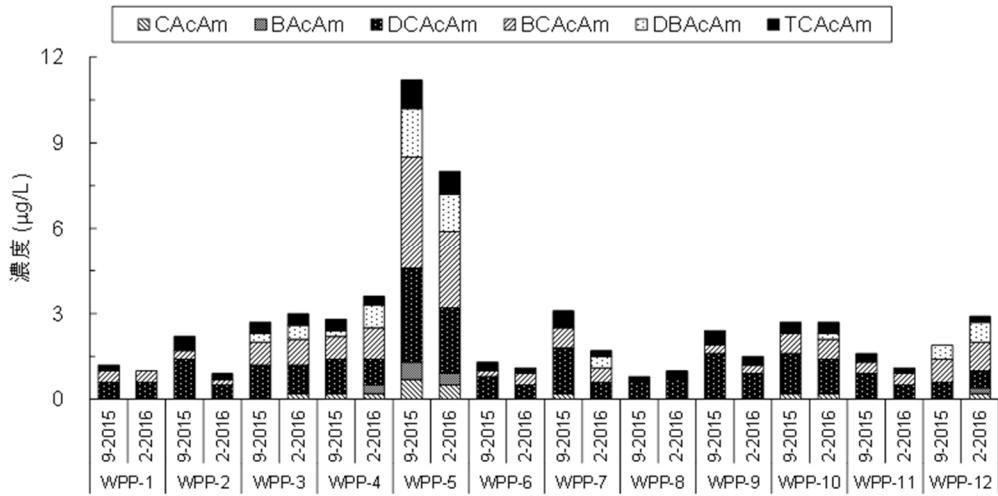


図8 全国 12 浄水場の原水の HACams 生成能

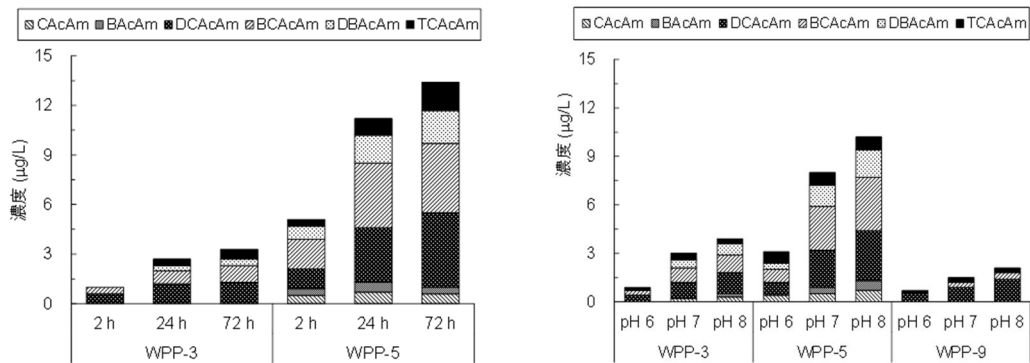


図9 原水の HACams 生成能におよぼす塩素処理時間 (左) と pH(右)の影響

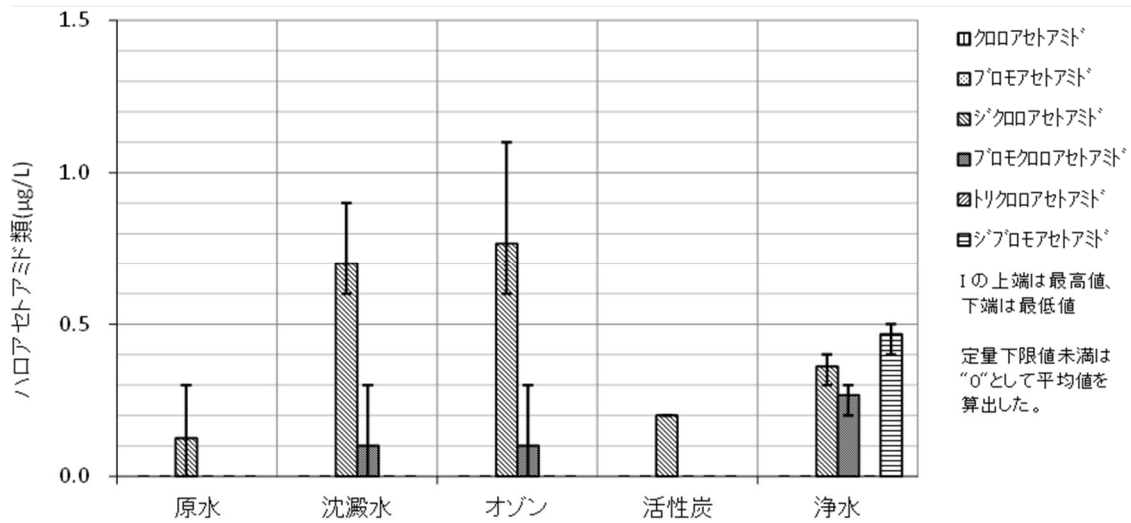


図 10 浄水処理過程における検出状況(平均値)

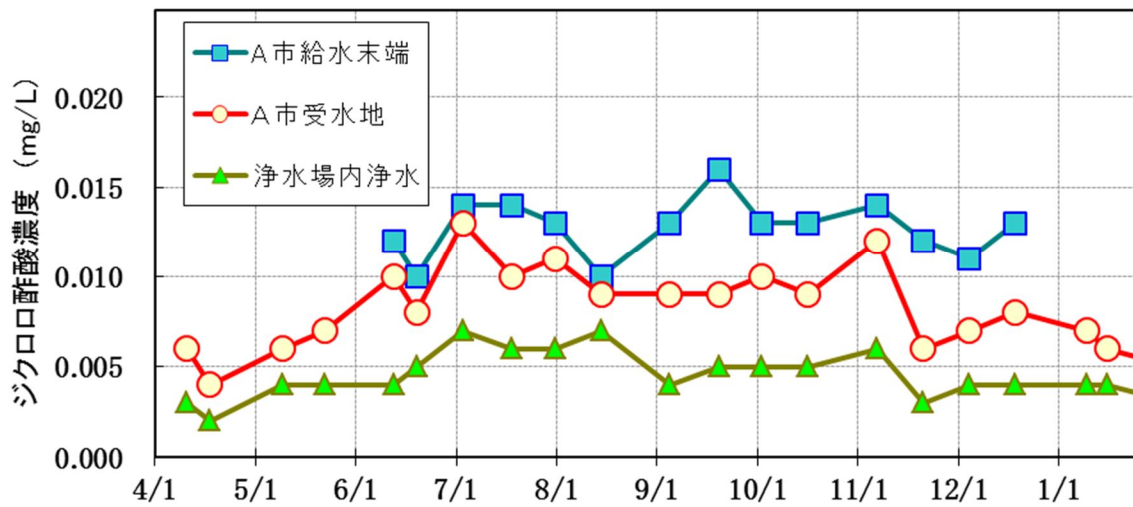


図 11 浄水場内浄水等のジクロロ酢酸濃度の推移

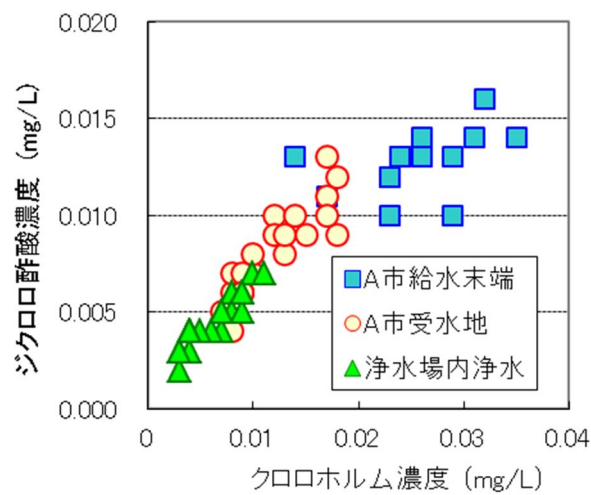


図 12 ジクロロ酢酸とクロロホルムの相関

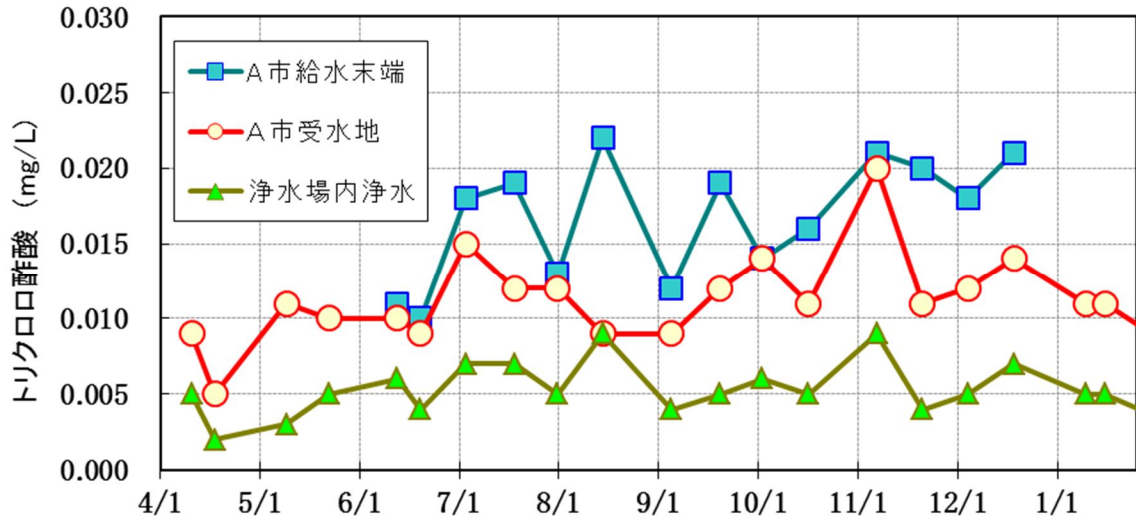


図 13 浄水場内浄水等のトリクロロ酢酸の推移

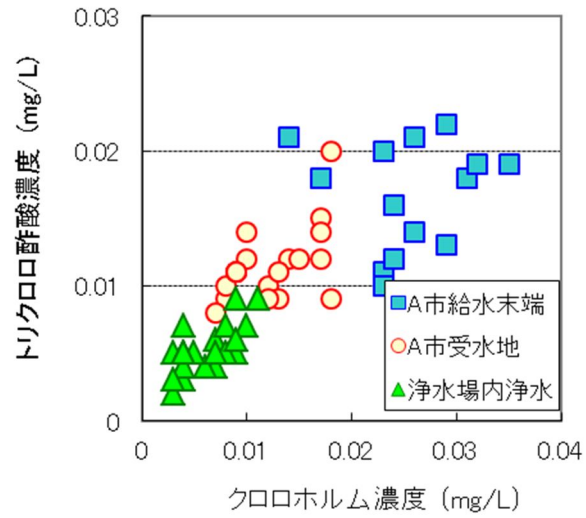


図 14 トリクロロ酢酸とクロロホルムの相関

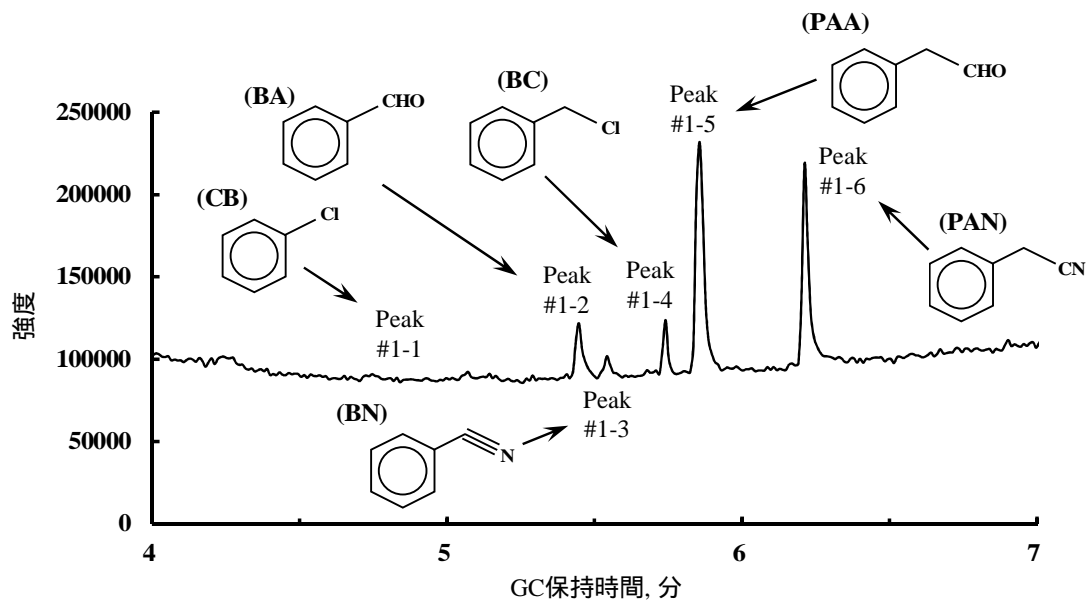


図 15 フェニルアラニン塩素処理溶液のパージ&トラップ - GC/MS トータルイオンクロマトグラム  
 CB, クロロベンゼン; BA, ベンズアルデヒド; BN, ベンゾニトリル; BC, 塩化ベンジル; PAA, フェニルアセト  
 アルデヒド; PAN, フェニルアセトニトリル

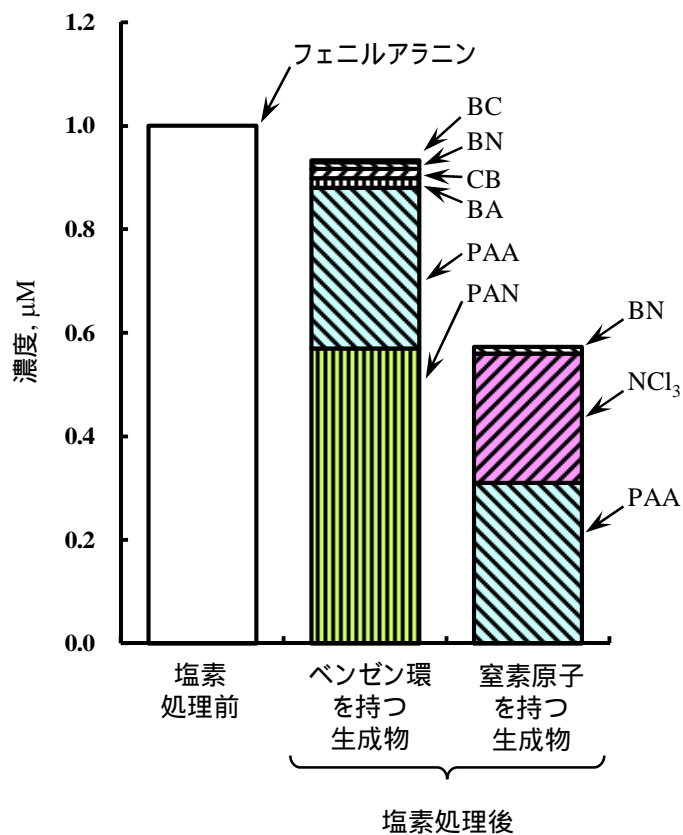


図 16 フェニルアラニン塩素処理溶液中での生成物 PAA, フェニルアセトアルデヒド; PAN, フェニルアセトニトリル;  $NCl_3$ , トリクロロアミン; BA, ベンズアルデヒド; CB, クロロベンゼン; BN, ベンゾニトリル; BC, 塩化ベンジル



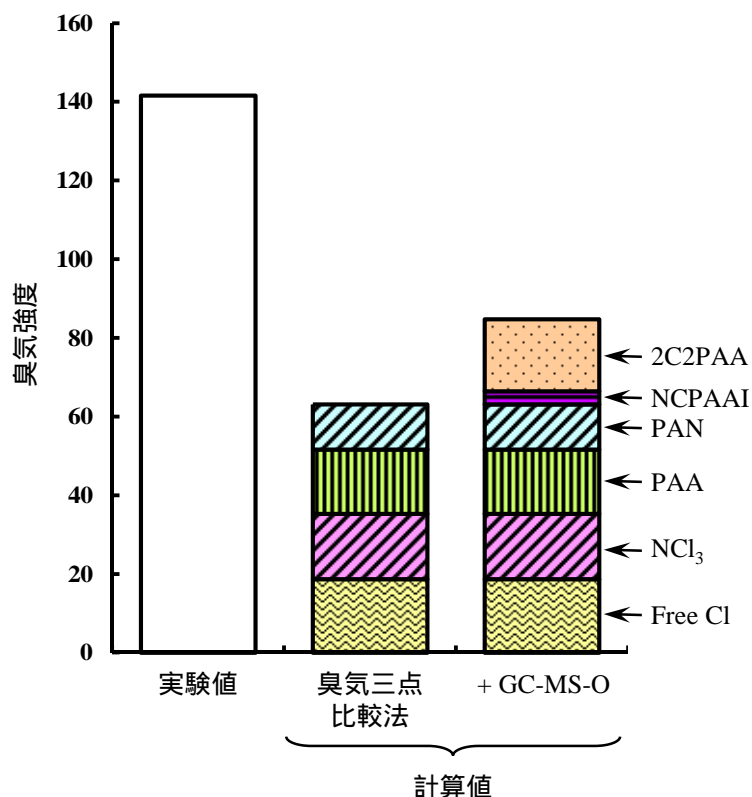


図 17 フェニルアラニン塩素処理溶液の臭気強度の実験値と計算値の比較 2C2PAA, 2-クロロ-2-フェニルアセトアルデヒド; NCPAAI, N-クロロフェニルアセトアルドイミン; PAN, フェニルアセトニトリル; PAA, フェニルアセトアルデヒド; NCl<sub>3</sub>, トリクロラミン; free Cl, 遊離塩素

表 5 三点比較法により評価した標準品が入手可能な分解生成物の臭気閾値とそれらのフェニルアラニン塩素処理溶液中での濃度

生成物など	検出濃度	本研究での臭気閾値		既存文献における臭気閾値
	μM	μM	μg/L	μg/L
フェニルアセトアルデヒド (PAA)	0.57	0.035	4.2	4 <sup>1-3)</sup> , 1200 <sup>4)</sup>
フェニルアセトニトリル (PAN)	0.31	0.027	3.2	30 <sup>4)</sup>
トリクロラミン	0.25	0.015	3.1 (μg-Cl <sub>2</sub> /L)	20 (μg-Cl <sub>2</sub> /L) <sup>5)</sup>
ベンズアルデヒド (BA)	0.019	0.79	84	350 <sup>3)</sup> , 2000 <sup>6)</sup> , 4600 <sup>1)</sup>
クロロベンゼン (CB)	0.018	0.31	35	50 <sup>7)</sup>
ベンゾニトリル (BN)	0.013	0.26	27	
塩化ベンジル (BC)	0.0034	0.025	3.1	12 <sup>7)</sup>
遊離塩素	14	0.75	53 (μg-Cl <sub>2</sub> /L)	50 (μg-Cl <sub>2</sub> /L) <sup>8)</sup>

<sup>1)</sup> Noguero-Pato et al. (2013); <sup>2)</sup> Eduardo et al. (2010); <sup>3)</sup> Buttery et al. (1988); <sup>4)</sup> Freuze et al. (2005); <sup>5)</sup> Kransner and Barrett (1984); <sup>6)</sup> Noguero-Pato et al. (2011); <sup>7)</sup> Amore and Hautala (1983); <sup>8)</sup> Piriou et al. (2004).

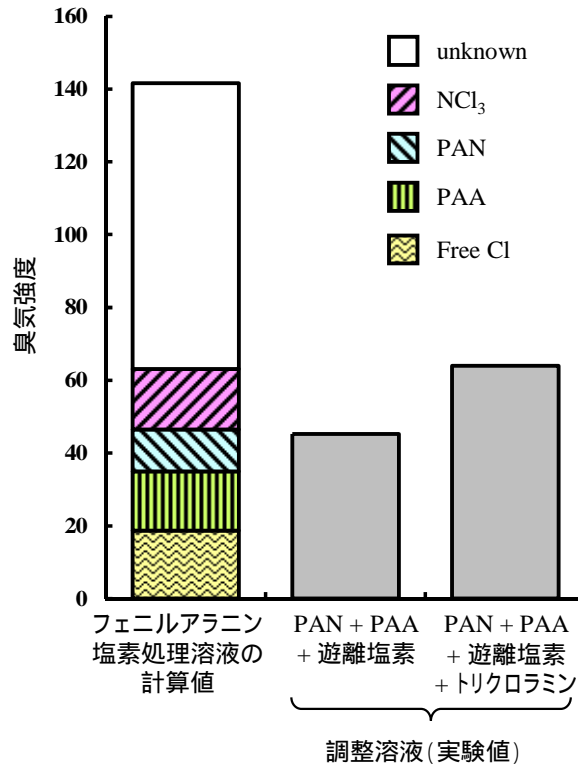


図 18 フェニルアラニン塩素処理溶液と同濃度の生成物（標準品）を含む調整溶液の臭気強度（臭気三点比較法による実験値）と計算値の比較 PAN, フェニルアセトニトリル; PAA, フェニルアセトアルデヒド; NCl<sub>3</sub>, トリクロラミン; free Cl, 遊離塩素

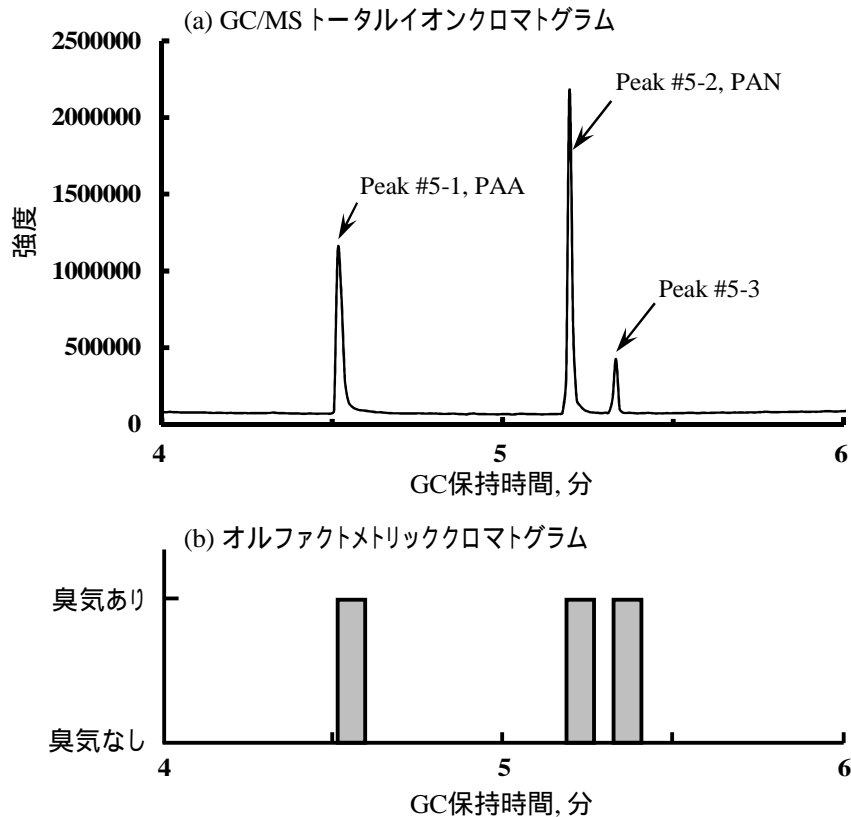


図 19 (a)フェニルアラニン塩素処理溶液をクロロホルムにより溶媒抽出したサンプルの GC/MS トータルイオンクロマトグラム (b)高濃度 NCPAAI 溶液をクロロホルムにより溶媒抽出したサンプルのオルファクトメトリッククロマトグラム PAA, フェニルアセトアルデヒド; PAN, フェニルアセトニトリル

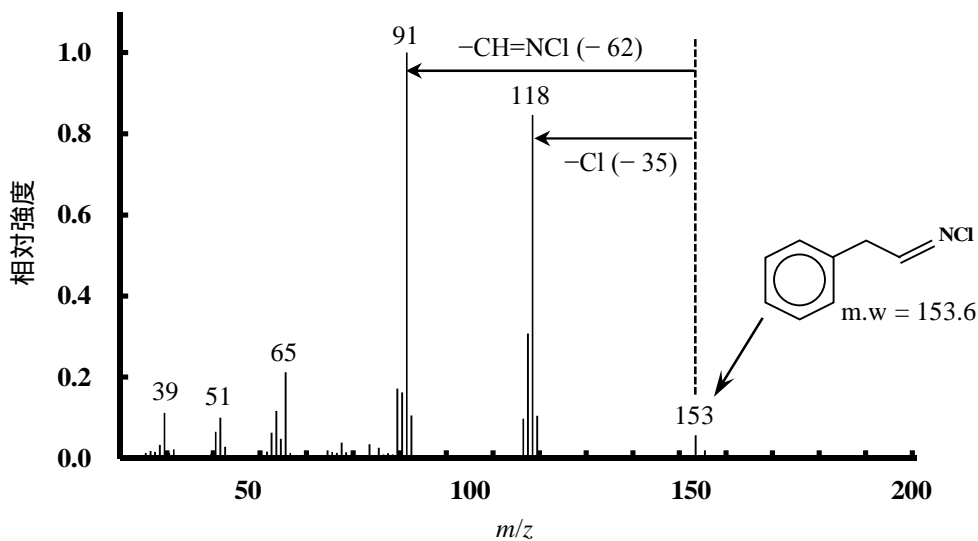


図 20 Peak#5-3 のマススペクトル

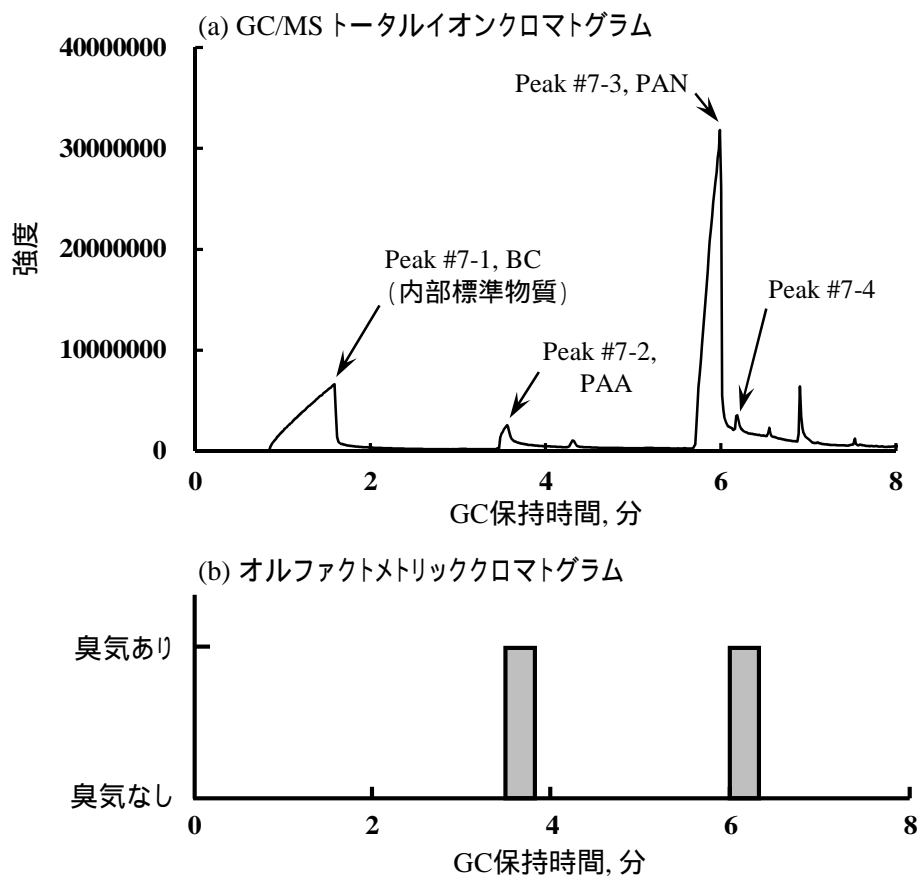


図 21 TP 探索用溶液の (a)GC/MS トータルイオンクロマトグラムと (b)オルファクトメトリッククロマトグラム BC, 塩化ベンジル(内部標準物質); PAA, フェニルアセトアルデヒド; PAN, フェニルアセトニトリル

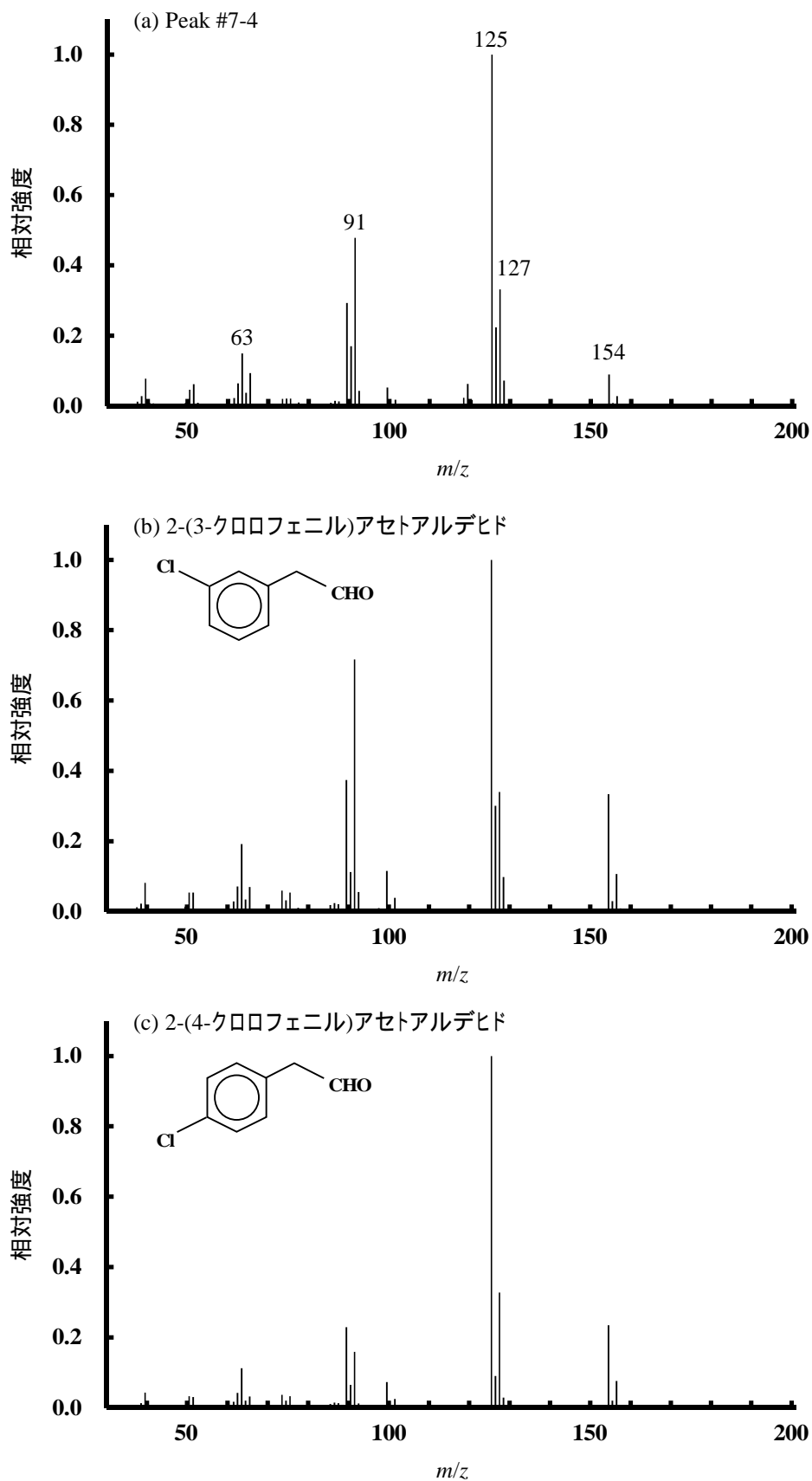
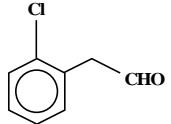
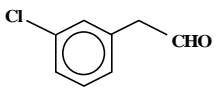
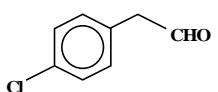
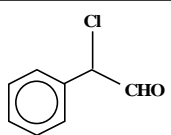


図 22 (a)Peak#7-4(保持時間 6.2 分), (b)2-(3-クロロフェニル)アセトアルデヒド標準品(保持時間 6.7 分), (c) 2-(4-クロロフェニル)アセトアルデヒド標準品(保持時間 6.8 分)のマススペクトルの比較

表6 Peak#7-4として検出された生成物の構造の候補 沸点はChemSpider web (<http://www.chemspider.com/>)上でのACD/Labs PredictorとEPISuiteによる予測値

	沸点, °C	
	ACD/Labs	EPISuite
 <p>(a) 2-(2-クロロフェニル)アセトアルデヒド</p>	235.8 ± 15	232.30
 <p>(b) 2-(3-クロロフェニル)アセトアルデヒド</p>	235.8 ± 15	232.30
 <p>(c) 2-(4-クロロフェニル)アセトアルデヒド</p>	235.8 ± 15	232.30
 <p>(d) 2-クロロ-2-フェニルアセトアルデヒド</p>	219.0 ± 20	220.81

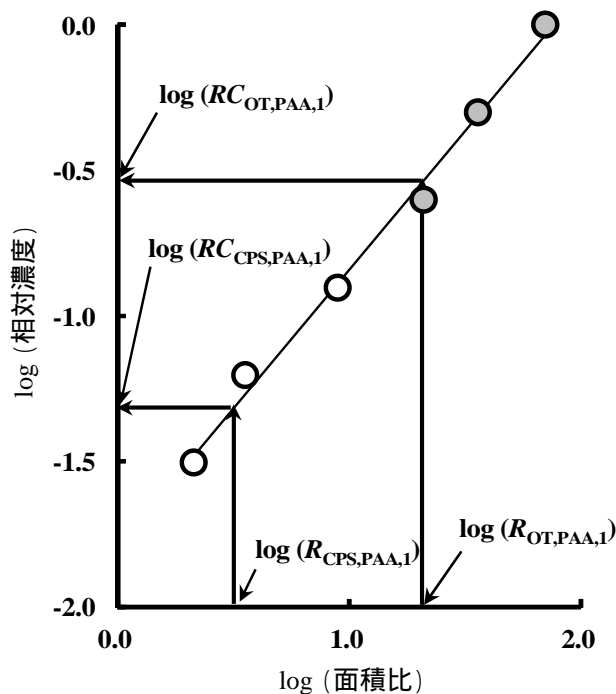


図23 パネラー#1に対する溶媒抽出 - GC-MS-0分析におけるPAAの面積比の常用対数値と相対濃度の常用対数値の関係 灰丸,パネラーがPAAの臭気を感じ;白丸。パネルがPAAの臭気を否感知; $R_{CPS,PAA,1}$ ,フェニルアラニン塩素処理溶液中のPAAの面積比;  $R_{OT,PAA,1}$ ,パネラーが臭気を感じできた最小のPAAの面積比

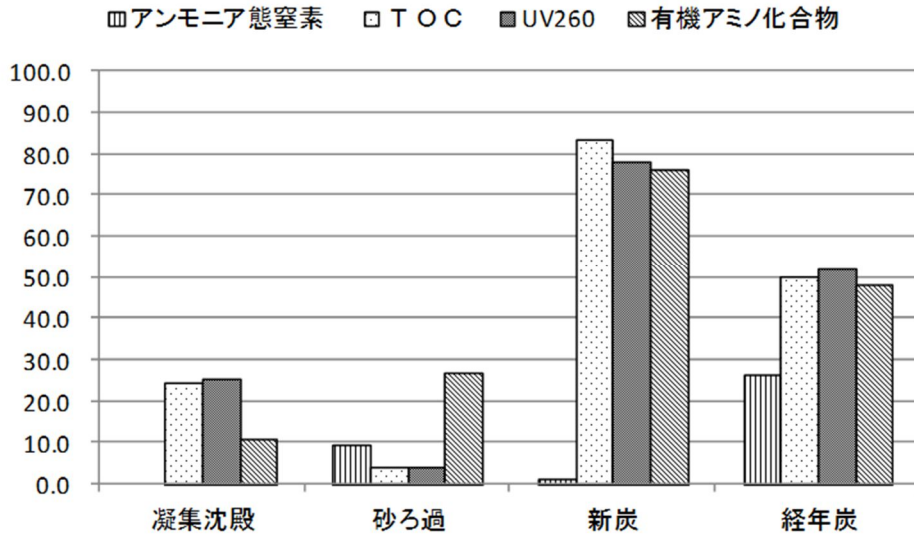


図 24 処理工程における各項目の除去率

表 7 クロラミン生成能

単位：mg/L

試料名	モノクロラミン	ジクロラミン	トリクロラミン
砂ろ過水	0.12	0.50	0.15
新炭処理水	0.02	0.11	0.33
経年炭処理水	0.06	0.24	0.24

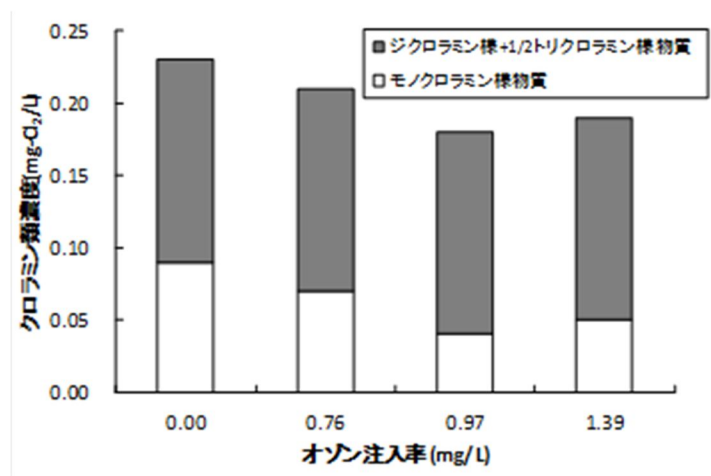


図 25 オゾン処理によるクロラミン類生成状況の変化 (大場川)

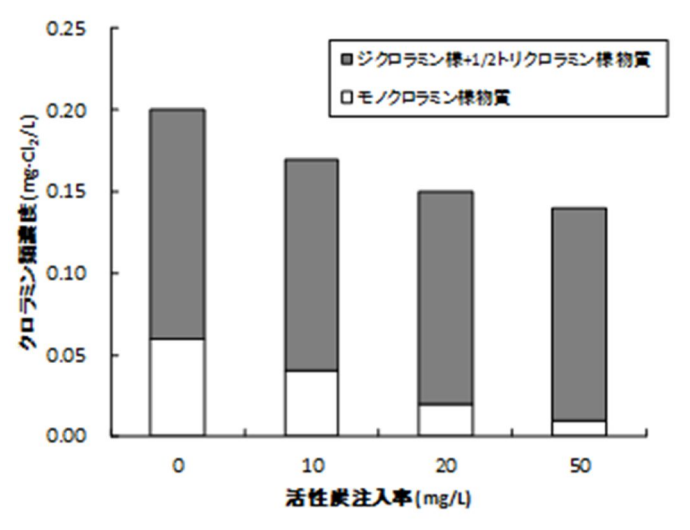


図 26 PAC 処理によるクロラミン類生成状況の変化 (対象水: 大場川と江戸川の混合水 (1:7))

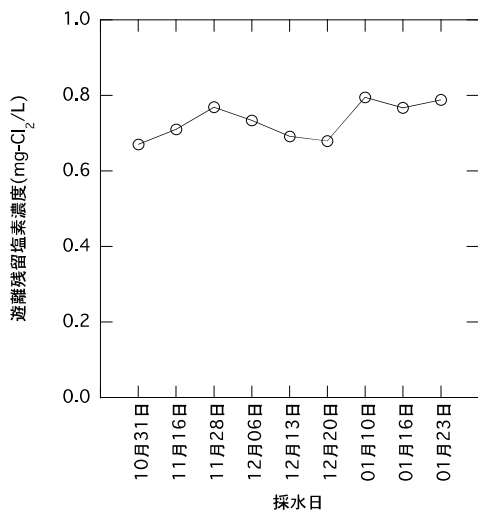


図 27 遊離残留塩素濃度の推移

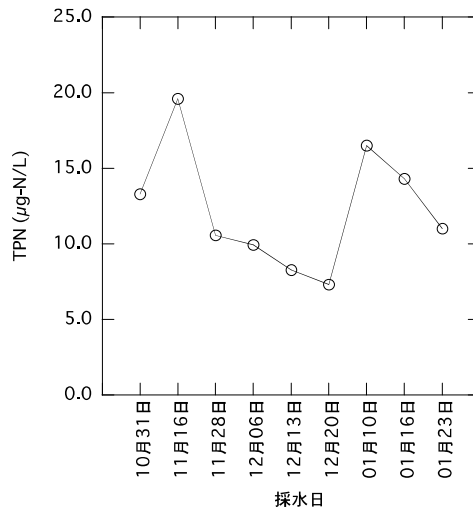


図 28 TPN の推移

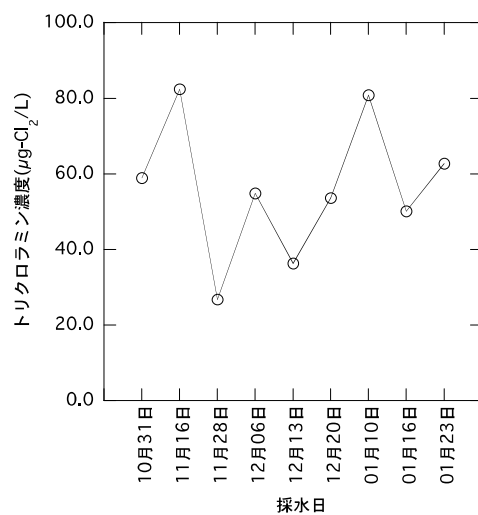


図 29 トリクロラミン濃度の推移

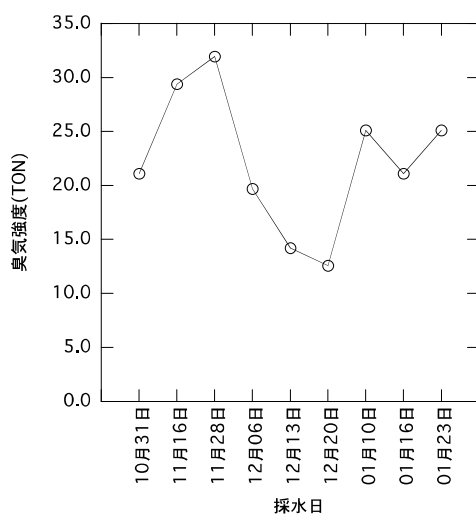


図 30 臭気強度の推移

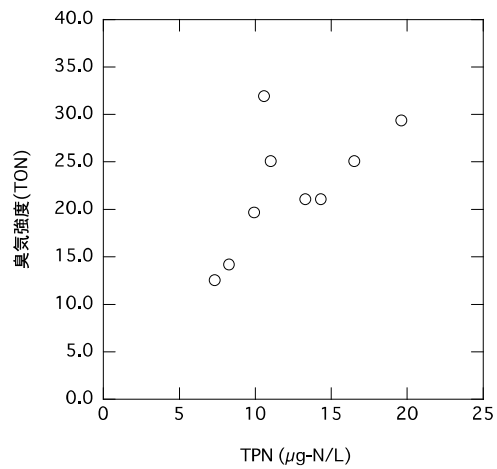


図 31 TPN と臭気強度の比較

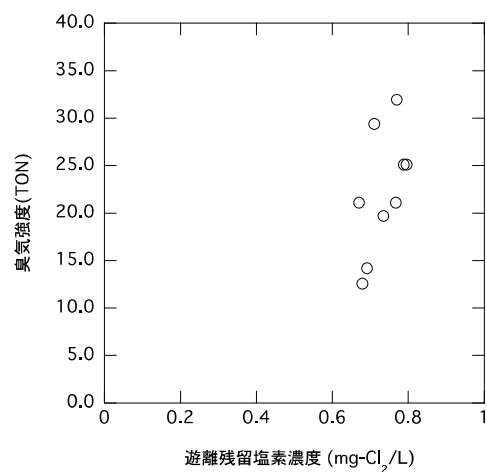
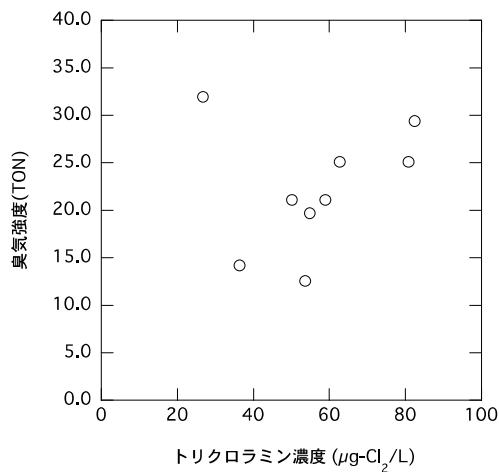


図 32 トリクロラミン濃度と臭気強度の比較 図 33 遊離残留塩素濃度と臭気強度の比較

表 8 塩素注入率を変化させた場合の TPN、トリクロラミン濃度、臭気強度の変化

	採水日	24時間後	TPN (µg-N/L)	トリクロラミン 濃度 (µg-Cl <sub>2</sub> /L)	臭気強度 (TON)	アンモニウム イオン濃度 (µg-N/L)
		遊離残留塩素濃度 (mg-Cl <sub>2</sub> /L)				
1回目	10月24日	0.5	10	83	10	31
		1.0	12	117	34	
		1.5	25	204	71	
2回目	1月10日	0.5	8	95	45	48
		1.0	13	116	62	
		1.5	21	152	87	



表9 pHを変化させた場合のTPN, トリクロラミン濃度、臭気強度の変化

	採水日	24時間後 遊離残留塩素濃度 (mg-Cl <sub>2</sub> /L)	TPN (μg-N/L)	トリクロラミン 濃度 (μg-Cl <sub>2</sub> /L)	臭気強度 (TON)	pH (塩素処理後)
1回目	12月6日	0.6	12	89	22	41
		0.6	20	47	41	
		0.5	5	13	10	
2回目	1月16日	1.0	17	187	56	48
		1.1	22	132	67	
		1.0	9	86	47	