



The Emission and Distribution of PCDD/Fs in Municipal Solid Waste Incinerators and Coal-fired Power Plant

Wen-Yinn Lin¹, Yee-Lin Wu^{2,3*}, Li-Kai Tu², Lin-Chi Wang⁴, Xin Lu²

¹ *Institute of Environmental Engineering and Management, National Taipei University of Technology, Taipei, 10608, Taiwan*

² *Department of Environmental Engineering, National Cheng Kung University, Tainan, 70101, Taiwan*

³ *Sustainable Environment Research Center, National Cheng Kung University, Tainan, 70101, Taiwan*

⁴ *Super Micro Mass Research and Technology Center, Cheng Shiu University, Kaohsiung, 833, Taiwan*

ABSTRACT

The emission and distribution of polychlorinated dibenzo-*p*-dioxins/dibenzofurans (PCDD/Fs) was investigated in two municipal solid waste incinerators (MSWIs) and one coal-fired power plant (PP) in southern Taiwan. Samples were collected from stack flue gases (SFG), bottom residues (BR), super heater (SH), economizer (EC), semi-dry absorber (SDA), bag filter (BF), and fly ash pit (FAP) in MSWIs. Stack flue gases, bottom residues and electrostatic dust collectors (ESD) in PP were also collected. In order to compare the difference between the results of MSWIs and PP, samples from SFG, BS, and FAP in a PP were also determined. Seventeen congeners of PCDD/Fs were analyzed by utilizing a high-resolution gas chromatograph/high-resolution mass spectrometer (HRGC/HRMS). Distributions of total PCDD/F-I-TEQ in each unit of MSWI-A and MSWI-B were SFG (0.3%, 0.07%), BR (3.9%, 0.62%), SH (0.17%, 0.24%), EC (4.2%, 0.05%), SDA (1.29%, 7.06%), and BF (90.14%, 91.97%), respectively. However, those in SFG, BS, and FAP of PP were 99.58%, 0.17%, and 0.25%, respectively. The above results indicated 99.5% PCDD/Fs were trapped in the fly ash of MWSI. On the other hand, 99.7% PCDD/Fs was emitted to the atmosphere from PP. The results of this study provide useful information for controlling PCDD/Fs in MSWIs and PP.

Keywords: PCDD/Fs; Coal-fired power plant; Municipal solid waste incinerator.

INTRODUCTION

The sources of PCDD/Fs are mainly human activities, including industrial and heat-treatment processes (Oh *et al.*, 1999; Baker *et al.*, 2000; Tame *et al.*, 2007; Hu *et al.*, 2009; Hu *et al.*, 2009), and are known to be persistent in the environment and animal tissue (Chao *et al.*, 2007). The formation of PCDD/Fs in municipal solid waste incinerators had been researched in many countries (McKay, 2002; Altarawneh *et al.*, 2009; Aurell and Marklund, 2009). With the continuous studying, three known PCDD/Fs formation mechanisms were found for PCDD/Fs in stack flue gases; (i) PCDD/Fs originally present in the feedstock of MSWIs; (ii) from precursor compounds in the MSW feed; (iii) from de novo synthesis of relatively innocuous chemical molecules combining together to form dioxins (McKay, 2002). However, the favourable temperature range of de novo synthesis is 250°C–400°C, and can be negligible when fly

ash was heated at 400°C or higher temperature (Kakuta *et al.*, 2007; Chen *et al.*, 2008). Chlorine, although it is not a precursor of PCDD/Fs, was found to predominate the forming tendency toward dibenzo-*p*-dioxins (PCDDs) or dibenzofurans (PCDFs) with a 0.8–1.1% threshold in the wastes (Wang *et al.*, 2002). The major distributing pathway of PCDD/Fs is through air (Lohmann and Jones, 1998). The amount of PCDD/Fs transported through air is so high that some of the water treatment plants have been suggested to put on a cover (Lin *et al.*, article in press). In order to discover the PCDD/Fs contribution to the environment of several PCDD/Fs emitting activities, the fates of PCDD/F-like compound and PCDD/Fs have been studied for years (Tsai *et al.*, 2001; Kuo *et al.*, 2003; Lee *et al.*, 2004; Van Caneghem *et al.*, 2010). Since tighter emission limits have been applied to incinerators, sinter plants have become the dominating PCDD/F emission sources (Wang *et al.*, 2003). Not only sinter plants, but also secondary aluminum smelters and electric arc furnaces emit more PCDD/Fs to the environment than MSWIs (Chen *et al.*, 2004; Lee *et al.*, 2005). Electric arc furnace dust treatment plant was found to provide PCDD/Fs to the downwind duck farms significantly higher than upwind farms (Lee *et al.*, 2009). Attempting to get rid

* Corresponding author. Tel.: 886-6-2386764;
Fax: 886-6-275-2790
E-mail address: ylwu@aerosol.ev.ncku.edu.tw

to stubborn PCDD/Fs from the soil, thermal treatment had been developed, and is proven as an effective technology to remove 99% PCDD/Fs from heavily contaminated soils when the treatment process was above 750°C (Lee *et al.*, 2008). Coal-fired power plant is the dominating PCDD/Fs emission source in southern Taiwan (Lin *et al.*, 2007). The coal ash contains organic constituents of potential environmental concern just like fly ash of MSWIs (Chen *et al.*, 2006; Sahu *et al.*, 2009). Comparison between MSWIs and coal-fired power plant could give us a comprehensive scope of the distribution of PCDD/Fs in different units of each plant due to different types of air pollution control devices were equipped.

THE SAMPLING INFORMATION

Basic Information of Sampling Sites

The stack flue gas (SFG) samples and ash samples were collected from MSWI-A, MSWI-B, and PP. The three sites are all located in southern Taiwan. During the sampling, there were three furnaces operating in MSWI-A and MSWI-B, the active carbon injected for each furnace in MSWI-A and MSWI-B was 9 kg/hr and 7 kg/hr, respectively; the wastes burnt were 1140 and 1185 metric tonnes/day. The municipal and industrial wastes treated in MSWI-A and B were 40% vs. 60% and 56% vs. 44%, respectively. Each furnace is two-stage and starved-air modular type. The air pollution control devices (dry scrubber, activated carbon injection, and fabric filter) which is the most universal combination in MSWIs of Taiwan installed in each MSWI were recognized as the most effective techniques for PCDD/Fs control (Wang, *et al.*, 2009). For PP, two furnaces and four electrostatic dust collectors (ESD) were functioning, and 4656 metric tonnes/day of coals were consumed as fuel during the sampling.

The Stack Flue Gas Sample Collection

Totally fifteen stack flue gas (SFG) samples, five samples from each site, were collected for PCDD/Fs analysis. The sampling method was in compliance with the standard sampling procedure of Dioxin and Furan in flue pipe, NIEA A807.74C, which was issued by Environmental Analysis laboratory EPA, Executive Yuan, (R.O.C). The stack flue gases were collected isokinetically and then the probe was cleansed with the order of acetone, dichloromethane, and toluene for the sample collecting. The sampling train adopted in this study was comparable with which specified by U.S. EPA modified method 5. No sealing grease was used during the train parts assembling. The gas density determination equipment is qualified with US EPA methods 3 and 4. Amberlite XAD-2[®] was used as the adsorbent. Prior to sampling, 20–40 g XAD-2[®] was loaded in the cartridge, and was spiked with PCDD/F surrogate standards pre-labeled with isotopes, ³⁷Cl₄-2,3,7,8-TCDD, ¹³C₁₂-1,2,3,4,7,8-HxCDD, ¹³C₁₂-2,3,4,7,8-PeCDF, ¹³C₁₂-1,2,3,4,7,8-HxCDF, and ¹³C₁₂-1,2,3,4,7,8,9-HpCDF. The PCDD/F surrogate standard recoveries were, 72.4–113.4%, reaching the criteria within 70–130%. After

the samples were collected, they were preserved under 10°C and shipped back the lab for further analysis.

The Ash Sampling in MSWI Units

In order to find out the characteristics of PCDD/Fs in MSWI ashes, those in MSWI units and the stack gas flue samples were collected at the same time. The typical ash samples were collected from six different MSWI units and two different PP units, which were bottom residues (BS), super heater (SH), economizer (EC), semi-dryer absorber (SDA), bag filter (BF), and fly ash pit (FAP) in MSWIs; bottom residues (BS) and electrostatic dust collectors (ESD) in PP. The ashes in FAP were the mixture of those in SH, EC, SDA, and BF. Solid waste sample collecting method (NIEA R119.00C) issued by NIEA was enforced to ensure quality of the sample. For each ton of solid waste burnt, ashes collected in each unit of MSWIs were 15%, 1%, 1%, 1%, 9%, and 12% for MSWI-A; and 12.5%, 0.83%, 0.83%, 0.83%, 7.5% and 9.99% for MSWI-B. For each ton of coal burnt, ashes equal to 1.46% and 5.86% weight of burnt coal will be generated in BR and ESD of PP. The sampling method of every chosen MSWI and PP part was the same. Ashes were collected every 12 hours for 3 days, and 200 grams were collected each time with the total of 1.2 kilograms. The collected samples were stored in properly sealed containers to prevent the amount of moisture in the samples being affected by air circulation. During the transportation, the samples were preserved under 4 ± 1°C, except the solidified samples.

PCDD/Fs Analysis

The samples had been pretreated before analysis. The ash samples were put on clean utensils or clean section of the foil, removed the impurities, and then wind-dried naturally or freeze dried. The pellets need to be shattered in order to prevent the dehydrated solid samples from cemented tightly during natural wind-dried process, if the diameter of pellets was greater than 15 mm. Solidified samples could be cracked and crushed to make them smaller than 5 mm before wind-dried naturally. After the drying, the samples were first sieved with 2 mm (10 mesh) standard sieve and then grinded to make them passing the 18 mesh (aperture < 1 mm) version. The sieved ashes were mixed properly and put into the flask to wait for being extracted.

The stack flue gas samples were put in a Soxhlet extractor with internal standard spiking solution (23IS) 30 µL and extracted for 18 ± 2 hours. The extract was then evaporated till almost dry out and was dissolved in dichloromethane three times in order to transfer to a clean tube. Each extract was separated equally into A and B flask. Flask A was taken to be acid-washed and flask B was stored.

The ash samples were put in thimble filters and were then moved in a Soxhlet extractor with internal standard spiking solution (1613LCS) and extracted on heat for the 22 ± 2 hours. The extract was cooled to room temperature and was then evaporated to near dryness.

The extracts were treated with sulfuric acid, and vibrated in an ultrasonic oscillator. A series of sample cleanup and fraction procedures, including acidic silica gel

column, acidic alumina column, and activated carbon chromatography, were used to treat the extract. The final extracts were blown with nitrogen to near dryness and RS (for stack flue gas samples) or ISS (for ash samples) were poured in the concentrates.

The analyses of PCDD/Fs were carried out by a high-resolution gas chromatographer/high-resolution mass spectrometer (HRGC/HRMS). Seventeen PCDD/F congeners were analyzed. The column equipped by HRGC was heated up from 150°C to 190°C with a raise of 20 °C/min, and was then raised up to 220°C with 1.5 °C/min. The temperature was then went up to 310°C with 3 °C/min and was maintained for 2 minutes. The HRMS was equipped with an electron impact (EI+) source. The analytical mode of selected ion monitoring (SIM) had a resolving power of 10,000. The temperature of the ion source was 250°C. The more details of analysis procedure could be found in Wang *et al.* (2003) and Chen *et al.* (2008).

RESULTS AND DISCUSSION

PCDD/Fs Characteristics in Stack Flue Gas Samples

Table 1 shows the averaged PCDD/F concentration and relative standard deviations (RSDs) for samples collected from MSWI-A, MSWI-B and PP. OCDD (0.3762 ng/Nm³ in MSWI-A, 0.4434 ng/Nm³ in MSWI-B, and 0.0785 ng/Nm³ in PP), and 1,2,3,4,6,7,8-HpCDD (0.248 ng/Nm³

in MSWI-A, 0.336 ng/Nm³ in MSWI-B, and 0.0683 ng/Nm³ in PP) were the dominating congeners, the above results could be compared with the results of Chao *et al.* (2004) and Jin *et al.* (2009). Ratio of PCDFs/PCDDs was greater than 1 in MSWI-B, indicating that the de nova synthesis were the primary PCDD/Fs formation pathway during the combustion (Huang and Buekens, 1995; Everaert and Baeyens, 2002). However, the ratio of PCDFs/PCDDs was 0.382 and 0.988 in MSWI-A and PP which were both smaller than 1, indicating precursors were the primary PCDD/F formation pathway in MSWI-A and PP. The mean PCDD/F toxic equivalent concentrations were 0.0327 I-TEQ/Nm³, 0.0784 I-TEQ/Nm³, and 0.015 I-TEQ/Nm³ which were all lower than the regulation 0.1 I-TEQ/Nm³ for PCDD/F emission in large MSWIs and PP, respectively. The main toxic equivalent in the stacks of MSWI-A and PP were PCDFs which were indicated by the ratio of PCDD equivalent to PCDF equivalent. PP and MSWIs had a similar congener profile, but the PCDD/Fs concentration in the stack flue gas of PP (0.332 ng/Nm³) was much lower than those of MSWIs (0.947 ng/Nm³ for MSWI-A, 1.78 ng/Nm³ for MSWI-B). Table 2 shows the emission rates of PCDD/Fs which were 68.7, 86.9, and 739 µg/hr for MSWI-A, MSWI-B, and PP. From PCDD/Fs equivalent perspective, the emission rates were 2.37, 3.83, and 38.3 µg I-TEQ/hr for MSWI-A, MSWI-B, and PP. In both view points, PP is the major contributor of PCDD/Fs.

Table 1. Concentration of PCDD/Fs in the stack flue gas of MSWI-A, MSWI-B, and coal-fired power plant.

PCDD/Fs	MSWI-A		MSWI-B		PP	
	Mean (n = 5)	RSD (%)	Mean (n = 5)	RSD (%)	Mean (n = 5)	RSD (%)
2,3,7,8-TeCDD	0.00116	14.9	0.0031	54.8	0.000481	103
1,2,3,7,8-PeCDD	0.00582	14.8	0.0128	51.4	0.00265	72
1,2,3,4,7,8-HxCDD	0.00854	19	0.0152	34.1	0.00374	77.7
1,2,3,6,7,8-HxCDD	0.0294	25	0.0409	22.5	0.00836	81.9
1,2,3,7,8,9-HxCDD	0.016	25.3	0.0243	28.3	0.00766	84.6
1,2,3,4,6,7,8-HpCDD	0.248	46	0.336	15.7	0.0683	96.3
OCDD	0.376	87.7	0.443	15	0.0758	71.8
2,3,7,8-TeCDF	0.00594	14.9	0.0183	51.2	0.00316	32.8
1,2,3,7,8-PeCDF	0.0102	13.9	0.0361	52.6	0.00443	54
2,3,4,7,8-PeCDF	0.0208	12.4	0.0506	72.9	0.0095	70.1
1,2,3,4,7,8-HxCDF	0.0191	14.7	0.065	55.8	0.00966	75.7
1,2,3,6,7,8-HxCDF	0.0214	15.9	0.0698	56.3	0.0111	77
1,2,3,7,8,9-HxCDF	0.00246	33.2	0.00784	49	0.00172	97.2
2,3,4,6,7,8-HxCDF	0.0356	18.3	0.103	50.9	0.023	92.4
1,2,3,4,6,7,8-HpCDF	0.0743	23.7	0.265	47.6	0.0463	95.2
1,2,3,4,7,8,9-HpCDF	0.0155	19.8	0.0662	32.3	0.014	87.5
OCDF	0.0567	28.2	0.222	32.5	0.0425	93.9
PCDDs	0.685	64.7	0.876	16.9	0.167	78.7
PCDFs	0.262	19.7	0.904	45.9	0.165	87.6
PCDF/PCDDs	0.382	29.8	1.03	23.6	0.988	27.9
Total PCDD/Fs (ng/Nm ³)	0.947	50.8	1.78	31.5	0.332	82.4
PCDDs (ng I-TEQ/Nm ³)	0.0123	23.9	0.0214	35.9	0.00454	80.9
PCDFs (ng I-TEQ/Nm ³)	0.0203	13.7	0.0571	61.3	0.0105	76.6
PCDFs/PCDDs (TEQ)	1.65	11.8	2.67	18.9	2.31	13.1
Total TEQ (ng I-TEQ/Nm ³)	0.0327	16.9	0.0784	54.3	0.015	77.8

Table 2. Emission Rate of PCDD/Fs in MSWI-A, MSWI-B, and coal-fired power plant.

PCDD/Fs	MSWI-A		MSWI-B		PP	
	Mean (n = 5)	RSD (%)	Mean (n = 5)	RSD (%)	Mean (n = 5)	RSD (%)
2,3,7,8-TeCDD	0.0839	14.9	0.151	55.4	1.07	105
1,2,3,7,8-PeCDD	0.422	14.2	0.628	51.9	5.87	74.9
1,2,3,4,7,8-HxCDD	0.619	18.4	0.744	34.6	8.29	80.7
1,2,3,6,7,8-HxCDD	2.13	24.5	2	23	18.5	84.9
1,2,3,7,8,9-HxCDD	1.16	24.7	1.19	28.7	17	87.6
1,2,3,4,6,7,8-HpCDD	18	45.7	16.4	16.2	152	99.3
OCDD	27.3	88	21.6	15.4	168	74.4
2,3,7,8-TeCDF	0.43	14.1	0.895	51.7	6.96	34.7
1,2,3,7,8-PeCDF	0.737	13.3	1.76	53.2	9.78	56.8
2,3,4,7,8-PeCDF	1.51	12	2.47	73.5	21.1	73
1,2,3,4,7,8-HxCDF	1.39	14.1	3.18	56.4	21.4	78.6
1,2,3,6,7,8-HxCDF	1.56	15.6	3.41	56.9	24.7	80.1
1,2,3,7,8,9-HxCDF	0.178	33.2	0.383	49.6	3.82	100
2,3,4,6,7,8-HxCDF	2.58	17.7	5.05	51.4	51.2	95.5
1,2,3,4,6,7,8-HpCDF	5.39	23	13	48.1	103	98.3
1,2,3,4,7,8,9-HpCDF	1.12	19.3	3.23	32.8	31.1	90.4
OCDF	4.11	27.8	10.8	33	94.6	96.9
PCDDs ($\mu\text{g/hr}$)	49.7	64.9	42.7	17.3	371	81.7
PCDFs ($\mu\text{g/hr}$)	19	19.1	44.2	46.4	368	90.6
PCDFs/PCDDs	0.382	29.7	1.04	23.4	0.99	27.8
Total PCDD/Fs ($\mu\text{g/hr}$)	68.7	50.8	86.9	32	739	85.5
PCDDs ($\mu\text{g I-TEQ/hr}$)	0.892	23.5	1.04	36.4	10.1	83.9
PCDFs ($\mu\text{g I-TEQ/hr}$)	1.47	13.2	2.79	61.8	23.3	79.6
PCDFs/PCDDs (TEQ)	1.65	11.6	2.68	18.8	2.31	13.1
Total I-TEQ ($\mu\text{g I-TEQ/hr}$)	2.37	16.4	3.83	54.9	33.3	80.8

Therefore, air pollution control devices (APCDs) in PP must be improved in order to the reduce PCDD/F emission.

PCDD/F Characteristics in Ash Samples

TEQs of PCDD/Fs and the averaged PCDD/F concentrations of ash samples collected from different units of MSWIs and PP could be found in Table 3(a1), 3(a2), 3(b1), 3(b2), and 3(c). Fig. 1, Fig. 2, and Fig. 3 show PCDD/F congener profiles in the ashes of each unit and stack flue gases. The averaged PCDD/F concentration in each unit was 0.318, 0.101, 12.9, 2.39, 7.40, and 13.3 ng/g for BR, SH, EC, SDA, BF, and FAP in MSWI-A; 0.405, 5.16, 0.49, 156, 228, and 45.8 ng/g in MSWI-B. In PP, the average PCDD/F concentration was 0.00997 ng/g in bottom residue, and 0.00261 ng/g in electrostatic dust collectors. The bag filter has the first and the second highest PCDD/F content in two MSWIs. In a recent study, bag filter was found having the highest PCDD/Fs concentration (Chang *et al.*, 2006). From PCDD/F equivalent, TEQs of PCDD/Fs were 0.0127, 0.00842, 0.206, 0.0634, 0.491, and 0.639 ng I-TEQ/g for MSWI-A; 0.0164, 0.0095, 0.0202, 2.84, 4.08, and 1.39 ng I-TEQ/g for MSWI-B. The averaged TEQs were 0.000205 and 0.0000746 for PP. The dominating congener in each unit was still OCDD (0.112, 0.122 ng/g in BR; 0.0132, 2.55 ng/g in SH, 7.5, 0.123 ng/g in EC, 1.03, 91.5 ng/g in SDA, and 1.55, 131 ng/g in BF for MSWI-A and MSWI-B; 3.71

and 19.8 ng/g in FAP for MSWI-A and MSWI-B, respectively; 0.00656, 0.00130 ng/g in BR and ESD for PP, respectively). However, the second dominating congener varies in different units. The control strategies of ashes generated from the MSWI-A, MSWI-B and PP are different, because the toxic content varies. In Taiwan, fly ashes from MSWI are treated as hazardous industrial waste and need to be solidified before burying in landfills. However, fly ashes from PP could be used as resource in embankment.

The Distribution of PCDD/Fs in MSWIs and PP

In MSWI-A and MSWI-B, weights of bottom residue were 15% and 12.5% of solid waste feed; weights of fly ash were 8.3% and 6.3% of solid waste feed. In PP, 1.46% weight of coal became bottom residue and 5.56% became fly ash. The emission factors of all congeners could be found in Table 4(a) and 4(b). Fig. 4 and Fig. 5 illustrate the distribution of PCDD/F mass and I-TEQ in MSWI-A, MSWI B and PP, respectively. According to Fig. 4 and Fig. 5, different PCDD/F emitting scenario of MSWIs and PP could be discovered. The concentration of PCDD/Fs in PP was lower than those in MSWIs; however, the emission rate of PCDD/Fs was the highest in PP than those in MSWIs. The extreme outcomes told us that even the PCDD/Fs concentration in PP was low, but the emission does still need to be noticed. The absence of air pollution

Table 3(a1). PCDD/F content in each unit of MSWI-A

PCDD/Fs	MSWI-A					
	BS		SH		EC	
	Mean (n = 2)	RPD (%)	Mean (n = 2)	RPD (%)	Mean (n = 2)	RPD (%)
2,3,7,8-TeCDD	0.000598	33	0.000491	61.8	0.0178	58.4
1,2,3,7,8-PeCDD	0.00203	41	0.00141	39.1	0.048	8.97
1,2,3,4,7,8-HxCDD	0.00175	37.2	0.00095	3.37	0.0485	30.9
1,2,3,6,7,8-HxCDD	0.00422	46	0.000981	8.06	0.0803	34.4
1,2,3,7,8,9-HxCDD	0.00373	55	0.00132	6.06	0.0885	15.6
1,2,3,4,6,7,8-HpCDD	0.0334	18	0.00561	11.6	1.14	48.9
OCDD	0.112	43.1	0.0132	66.4	7.5	72
2,3,7,8-TeCDF	0.00529	4.35	0.00357	108	0.0235	3.84
1,2,3,7,8-PeCDF	0.00639	25.4	0.00683	69.4	0.0621	12.9
2,3,4,7,8-PeCDF	0.00995	37.2	0.00707	62	0.0793	25.4
1,2,3,4,7,8-HxCDF	0.0095	40.1	0.00721	41.8	0.146	30.1
1,2,3,6,7,8-HxCDF	0.00957	44.6	0.00802	46.1	0.17	28.9
1,2,3,7,8,9-HxCDF	0.00601	129	0.00491	151	0.128	145
2,3,4,6,7,8-HxCDF	0.00802	177	0.00332	98.6	0.155	189
1,2,3,4,6,7,8-HpCDF	0.043	25.8	0.0197	27	1.55	49.8
1,2,3,4,7,8,9-HpCDF	0.00553	40.5	0.00493	36.5	0.137	25.6
OCDF	0.0574	14.8	0.012	54.6	1.32	43.9
PCDDs	0.158	31.1	0.024	43	8.91	67.3
PCDFs	0.161	25.5	0.0776	48.4	3.77	42.8
PCDFs/PCDDs	1.07	55.6	3.22	5.64	0.442	26.4
Total PCDD/Fs (ng/g)	0.318	1.89	0.101	46.7	12.7	60.1
PCDDs (ng I-TEQ/g)	0.00303	32.1	0.00159	37.2	0.0825	35.3
PCDFs (ng I-TEQ/g)	0.00966	36	0.00684	58.1	0.123	32.5
PCDFs/PCDDs (TEQ)	3.18	4.08	4.23	22.1	1.5	2.86
Total TEQ (ng I-TEQ/g)	0.0127	34.6	0.00842	54.2	0.206	33.6

Note: BR (bottom residues); SH (super heater); EC (economizer).

Table 3(a2). PCDD/F content in each unit of MSWI-A.

PCDD/Fs	MSWI-A					
	SDA		BF		FAP	
	Mean (n = 2)	RPD (%)	Mean (n = 2)	RPD (%)	Mean (n = 2)	RPD (%)
2,3,7,8-TeCDD	0.00338	43	0.0331	12.4	0.0322	2.8
1,2,3,7,8-PeCDD	0.0121	70.9	0.0928	15.5	0.117	21.5
1,2,3,4,7,8-HxCDD	0.0114	78	0.06	19.5	0.107	79.7
1,2,3,6,7,8-HxCDD	0.0157	75.2	0.0859	7.57	0.175	99.7
1,2,3,7,8,9-HxCDD	0.0199	76.1	0.0776	20.1	0.148	69.7
1,2,3,4,6,7,8-HpCDD	0.187	86.6	0.541	24.2	1.32	120
OCDD	1.03	98.2	1.55	25.8	3.71	134
2,3,7,8-TeCDF	0.0144	63.8	0.181	43.8	0.168	44
1,2,3,7,8-PeCDF	0.0295	61	0.338	13.6	0.348	27.9
2,3,4,7,8-PeCDF	0.0354	68.7	0.356	35.4	0.441	67.6
1,2,3,4,7,8-HxCDF	0.0519	79.7	0.419	17	0.543	68.5
1,2,3,6,7,8-HxCDF	0.0596	82.8	0.476	16.2	0.629	70.2
1,2,3,7,8,9-HxCDF	0.0723	187	0.381	179	0.63	187
2,3,4,6,7,8-HxCDF	0.0253	161	0.224	167	0.215	135
1,2,3,4,6,7,8-HpCDF	0.395	105	1.6	18.2	2.67	118
1,2,3,4,7,8,9-HpCDF	0.0515	103	0.209	23	0.355	118
OCDF	0.373	116	0.772	24.6	1.69	147
PCDDs	1.28	95.2	2.44	20.5	5.61	123
PCDFs	1.11	103	4.96	24.8	7.68	107

Table 3(a2). (continued).

PCDD/Fs	MSWI-A					
	SDA		BF		FAP	
	Mean (n = 2)	RPD (%)	Mean (n = 2)	RPD (%)	Mean (n = 2)	RPD (%)
PCDFs/PCDDs	0.845	10.3	2.08	44.7	1.48	24.0
Total PCDD/Fs (ng/g)	2.39	98.7	7.4	9.87	13.3	114
PCDDs (ng I-TEQ/g)	0.017	70.6	0.109	9.17	0.15	45.3
PCDFs (ng I-TEQ/g)	0.0464	81.3	0.382	31.9	0.488	75.8
PCDFs/PCDDs (TEQ)	2.67	12.6	3.49	22.9	3.13	33.4
Total TEQ (ng I-TEQ/g)	0.0634	78.1	0.491	26.9	0.639	68.8

Note: SDA (semi-dryer absorber); BF (bag filter); FAP (fly ash pit).

Table 3(b1). PCDD/F content in each unit of MSWI-B.

PCDD/Fs	MSWI-B					
	BS		SH		EC	
	Mean (n = 2)	RPD (%)	Mean (n = 2)	RPD (%)	Mean (n = 2)	RPD (%)
2,3,7,8-TeCDD	0.000476	146	0.00636	199	0.000604	161
1,2,3,7,8-PeCDD	0.00232	151	0.0219	200	0.00259	200
1,2,3,4,7,8-HxCDD	0.00231	148	0.0248	199	0.00319	188
1,2,3,6,7,8-HxCDD	0.00584	153	0.038	199	0.00681	193
1,2,3,7,8,9-HxCDD	0.00436	151	0.0405	200	0.00457	187
1,2,3,4,6,7,8-HpCDD	0.0531	161	0.531	200	0.0491	192
OCDD	0.122	162	2.55	200	0.123	186
2,3,7,8-TeCDF	0.00464	118	0.0109	197	0.00546	185
1,2,3,7,8-PeCDF	0.00698	130	0.0334	199	0.0192	194
2,3,4,7,8-PeCDF	0.0122	137	0.0407	198	0.0129	189
1,2,3,4,7,8-HxCDF	0.0142	136	0.0776	199	0.0173	191
1,2,3,6,7,8-HxCDF	0.0146	140	0.0887	199	0.0213	192
1,2,3,7,8,9-HxCDF	0.00587	11.4	0.0118	192	0.00637	161
2,3,4,6,7,8-HxCDF	0.017	195	0.113	200	0.0275	200
1,2,3,4,6,7,8-HpCDF	0.0622	144	0.591	199	0.0775	187
1,2,3,4,7,8,9-HpCDF	0.0109	145	0.0942	199	0.0259	192
OCDF	0.0671	167	0.876	199	0.0872	187
PCDDs	0.19	161	3.21	200	0.19	188
PCDFs	0.216	150	1.94	199	0.3	189
PCDFs/PCDDs	1.28	28.1	1.02	82.3	1.48	14.4
Total PCDD/Fs (ng/g)	0.405	155	5.16	200	0.49	189
PCDDs (ng I-TEQ/g)	0.00354	152	0.0354	200	0.00396	189
PCDFs (ng I-TEQ/g)	0.0129	139	0.0602	199	0.0163	191
PCDFs/PCDDs (TEQ)	4.07	28	2.99	86.7	3.83	15.3
Total TEQ (ng I-TEQ/g)	0.0164	142	0.0952	199	0.0202	190

Note: BR (bottom residues); SH (super heater); EC (economizer).

Table 3(b2). PCDD/F content in each unit of MSWI-B.

PCDD/Fs	MSWI-B					
	SDA		BF		FAP	
	Mean (n = 2)	RPD (%)	Mean (n = 2)	RPD (%)	Mean (n = 2)	RPD (%)
2,3,7,8-TeCDD	0.0498	179	0.0462	18.6	0.0367	40.7
1,2,3,7,8-PeCDD	0.349	190	0.451	14.4	0.206	49.5
1,2,3,4,7,8-HxCDD	0.555	193	0.756	10.2	0.281	47.4
1,2,3,6,7,8-HxCDD	3.25	198	4.45	59.2	1.03	6.96
1,2,3,7,8,9-HxCDD	1.72	196	2.18	33.6	0.574	13.4
1,2,3,4,6,7,8-HpCDD	37.3	198	54.5	67.9	8.76	51.1
OCDD	91.5	196	131	33.6	19.8	70.7

Table 3(b2). (continued).

PCDD/Fs	MSWI-B					
	SDA		BF		FAP	
	Mean (n = 2)	RPD (%)	Mean (n = 2)	RPD (%)	Mean (n = 2)	RPD (%)
2,3,7,8-TeCDF	0.44	193	0.516	51.8	0.244	20.1
1,2,3,7,8-PeCDF	0.823	191	1.12	37	0.508	36.6
2,3,4,7,8-PeCDF	1.67	193	2.39	44.9	0.914	45.2
1,2,3,4,7,8-HxCDF	1.44	187	2.3	6.09	0.972	46.9
1,2,3,6,7,8-HxCDF	1.78	189	2.77	5.42	1.12	35.4
1,2,3,7,8,9-HxCDF	2.63	199	2.52	162	0.829	147
2,3,4,6,7,8-HxCDF	0.163	42.9	2.12	179	0.949	184
1,2,3,4,6,7,8-HpCDF	5.56	177	9.85	43.7	4.52	42.5
1,2,3,4,7,8,9-HpCDF	1.27	189	2.21	63	0.844	9.72
OCDF	5.21	177	8.69	94.6	4.14	20.6
PCDDs	135	196	194	43.9	30.8	58.9
PCDFs	21	184	34.5	35.1	15.1	31.2
PCDFs/PCDDs	0.421	128	0.194	76.1	0.56	86.1
Total PCDD/Fs (ng/g)	156	195	228	31.6	45.8	29.1
PCDDs (ng I-TEQ/g)	1.24	195	1.69	46.9	0.436	8.26
PCDFs (ng I-TEQ/g)	1.59	191	2.4	21.7	0.952	39.6
PCDFs/PCDDs (TEQ)	1.87	64.6	1.47	25.9	2.17	31.6
Total TEQ (ng I-TEQ/g)	2.84	193	4.08	31.9	1.39	29.6

Note: SDA (semi-dryer absorber); BF (bag filter); FAP (fly ash pit).

Table 3(c). PCDD/F content in each different unit of the coal-fired power plant.

PCDD/Fs	PP			
	BS		ESD	
	Mean (n = 2)	RPD (%)	Mean (n = 2)	RPD (%)
2,3,7,8-TeCDD	ND	-	ND	-
1,2,3,7,8-PeCDD	0.0000465	153	0.0000800	200
1,2,3,4,7,8-HxCDD	0.0000045	200	0.0000850	200
1,2,3,6,7,8-HxCDD	0.0000615	122	0.0000950	200
1,2,3,7,8,9-HxCDD	0.0000465	131	0.000105	200
1,2,3,4,6,7,8-HpCDD	0.000687	81.8	0.000214	75.7
OCDD	0.00656	148	0.00130	143
2,3,7,8-TeCDF	0.000294	169	0.0000320	18.8
1,2,3,7,8-PeCDF	0.000186	164	0.0000490	16.3
2,3,4,7,8-PeCDF	0.000162	137	0.0000670	2.99
1,2,3,4,7,8-HxCDF	0.000104	103	0.0000715	23.8
1,2,3,6,7,8-HxCDF	0.000106	108	0.0000675	28.1
1,2,3,7,8,9-HxCDF	0.000022	200	0.0000390	200
2,3,4,6,7,8-HxCDF	0.000085	191	0.0000330	200
1,2,3,4,6,7,8-HpCDF	0.000433	132	0.000291	27.2
1,2,3,4,7,8,9-HpCDF	0.000049	118	0.0000335	2.99
OCDF	0.00113	181	0.000375	103
PCDDs	0.00739	141	0.00155	126
PCDFs	0.00258	157	0.00106	40.3
PCDFs/PCDDs	0.309	36	1.34	98.0
Total PCDD/Fs (ng/g)	0.00997	145	0.00261	91.2
PCDDs (ng I-TEQ/g)	0.000048	132	0.0000103	99.2
PCDFs (ng I-TEQ/g)	0.000157	139	0.0000639	5.79
PCDFs/PCDDs (TEQ)	3.16	11.5	0.159	94.7
Total TEQ (ng I-TEQ/g)	0.000205	137	0.0000742	18.7

Note: BR (bottom residues); ESD (electrostatic dust collectors).

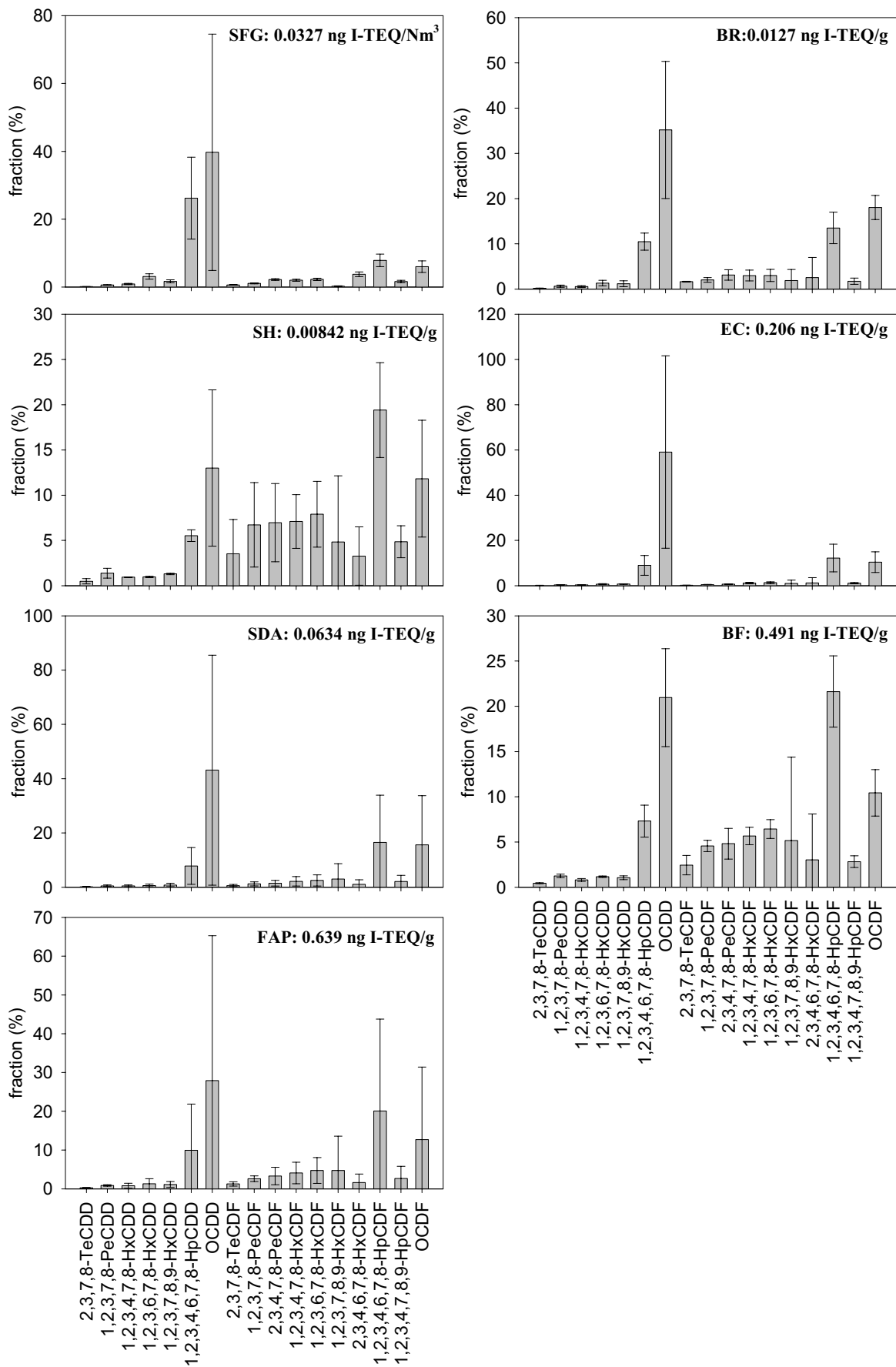


Fig. 1. PCDD/F congener profiles for MSWI A.

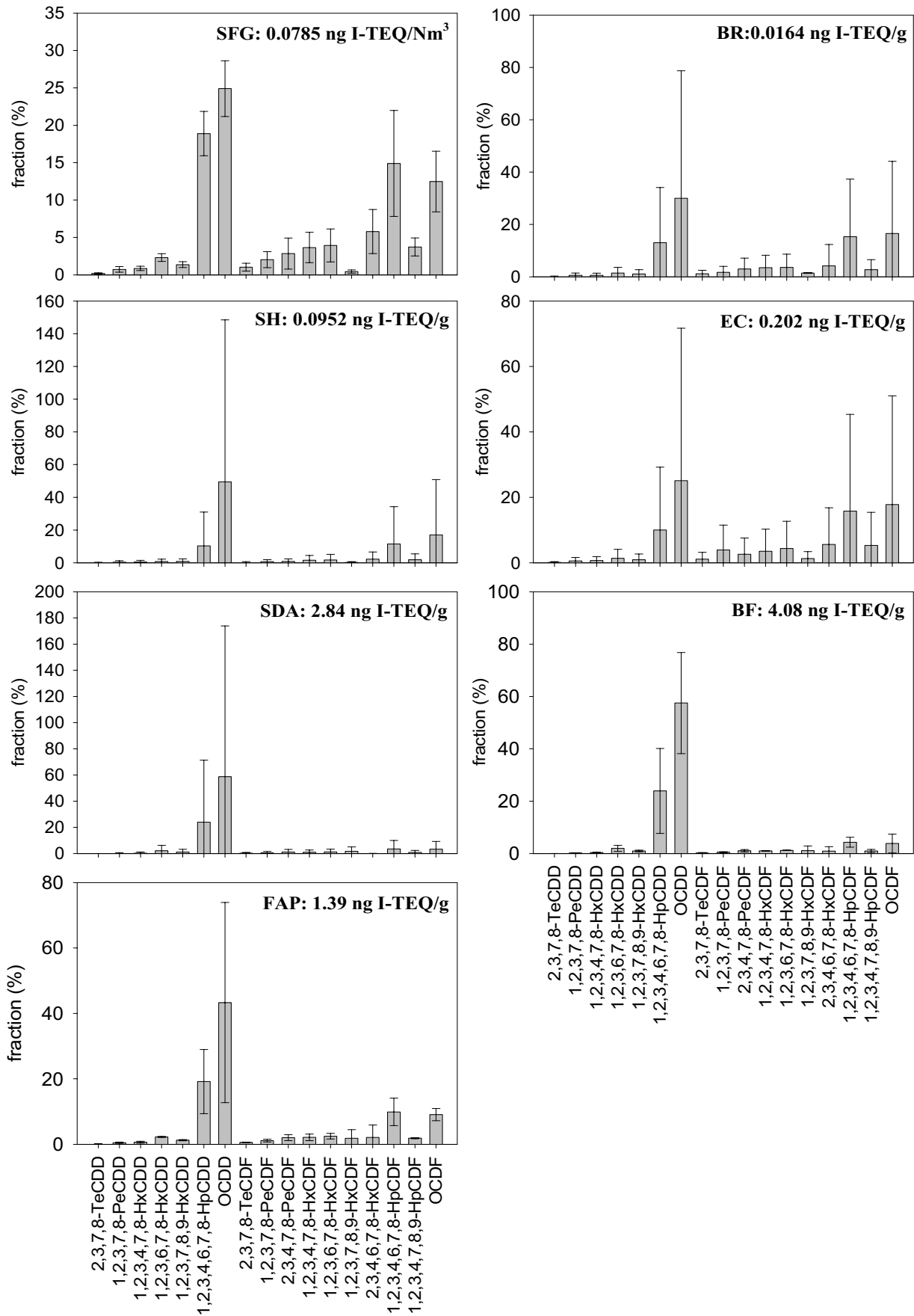


Fig. 2. PCDD/F congener profiles for MSWI B.

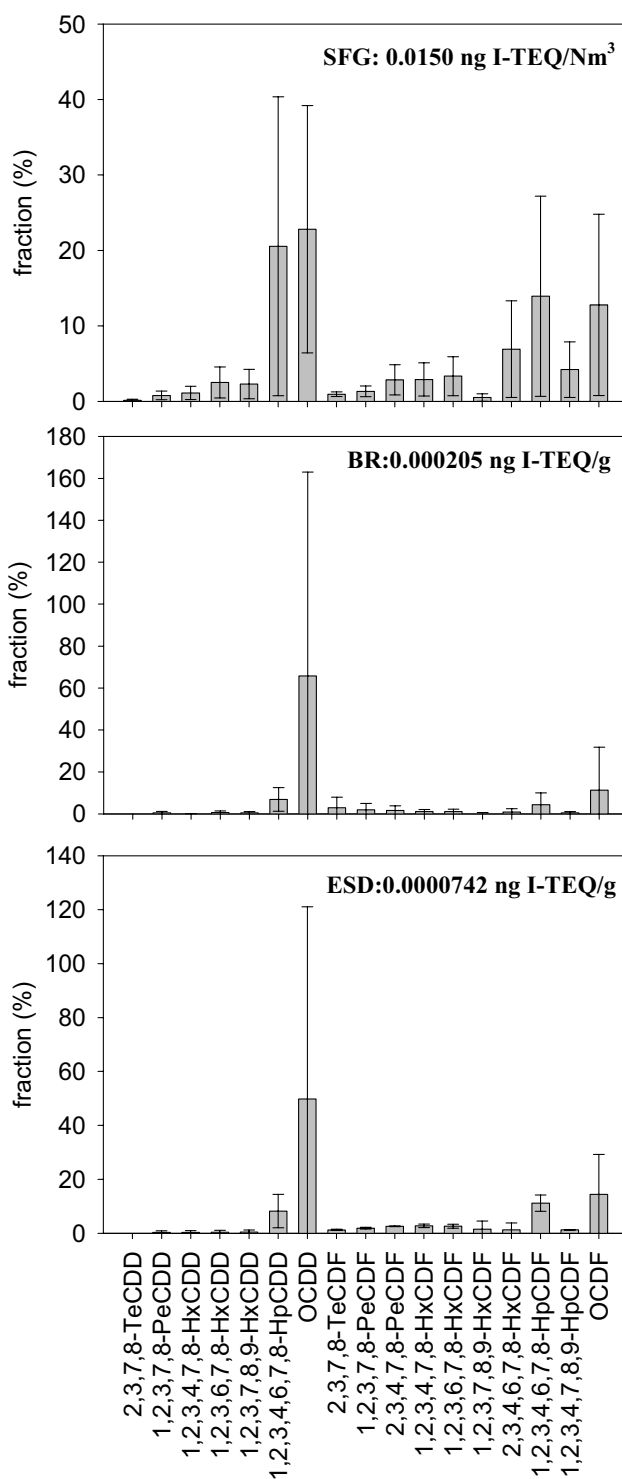


Fig. 3. PCDD/F congener profiles for PP.

control devices suitable for PCDD/F elimination in PP has made the situation worse, 99.58% PCDD/Fs I-TEQ has dispersed into the atmosphere for each ton-coal burnt, instead of trapped in the fly ash just like MWSIs. Only 0.5% or lower PCDD/F amount has been escaped to the environment for every ton-waste treated. The combination of dry scrubber, activated carbon injection, and bag filter installed in MSWIs has been recognized as the most

effective technique for PCDD/Fs control. 71.31% and 91.43% of PCDD/F mass and I-TEQ in MSWI A were trapped in the semi-dryer absorber and the bag filter; 99.45% and 99.03% PCDD/F mass and I-TEQ in MSWI B were found in the semi-dryer absorber and the bag filter, respectively. However, 99.69% of PCDD/F mass and I-TEQ were emitted to the atmosphere from PP. The necessity of installing effective PCDD/F control devices

Table 4(a). The emission factors of BS, SFG and FAP (ESD for PP) in MSWI-A, MSWI-B and the coal fired power plant (unit: $\mu\text{g}/\text{ton-waste}$; $\mu\text{g}/\text{ton-coal}$ for PP).

Mg/ton-waste	SFG			BS			FAP (ESD for PP)		
	MSWI-A	MSWI-B	PP	MSWI-A	MSWI-B	PP	MSWI-A	MSWI-B	PP
2,3,7,8-TeCDD	0.0053	0.0087	0.0545	0.0896	0.0593	ND	3.86	3.66	ND
1,2,3,7,8-PeCDD	0.0266	0.0361	0.308	0.304	0.289	0.000679	14.0	20.6	0.000469
1,2,3,4,7,8-HxCDD	0.0391	0.0429	0.435	0.262	0.287	0.0000657	12.8	28.0	0.000498
1,2,3,6,7,8-HxCDD	0.134	0.115	0.973	0.633	0.727	0.000898	21.0	103	0.000557
1,2,3,7,8,9-HxCDD	0.0732	0.0687	0.894	0.559	0.542	0.000679	17.7	57.3	0.000615
1,2,3,4,6,7,8-HpCDD	1.14	0.949	7.97	5.01	6.60	0.0100	158	875	0.0125
OCDD	1.72	1.25	8.59	16.8	15.1	0.0957	445	1,978.0	0.0764
2,3,7,8-TeCDF	0.0272	0.0516	0.363	0.793	0.577	0.00429	20.2	24.3	0.00188
1,2,3,7,8-PeCDF	0.0465	0.101	0.51	0.959	0.868	0.00271	41.7	50.7	0.00287
2,3,4,7,8-PeCDF	0.0953	0.142	1.1	1.49	1.52	0.00237	52.9	91.3	0.00393
1,2,3,4,7,8-HxCDF	0.0876	0.182	1.12	1.42	1.77	0.00151	65.2	97.1	0.00419
1,2,3,6,7,8-HxCDF	0.0982	0.196	1.29	1.43	1.81	0.00155	75.4	112	0.00396
1,2,3,7,8,9-HxCDF	0.0113	0.022	0.2	0.902	0.730	0.000321	75.6	82.8	0.00229
2,3,4,6,7,8-HxCDF	0.163	0.290	2.69	1.20	2.11	0.00124	25.8	94.8	0.00193
1,2,3,4,6,7,8-HpCDF	0.340	0.745	5.39	6.44	7.74	0.00632	320	452	0.0170
1,2,3,4,7,8,9-HpCDF	0.0709	0.186	1.64	0.830	1.36	0.000715	42.6	84.3	0.00196
OCDF	0.259	0.625	4.96	8.60	8.35	0.0166	202	413	0.0220
PCDDs	3.14	2.47	19.2	23.6	23.6	0.108	673	3,071.9	0.0910
PCDFs	1.2	2.54	19.3	24.1	26.9	0.0376	922	1,503.5	0.0619
PCDFs/PCDDs	0.568	0.193	0.24	1.02	1.14	0.349	1.37	0.489	0.681
Total PCDD/Fs	4.34	5.02	38.5	47.7	50.5	0.145	1,596.6	4,570.4	0.153
PCDDs ($\mu\text{g I-TEQ}/\text{ton-waste}$)	0.0563	0.0602	0.527	0.454	0.441	0.000700	18.0	43.6	0.000603
PCDFs ($\mu\text{g I-TEQ}/\text{ton-waste}$)	0.0931	0.160	1.22	1.45	1.60	0.00229	58.6	95.1	0.00374
PCDFs/PCDDs (TEQ)	0.157	0.0794	0.107	3.19	3.64	3.28	3.25	2.18	6.20
Total PCDD/Fs ($\mu\text{g I-TEQ}/\text{ton-waste}$)	0.149	0.220	1.75	1.91	2.05	0.00300	76.6	138	0.00435

Table 4(b). The emission factors of SH, EC, SDA, and BF in MSWI-A and MSWI-B (unit: $\mu\text{g}/\text{ton-waste}$; $\mu\text{g}/\text{ton-coal}$ for PP).

$\mu\text{g}/\text{ton-waste}$	SH		EC		SDA		BF	
	MSWI-A	MSWI-B	MSWI-A	MSWI-B	MSWI-A	MSWI-B	MSWI-A	MSWI-B
2,3,7,8-TeCDD	0.00491	0.0528	0.178	0.00501	0.0338	0.414	2.97	3.47
1,2,3,7,8-PeCDD	0.0141	0.181	0.480	0.0429	0.121	2.90	8.35	33.8
1,2,3,4,7,8-HxCDD	0.00950	0.206	0.485	0.0265	0.114	4.61	5.40	56.7
1,2,3,6,7,8-HxCDD	0.00981	0.316	0.803	0.0566	0.157	27.0	7.73	333
1,2,3,7,8,9-HxCDD	0.0132	0.336	0.885	0.0379	0.199	14.3	6.98	163
1,2,3,4,6,7,8-HpCDD	0.0561	4.40	11.4	0.408	1.87	310	48.6	4,088
OCDD	0.132	21.2	75.0	1.02	10.3	759	140	9,825
2,3,7,8-TeCDF	0.0357	0.0908	0.235	0.0453	0.144	3.65	16.2	38.7
1,2,3,7,8-PeCDF	0.0683	0.278	0.621	0.159	0.295	6.83	30.4	84.2
2,3,4,7,8-PeCDF	0.0707	0.338	0.793	0.107	0.354	13.9	32.0	179
1,2,3,4,7,8-HxCDF	0.0721	0.644	1.46	0.144	0.519	11.9	37.7	173
1,2,3,6,7,8-HxCDF	0.0802	0.736	1.70	0.177	0.596	14.8	42.8	207
1,2,3,7,8,9-HxCDF	0.0491	0.0977	1.28	0.0528	0.723	21.8	34.3	189
2,3,4,6,7,8-HxCDF	0.0332	0.939	1.55	0.457	0.253	1.35	20.2	159
1,2,3,4,6,7,8-HpCDF	0.197	4.90	15.5	0.643	3.95	46.2	144	739
1,2,3,4,7,8,9-HpCDF	0.0493	0.782	1.37	0.215	0.515	10.5	18.8	165
OCDF	0.120	7.27	13.2	0.724	3.73	43.2	69.5	652

Table 4(b). (continued).

µg/ton-waste	SH		EC		SDA		BF	
	MSWI-A	MSWI-B	MSWI-A	MSWI-B	MSWI-A	MSWI-B	MSWI-A	MSWI-B
PCDDs	0.240	26.7	89.1	1.58	12.8	1,118.4	220	14,513
PCDFs	0.776	16.1	37.7	2.49	11.1	174	446	2,584
PCDFs/PCDDs	3.24	0.603	0.423	1.58	0.866	0.156	2.03	0.178
Total PCDD/Fs	1.01	42.8	127	4.07	23.9	1,291.6	666	17,100
PCDDs (µg I-TEQ/ton-waste)	0.0159	0.294	0.825	0.0329	0.170	10.3	9.81	126
PCDFs (µg I-TEQ/ton-waste)	0.0684	0.499	1.23	0.135	0.464	13.2	34.4	180
PCDFs/PCDDs (TEQ)	4.31	1.70	1.49	4.11	2.73	1.28	3.50	1.42
Total PCDD/Fs (µg I-TEQ/ton-waste)	0.0842	0.790	2.06	0.168	0.634	23.5	44.2	306

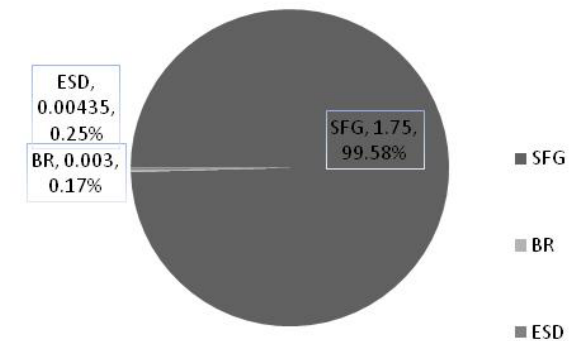
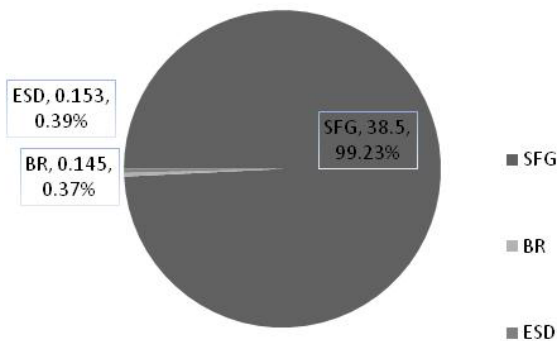
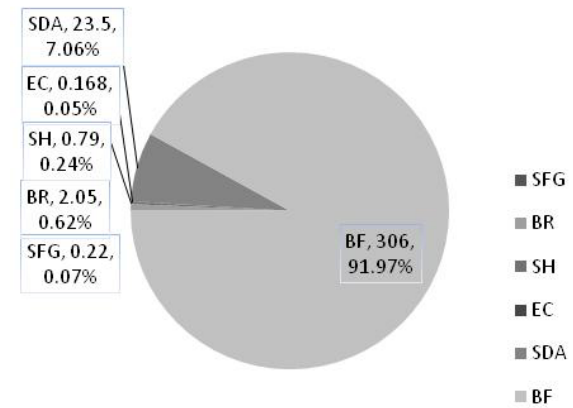
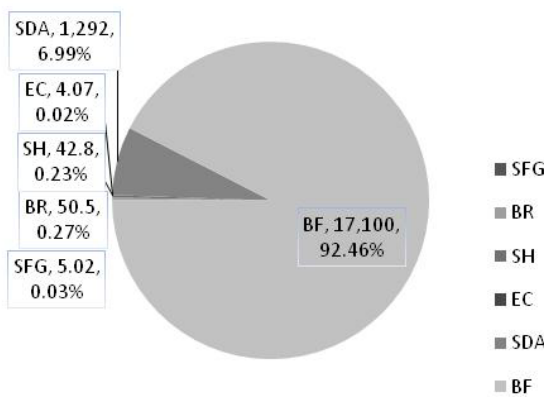
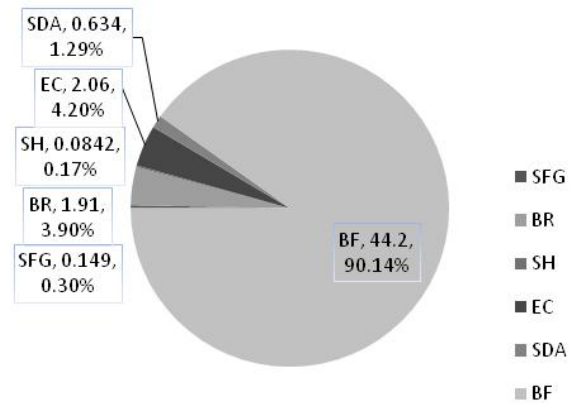
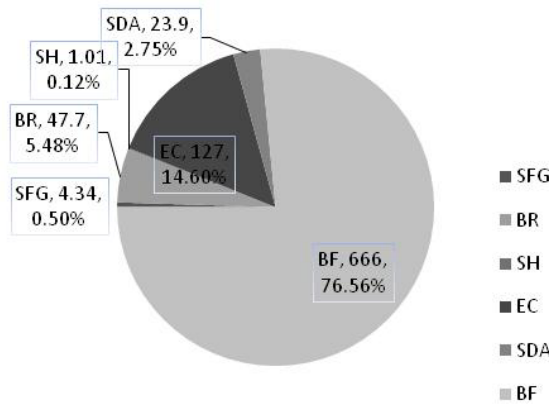


Fig. 4. PCDD/F mass distribution in municipal solid waste incinerator A, B, and coal-fired power plant.

Fig. 5. PCDD/F I-TEQ distribution in municipal solid waste incinerator A, B and coal-fired power plant.

such as activated carbon injection and bag filters in coal-fired power plant needs to investigate further in order to mitigate PCDD/F emission from PP. The pattern of PCDD/F mass and I-TEQ distribution was very similar in each MSWI; however, the resemblance between the MSWIs is unapparent. The different PCDD/F emission characteristics of MSWIs and coal-fired power plant should be noticed. The PCDD/F emission rate is high in coal-fired power plant and low in MSWIs, but the emitting PCDD/F concentration is high in MSWIs and low in the coal-fired power plant. Thus, the PCDD/F contribution of the coal-fired power plant still needs to be marked, because the average amount of coal burnt in PP were 16,442 ton/day, and the solid waste burnt in MSWIs was 1140–1185 ton/day. Although the emission factor of PCDD/Fs from PP is low. After multiplying with the coal burnt per day, the mass of PCDD/Fs provided by PP would be 29102 µg I-TEQ/day, which could be compared with those of PCDD/Fs emitted from MSWIs (84,360–97,170 µg I-TEQ/day). The PCDD/F emission provided by PP is 29.9%–34.5% of those provided by MSWIs.

Improving PCDD/F control devices for the coal-fired power plant is suggested.

CONCLUSIONS

According to the experimental results, the following conclusions could be made:

1. Total PCDD/F mass and I-TEQ distribution of municipal solid waste incinerator A are SFG (0.5%, 0.3%), BR (5.48%, 3.90%), SH (0.12%, 0.17%), EC (14.6%, 4.2%), SDA (22.75%, 12.9%), and BF (76.56%, 90.14%), respectively.
2. Total PCDD/F mass and I-TEQ distribution of municipal solid waste incinerator B are SFG (0.023%, 0.07%), BR (0.27%, 0.62%), SH (0.23%, 0.24%), EC (0.02%, 0.05%), SDA (6.99%, 7.06%), and BF (92.46%, 97.97%), respectively.
3. Total PCDD/F mass and I-TEQ distribution of the coal-fired power plant are SFG (99.23%, 99.58%), BR (0.37%, 0.17%) and ESD (0.39%, 0.25%), respectively. The patterns of total PCDD/F I-TEQ and mass distribution in each site represent a similarity. The semi-dryer absorber and the bag filter prevented 79.31% and 99.03% of PCDD/F mass and I-TEQ in MSWI A, 99.23% and 99.58% of PCDD/F mass and I-TEQ in MSWI B, escaping to the environment. Moreover, only 0.77% PCDD/F mass and 0.42% PCDD/F I-TEQ were found in the ashes of coal-fired plant. The improvement of air control devices for PCDD/F in the coal-fired power plant is advised.

REFERENCES

Altarawneh, M., dlugogorski, B.Z., Kennedy, E.M. and Mackie, J.C. (2009). Mechanisms for Formation, Chlorination, Dechlorination and Destruction of Polychlorinated Debenzo-*p*-dioxins and Dibenzofurans (PCDD/Fs). *Prog. Energy Combust. Sci.* 35: 245–274.

- Aurell, J. and Marklund, S. (2009). Effects of Varying Combustion Condition on PCDD/F Emissions and Formation during MSW Incineration. *Chemosphere* 75: 667–673.
- Baker, J.I. and Hites, R.A. (2000). Is Combustion the Major Source of Polychlorinated Debenzo-*p*-dioxins and Dibenzofurans to the Environment? A Mass Balance Investigation. *Environ. Sci. Technol.* 34: 2879–2886.
- Chang, M.B., Huang, H.C., Tsai, S.S., Chi, K.H. and Chang-Chien, G.P. (2006) Evaluation of the Emission Characteristics of PCDD/Fs from Electric Arc Furnaces; *Chemosphere* 62: 1761–1773.
- Chao, H.R., Wang, S.L., Lin, L.Y., Lee, W.J. and Pöpke, O. (2007). Placental Transfer of Polychlorinated Debenzo-*p*-dioxins, Dibenzofurans, and Biphenyls in Taiwanese Mothers in Relation to Menstrual Cycle Characteristics. *Food Chem. Toxicol.* 45: 259–265.
- Chao, M.R., Hu, C.W., Chen, Y.L., Chen, Chang-Chien, G.P., Lee, W.J., Chang, L.W., Lee, W.S. and Wu, K.Y. (2004). Approaching Gas-particle Partitioning Equilibrium of Atmospheric PCDD/Fs with Increasing Distance from an Incinerator: Measurements and Observations on Modeling. *Atmos. Environ.* 38: 1501–1510.
- Chen, C.K., Lin, C., Wang, L.C. and Chang-Chien, G.P. (2006). The Size Distribution of Polychlorinated Debenzo-*p*-dioxins and Dibenzofurans in the Bottom Ash of Municipal Solid Waste Incinerators. *Chemosphere* 65: 514–520
- Chen, C.K., Lin, C., Lin, Y.C., Wang, L.C. and Chang-Chien, G.P. (2008). Polychlorinated Debenzo-*p*-dioxins/ Dibenzofurans Mass Distribution in both Start-up and Normal Condition in the Whole Municipal Solid Waste Incinerator. *J. Hazard. Mater.* 160: 37–44.
- Chen, S.J., Lee, W.S., Chang-Chien, G.P., Wang, L.C., Lee, W.J., Kao, J.H. and Hu, M.T. (2004). Characterizing Polychlorinated Dibenzofurans and Dibenzofurans in the Surrounding Environment and Workplace of a Secondary Aluminum Smelter. *Atmos. Environ.* 38: 3729–3732.
- Everaert, K. and Baeyens, J. (2002). The Formation and Emission of Dioxins in Large Scale Thermal Processes. *Chemosphere* 46: 439–448
- Hu, M.T., Chen, S.J., Huang, K.L., Lin, Y.C., Lee, W.J., Chang-Chien, G.P., Tsai, J.H., Lee, J.T. and Chiu, C.H. (2009). Characterization of, and Health Risks from, Polychlorinated Dibenzofurans/Dibenzofurans from Incense Burned in a Temple. *Sci. Total Environ.* 407: 4870–4875.
- Hu, M.T., Chen, S.J., Lai, Y.C. Huang, K.L., Chang-Chien, G.P. and Tsai, J.H. (2009). Characteristics of Polychlorinated Dibenzofurans/Dibenzofuran from Joss Paper Burned in Taiwanese Temples. *Aerosol Air Qual. Res.* 9:369–377
- Huang, H. and Buekens, A. (1995). On the Mechanisms of Dioxin Formation in Combustion Processes. *Chemosphere* 31: 4099–4117.
- Jin, G.Z., Lee, S.J., Park, H., Lee, J.E., Shin, S.K. and Chang, Y.S. (2009). Characteristics and Emission Factors of PCDD/Fs in Various Industrial Wastes in

- South Korea. *Chemosphere* 75: 1226–1331.
- Kakuta, Y., Matsuto, T., Tojo, Y. and Tomikawa, H. (2007). Characterization of Residual Carbon Influencing on De Novo Synthesis of PCDD/Fs in MSWI Fly Ash. *Chemosphere* 68: 880–886.
- Kuo, Y.M., Lin, T.C., Tsai, P.J., Lee, W.J. and Lin, H. Y. (2003). Fate of Polycyclic Aromatic Hydrocarbons during Vitrification of Incinerator Ash in a Coke Bed Furnace. *Chemosphere* 51: 313–319.
- Lee, W.S., Chang-Chien, G.P., Wang, L.C., Lee, W.J., Wu, K.Y. and Tsai, P.J. (2005). Emissions of Polychlorinated Dibenzop-dioxins and Dibenzofurans from Stack Gases of Electric Arc Furnaces and Secondary Aluminum Smelters. *J. Air Waste Manage. Assoc.* 55: 219–226.
- Lee, W.S., Chang-Chien, G.P., Wang, L.C., Lee, W.J., Tsai, P.J., Wu, K.Y. and Lin, C. (2004). Source Identification of PCDD/Fs for Various Atmospheric Environments in a Highly Industrialized City. *Environ. Sci. Technol.* 38: 4937–4944.
- Lee, W.J., Shih, S.I., Chang, C.Y., Lai, Y.C., Wang, L.C. and Chang-Chien, G.P. (2008). Thermal Treatment of Polychlorinated Dibenzop-dioxins and Dibenzofurans from Contaminated Soils. *J. Hazard. Mater.* 160: 220–227.
- Lee, W.J., Shih, S.I., Li, H.W., Lin, L.F., Yu, K.M., Lu, K., Wang, L.C., Chang-Chien, G.P., Fang, K. and Lin, M. (2009). Assessment of Polychlorinated Dibenzop-dioxins and Dibenzofurans Contribution from Different Media to Surrounding Duck Farms. *J. Hazard. Mater.* 163: 1185–1193.
- Lin, L.F., Lee, W.J., Li, H.W., Wang, M.S. and Chang-Chien, G.P. (2007). Characterization and Inventory of PCDD/F Emissions from Coal-fired Power Plants and Other Sources in Taiwan. *Chemosphere* 68: 1642–1649.
- Lin, L.F., Shih, S.I., Su, J.W., Shih, M., Lin, K.C., Wang, L.C., and Chang-Chien, G.P. (in press). Dry and Wet Deposition of Polychlorinated Dibenzop-dioxins and Dibenzofurans on the Drinking Water Treatment Plant. *Aerosol Air Qual. Res.* (in press).
- Lohmann, R. and Jones, K.C. (1998). Dioxins and Furans in Air and Deposition: A Review of Levels, Behaviour and Processes. *Sci. Total Environ.* 219: 53–81.
- McKay, G. (2002). Dioxin Characterization, Formation and Minimization during Municipal Solid Waste (MSW) Incineration: Review. *Chem. Eng. J.* 86: 343–368.
- Oh, J.E., Lee, K.T., Lee, J.W. and Chang, Y.S. (1999). The Evaluation of PCDD/Fs from Various Korean Incinerators. *Chemosphere* 38: 2097–2108.
- Sahu, S.K., Bhangare, R.C., Ajmal, P.Y., Sharma, S., Pandit, G.G. and Puranik, V.D. (2009). Characterization and Quantification of Persistent Organic Pollutants in Fly Ash from Coal Fueled Thermal Power Station in India. *Microchem. J.* 92: 92–96.
- Tame, N.W., Dlugogorski, B.Z. and Kennedy, E.M. (2007). Formation of Dioxins and Furans during Combustion of Treated Wood. *Prog. Energy Combust. Sci.* 33: 384–408.
- Tsai, P.J., Shieh, H.Y., Hsieh, L.T. and Lee, W.J. (2005). The Fate of PAHs in the Carbon Black Manufacturing Process. *Atmos. Environ.* 35: 3495–3501.
- Van Caneghem, J., Block, C., Van Brecht, A., Wauters, G. and Vandecasteele, C. (2010). Mass Balance for POPs in Hazardous and Municipal Solid Waste Incinerators. *Chemosphere* 78: 701–708.
- Wang, L.C., Lee, W.J., Lee, W.S., Chang-Chien, G.P. and Tsai, P.J. (2003). Characterizing the Emission of Polychlorinated Dibenzop-dioxins and Dibenzofurans from Crematories and their Impacts to the Surrounding Environment. *Environ. Sci. Technol.* 37: 62–67.
- Wang, L.C., Lee, W.J., Tsai, P.J., Lee, W.S. and Chang-Chien, G.P. (2003). Emissions of Polychlorinated Dibenzop-dioxins and Dibenzofurans from Stack Flue Gases of Sinter Plants. *Chemosphere* 50: 1123–1129.
- Wang, L.C., Lee, W.J., Lee, W.S., Chang-Chien, G.P. and Tsai, P.J. (2003). Effect of Chlorine Content in Feeding Wastes of Incineration on the Emission of Polychlorinated Dibenzop-dioxins and Dibenzofurans. *Sci. Total Environ.* 302: 185–198.
- Wang, Y.H., Lin, C. and Chang-Chien, G.P. (2009). Characteristics of PCDD/Fs in a Particles Filtration Device with Activated Carbon Injection. *Aerosol Air Qual. Res.* 9: 317–322.

Received for review, March 12, 2010

Accepted, August 18, 2010