

# CHAPTER 6

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## CONSTRUCTED WETLANDS FOR WASTEWATER TREATMENT

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## 74 **6 CONSTRUCTED WETLANDS FOR** 75 **WASTEWATER TREATMENT**

### 76 **6.1 INTRODUCTION**

#### 77 **6.1.1 Constructed Wetlands for Wastewater Treatment**

78 Wetland ecosystems can act as sources, sinks, or transformers of nutrients and carbon (C) (Mitsch and  
79 Gossenlink, 1993). This ability of wetlands has led to a widespread use of natural and constructed wetlands  
80 (CWs) for water quality improvement (Brix, 1997).

81 Constructed wetlands systems are fully human-made wetlands for wastewater treatment, which apply various  
82 technological designs, using natural wetland processes, associated with wetland hydrology, soils, microbes and  
83 plants. Thus, CWs are engineered systems that have been designed and constructed to utilize the natural  
84 processes involving wetland vegetation, soils, and their associated microbial assemblages to assist in treating  
85 wastewater. Synonymous terms to “constructed” include “man-made”, “engineered” or “artificial” (Vymazal,  
86 2007).

87 "Semi-natural treatment wetlands" (SNTWs) for wastewater treatment are natural wetland systems that have  
88 been modified for this purpose. The modifications made within these systems usually are based on increasing the  
89 volume of water reserved (i.e. dams) and constructing channels for targeting the influent and effluent. These  
90 systems can be found in both freshwater and coastal wetlands. The functioning of SNTWs is similar to that of  
91 surface flow CWs.

92 This chapter only provides guidance for CWs and SNTWs for wastewater treatment. Decision tree for finding  
93 the appropriate guidance chapter within this supplement or the *2006 IPCC Guidelines for National Greenhouse*  
94 *Gas Inventories (2006 IPCC Guidelines)* is provided as Figure 1.1 in Chapter 1 of this supplement.

95 Emissions from CWs and SNTWs must be reported in waste sector. If freshwater and coastal wetlands are  
96 modified to SNTWs, inventory compilers should check with relevant land-use category in this supplement to  
97 avoid double-counting.

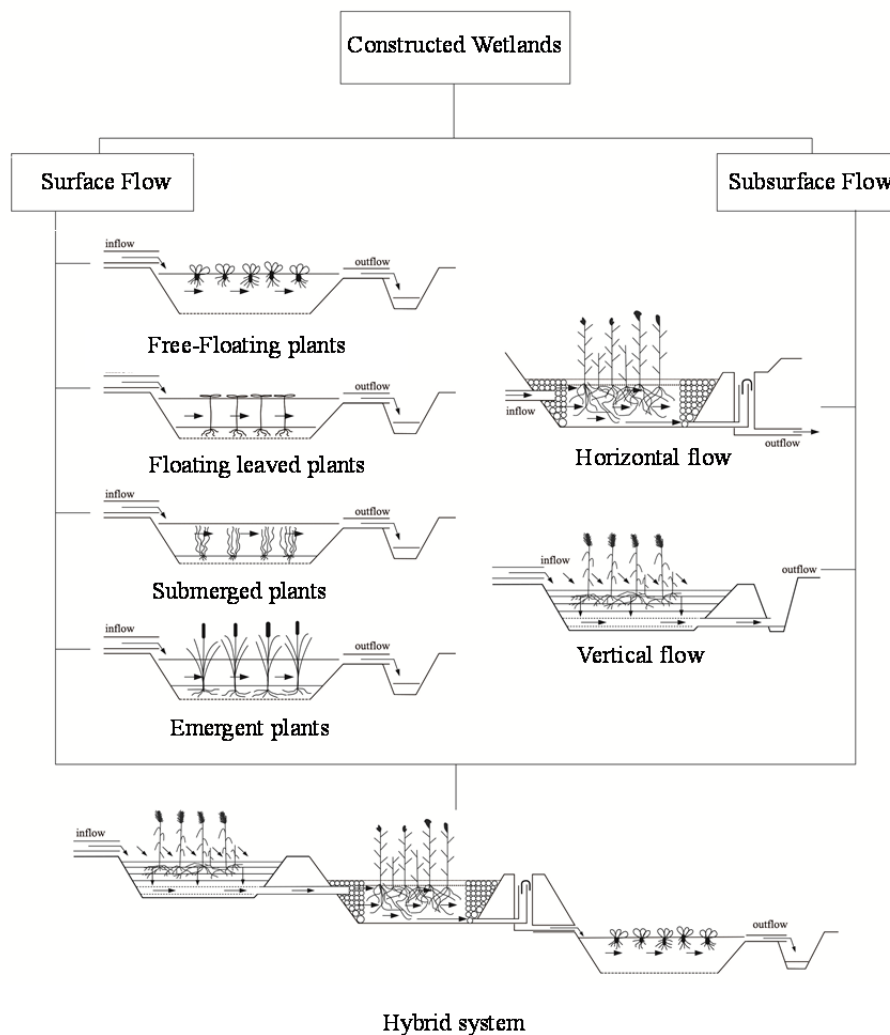
98 Constructed wetlands and SNTWs can be used to improve the quality of collected wastewater including  
99 domestic wastewater, industrial wastewater such as wastewater from processing factories of agricultural products  
100 and dairy farm, collected runoff from agricultural land and leachate from landfill. For some wastewaters, CWs  
101 are the sole treatment; for others, they are one component in a sequence of treatment processes (US EPA, 1995).

102 There are various types of CWs used for treatment of wastewater, and the following paragraphs highlight the  
103 main classification of CWs.

#### 104 **TYPE OF CONSTRUCTED WETLANDS FOR WASTEWATER TREATMENT**

105 Constructed wetlands may be categorized according to the various design parameters, but the three most  
106 important criteria are hydrology (water surface flow and subsurface flow), macrophyte growth form (emergent,  
107 submerged, free-floating, and floating leaved plants) and flow path (horizontal and vertical) (see Figure 6.1;  
108 Vymazal 2007, 2011). Different types of CWs may be combined (which are called hybrid or combined systems)  
109 to utilize the specific advantages of the different systems. For instance, to guarantee more effective removal of  
110 ammonia and total nitrogen (N), during the 1990s and 2000s an enhanced design approach combined vertical and  
111 horizontal flow CWs to achieve higher treatment efficiency (Vymazal, 2011).

112 **Figure 6.1** Classification and configuration of constructed wetlands for wastewater  
 113 treatment



114

115 Note: Adapted from Vymazal, 2007, 2011. Lower part is original. Most of SNTWs represent surface flow type wetlands.

### 116 **Constructed Wetlands with Surface Flow**

117 Constructed wetlands with *surface flow* (SF), known as *free water surface CWs*, contain areas of open water and  
 118 floating, submerged, and emergent plants (Kadlec and Wallace 2008). The shallow water depth, low flow  
 119 velocity, and presence of the plant stalks and litter regulate water flow and, especially in long, narrow channels  
 120 (Crites *et al.* 2005), ensure better water purification. The most common application for SF CWs is for tertiary  
 121 treatment of municipal wastewater and also for stormwater runoff and mine drainage waters (Kadlec and Knight  
 122 1996; Kadlec and Wallace 2008). SF CWs are suitable in all climates, including the far north (Mander and  
 123 Janssen 2003).

### 124 **Constructed Wetlands with Subsurface Flow**

125 In *horizontal subsurface flow constructed wetlands* (HSSF CWs), the wastewater flows from the inlet and flows  
 126 slowly through the porous medium under the surface of the bed planted with emergent vegetation to the outlet  
 127 where it is collected before leaving via a water level control structure (Vymazal *et al.*, 1998). During passage the  
 128 wastewater comes into contact with a network of aerobic, anoxic, and anaerobic zones. Most of the bed is  
 129 anoxic/anaerobic due to permanent saturation of the beds. The aerobic zones occur around roots and rhizomes  
 130 that leak oxygen into the substrate (Brix 1987). HSSF CWs are commonly sealed with a liner to prevent seepage  
 131 and to ensure the controllable outflow. HSSF CWs are commonly used for secondary treatment of municipal  
 132 wastewater but many other applications have been reported in the literature (Vymazal and Kröpfelova 2008).  
 133 The oxygen transport capacity in these systems is insufficient to ensure aerobic decomposition, thus, anaerobic  
 134 processes play an important role in HSSF CWs (Vymazal and Kröpfelova 2008). Some HSSF CWs, having the  
 135 ability to insulate the surface of the bed, are capable of operation under colder conditions than SF systems  
 136 (Mander and Janssen 2003).

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137 *Vertical subsurface flow constructed wetlands* (VSSF CWs) comprise a flat bed of graded gravel topped with  
138 sand planted with macrophytes. VSSF CWs are fed with large intermittent wastewater flows, which flood the  
139 surface of the bed, then percolate down through the bed and are collected by a drainage network at the bottom.  
140 The bed drains completely which allows air to refill the bed. Thus, VSSF CWs provide greater oxygen transfer  
141 into the bed, producing a nitrified (high  $\text{NO}_3^-$ ) effluent (Cooper *et al.*, 1996; Cooper 2005). Consequently, VSSF  
142 CWs do not provide suitable conditions for denitrification to complete conversion to gaseous nitrogen forms,  
143 which then escape to the atmosphere.

144 In recently developed tidal (“fill and drain”) flow systems better contact of wastewater with the microorganisms  
145 growing on the media is guaranteed. This significantly enhances the purification processes (Vymazal 2011).

#### 146 **Hybrid Constructed Wetlands**

147 Various types of CWs can be combined to achieve higher removal efficiency, especially for nitrogen. The design  
148 consists of two stages, several parallel vertical flow (VF) beds followed by 2 or 3 horizontal flow (HF) beds in  
149 series (VSSF-HSSF system). The VSSF wetland is intended to remove organics and suspended solids and to  
150 promote nitrification, while in HSSF wetland denitrification and further removal of organics and suspended  
151 solids occur.

152 Another configuration is a HSSF-VSSF system. The large HSSF bed is placed first to remove organics and  
153 suspended solids and to promote denitrification. An intermittently loaded small VF bed is used for additional  
154 removal of organics and suspended solids and for nitrification of ammonia into nitrate. To maximize removal of  
155 total N, however, the nitrified effluent from the VF bed must be recycled to the sedimentation tank (Vymazal  
156 2011).

157 The VSSF-HSSF and HSSF-VSSF CWs are the most common hybrid systems, but in general, any kind of CWs  
158 could be combined to achieve higher treatment effect (Vymazal 2007).

#### 159 **GREENHOUSE GASES EMISSIONS FROM VARIOUS TYPES OF** 160 **CONSTRUCTED WETLANDS**

161 Emissions of greenhouse gases such as methane ( $\text{CH}_4$ ) and nitrous oxide ( $\text{N}_2\text{O}$ ) are a byproduct of CWs, the  
162 importance of which has been increasing recently. Methane is produced in methanogenesis whereas  $\text{N}_2\text{O}$  is a  
163 product of denitrification and/or nitrification of N compounds by microorganisms. Among several environmental  
164 factors controlling the greenhouse gases emissions, availability of C and nutrients (especially N) which directly  
165 depend on wastewater loading, temperature, hydrological regime (pulsing vs steady-state flow), groundwater  
166 depth, moisture of filter material (water filled soil pores (WFSP)), and presence of aerenchyma plants play a  
167 significant role (see Table 6.1).

168 Soil temperature, oxidation reduction potential and the soil moisture (WFSP, depth of ground water level) are the  
169 most significant factors affecting emissions of  $\text{CH}_4$  from CWs (Mander *et al.*, 2003; Van der Zaag *et al.*, 2010).  
170 Several investigations show that a water table deeper than 20 cm from the surface of wetlands and/or water-  
171 logged soils oxidizes most  $\text{CH}_4$  fluxes (Soosaar *et al.*, 2011; Salm *et al.*, 2012). Fluxes of  $\text{N}_2\text{O}$ , however do not  
172 show a clear correlation with soil/air temperature, and significant emissions of  $\text{N}_2\text{O}$  from CWs have been  
173 observed in winter (Søvik *et al.*, 2006). Likewise, freezing and thawing cycles enhance  $\text{N}_2\text{O}$  emissions (Yu *et al.*,  
174 2011). Hydrological regime also plays a significant role in greenhouse gases emissions from CWs. Altor and  
175 Mitsch (2008) and Mander *et al.*, (2011) demonstrated that the intermittent loading (pulsing) regime and  
176 fluctuating water table in CWs enhance  $\text{CO}_2$  emissions and significantly decrease  $\text{CH}_4$  emissions.  $\text{N}_2\text{O}$  emissions,  
177 in contrast, do not show a clear pattern regarding pulsing regime.

178 Table 6.2 shows  $\text{CH}_4$  and  $\text{N}_2\text{O}$  conversion rates derived from the relationship between the initial (input) C and N  
179 loadings and respective  $\text{CH}_4$  and  $\text{N}_2\text{O}$  emissions from the main types of CWs. There is a significant positive  
180 correlation ( $p < 0.05$ ) between the initial loadings and  $\text{CH}_4$  and  $\text{N}_2\text{O}$  emissions from both SF and VSSF CWs,  
181 whereas no correlation was found for HSSF types. Seemingly, high variability of conditions and combination of  
182 several factors in HSSF CWs may be the reason for that. The limited number of available data did not allow  
183 derivation of reliable relationships for HSSF CWs. These shares (%) can be used as a base for the calculation of  
184 emission factors for Tier 1 and Tier 2 methodologies. The high emission factor for  $\text{CH}_4$  in SF CWs (Table 6.4)  
185 is thought to be due to the additional  $\text{CH}_4$  from sediments accumulated at the bottom of SF CWs.

186

**TABLE 6.1**  
**SELECTED FACTORS IMPACTING CH<sub>4</sub> AND N<sub>2</sub>O EMISSIONS IN CONSTRUCTED WETLANDS**

Factors/processes	CH <sub>4</sub>	N <sub>2</sub> O
Higher water/soil/air temperature	Increase in almost all cases <sup>1-6</sup> with few exceptions <sup>7</sup>	No clear relationship <sup>1-4, 7, 8</sup>
Higher moisture of soil or filter material (higher value of WFSP)	Clear increase <sup>9, 10</sup>	Decrease <sup>9, 10</sup>
Higher wastewater loading	Increase <sup>1-4, 11, 12</sup>	Increase <sup>1, 2, 4, 13</sup>
Presence of aerenchymal plants	Increase <sup>14-16</sup> Decrease (depends on conditions) <sup>17</sup>	Increase <sup>16, 18</sup> Decrease <sup>16, 19</sup>
Pulsing hydrological regime (intermittent loading)	Clear decrease <sup>9, 20</sup>	Increase <sup>9, 21, 22</sup> Decrease in some SF CWs <sup>23</sup>
Deeper water table (from surface) in HSSF CWs	Decrease <sup>9, 10</sup>	Increase <sup>9, 10</sup>

Source:  
<sup>1</sup> Mander and Jenssen 2003; <sup>2</sup> Mander *et al.*, 2005; <sup>3</sup> Teiter and Mander 2005; <sup>4</sup> Søvik *et al.*, 2006; <sup>5</sup> Kayranli *et al.*, 2010; <sup>6</sup> Van der Zaag *et al.*, 2010; <sup>7</sup> Søvik and Kløve 2007; <sup>8</sup> Fey *et al.*, 1999; <sup>9</sup> Mander *et al.*, 2011; <sup>10</sup> Yang *et al.*, 2013; <sup>11</sup> Tanner *et al.*, 1997; <sup>12</sup> Tai *et al.*, 2002; <sup>13</sup> Hunt *et al.*, 2009; <sup>14</sup> Inamori *et al.*, 2007; <sup>15</sup> Inamori *et al.*, 2008; <sup>16</sup> Wang *et al.*, 2008; <sup>17</sup> Maltais-Landry *et al.*, 2009; <sup>18</sup> Rückauf *et al.*, 2004; <sup>19</sup> Silvan *et al.*, 2005; <sup>20</sup> Altort and Mitsch 2008; <sup>21</sup> Jia *et al.*, 2011; <sup>22</sup> Van de Riet *et al.*, 2013; <sup>23</sup> Hernandez and Mitsch 2006

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**TABLE 6.2**  
**INFLUENT TOTAL ORGANIC CARBON (TOC) AND TOTAL NITROGEN (TN) VALUES, RELEVANT CH<sub>4</sub>-C AND N<sub>2</sub>O-N EMISSIONS, AND SHARE (%) OF CH<sub>4</sub>-C AND N<sub>2</sub>O-N IN THE INITIAL LOADING OF TOC AND TN IN CONSTRUCTED WETLANDS**

Type of CW	Influent TOC* (mg C m <sup>-2</sup> h <sup>-1</sup> )	CH <sub>4</sub> -C emission* (mg CH <sub>4</sub> -C m <sup>-2</sup> h <sup>-1</sup> ) <sup>1</sup>	CH <sub>4</sub> -C/ TOC** (%)	Influent TN* (mg N m <sup>-2</sup> h <sup>-1</sup> ) <sup>1</sup>	N <sub>2</sub> O-N emission* (mg N <sub>2</sub> O-N m <sup>-2</sup> h <sup>-1</sup> )	N <sub>2</sub> O-N/TN** (%)
SF	1.04-173.6 (10) <sub>1-11</sub>	0.15-181.0 (10.7) <sup>1</sup> <sub>11</sub>	42 (20)	0.76-202.8 (12) <sup>2, 3, 6-11, 21-23</sup>	0.009-0.65 (0.03) <sup>2, 6-11, 21-23</sup>	0.13 (0.02)
HSSF	15.0-2190.2 (177) <sup>8, 10-12, 15-20</sup>	0.048-17.5 (1.7) <sup>8, 10</sup> <sub>11, 15-20</sub>	12 (6.9)	1.04-295.20 (40) <sup>6, 10, 12, 15-17, 24, 25</sup>	0.014-0.89 (0.10) <sup>6, 10-12, 15-17, 25</sup>	0.79 (0.4)
VSSF	17.88-1417.50 (317) <sup>6, 8, 10, 12</sup>	0.3-5.4 (1.3) <sup>6, 8, 10, 12</sup>	1.17 (0.33)	102.5-2105.0 (155) <sup>6, 8, 10, 12-14</sup>	0.033-0.424 (0.03) <sup>6, 8, 10, 11, 12-14</sup>	0.023 (0.005)

\* Range and standard error (in bracket)  
 \*\* Average and standard error (in bracket)  
 Source: <sup>1</sup> Tanner *et al.*, 1997; <sup>2</sup> Wild *et al.*, 2001; <sup>3</sup> Tai *et al.*, 2002; <sup>4</sup> Johansson *et al.*, 2004; <sup>5</sup> Stadmark and Leonardson 2005; <sup>6</sup> Søvik *et al.*, 2006; <sup>7</sup> Søvik and Kløve 2007; <sup>8</sup> Gui *et al.*, 2007; <sup>9</sup> Ström *et al.*, 2006; <sup>10</sup> Liu *et al.*, 2009; <sup>11</sup> Van der Zaag *et al.*, 2010; <sup>12</sup> Teiter and Mander 2005; <sup>13</sup> Inamori *et al.*, 2007; <sup>14</sup> Wang *et al.*, 2008; <sup>15</sup> Mander *et al.*, 2003; <sup>16</sup> Mander *et al.*, 2008; <sup>17</sup> Liikanen *et al.*, 2006; <sup>18</sup> Garcia *et al.*, 2007; <sup>19</sup> Picek *et al.*, 2007; <sup>20</sup> Chiemchaisri *et al.*, 2009; <sup>21</sup> Xue *et al.*, 1999; <sup>22</sup> Johansson *et al.*, 2003; <sup>23</sup> Wu *et al.*, 2009; <sup>24</sup> Inamori *et al.*, 2008; <sup>25</sup> Fey *et al.*, 1999

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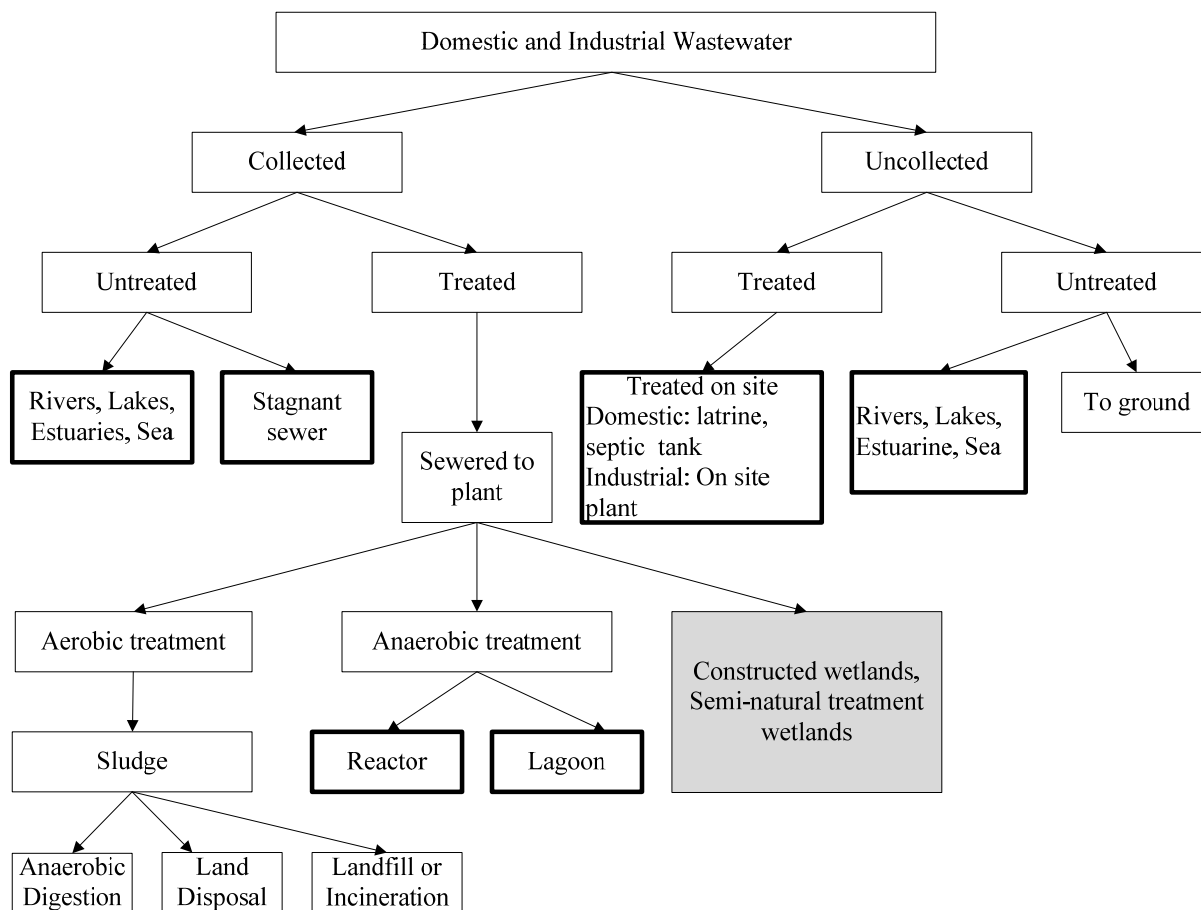
## 6.1.2 Relation to 2006 IPCC Guidelines

190

This chapter is a supplement to Chapter 6 Wastewater Treatment and Discharge of the Volume 5 of the 2006 IPCC Guidelines. The 2006 IPCC Guidelines include a section to estimate CH<sub>4</sub> emissions from uncollected wastewater. The section is expanded in this supplement to cover CH<sub>4</sub> emissions from CWs and SNTWs. This *Wetlands Supplement* includes guidance on estimation of N<sub>2</sub>O emissions from CWs and SNTWs. Emission factors of CH<sub>4</sub> and N<sub>2</sub>O emissions from CWs and SNTWs treating industrial wastewater are the same as those treating domestic wastewater. CO<sub>2</sub> emissions are not included in greenhouse gases emissions from wastewater treatment as CO<sub>2</sub> from wastewater is considered biogenic.

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197 **Figure 6.2 Wastewater treatment systems and discharge pathways**

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200 Note: This figure was modified from the 2006 IPCC Guidelines. Emissions from boxes with bold frames are accounted for in the 2006 IPCC  
 201 Guidelines. This supplement provides emission factors for gray-colored box: CWs and SNTWs for treatment of collected wastewater.

## 202 Coverage of wastewater types and gases

203 Chapter 6 of the Volume 5 of the 2006 IPCC Guidelines provides guidance on estimation of CH<sub>4</sub> and N<sub>2</sub>O  
 204 emissions from domestic wastewater with emission factors based on treatment technology. Constructed wetlands  
 205 in this supplement are an additional treatment technology. The emission factors provided in this chapter cover  
 206 CWs and SNTWs (collected and treated; see Figure 6.2).

207 The methodology is provided for estimation of CH<sub>4</sub> and N<sub>2</sub>O emissions from both domestic and industrial  
 208 wastewater (Table 6.3). The indirect N<sub>2</sub>O emissions from N leaching and runoff from agricultural land are  
 209 covered in Chapter 11, Volume 4 of the 2006 IPCC Guidelines. Emissions from processing factories of  
 210 agricultural products and dairy farm wastewater, collected runoff from agricultural land and leachate from  
 211 landfill are considered as industrial wastewater. According to Chapter 3 of the Volume 5 in the 2006 IPCC  
 212 Guidelines, all amount of degradable organic carbon (DOC) in solid waste is subjected to estimation of CH<sub>4</sub>  
 213 in landfill site, and carbon loss with leachate is not considered because of its low percentage. That means that CH<sub>4</sub>  
 214 emissions from leachate treatment are already covered, and are not included in Section 6.2, while N<sub>2</sub>O emissions  
 215 are considered in Section 6.3 of this supplement. If CH<sub>4</sub> emission from CWs is accounted, the amount of DOC  
 216 in leachate must be subtracted from that in solid waste to avoid double counting. Because C in leachate is normally  
 217 indicated in terms of COD, conversion rate from COD in leachate to TOC in solid waste is required in order to  
 218 subtract the amount of DOC entering CWs from that in solid waste. This logic can be applied in Tier 2 or 3  
 219 estimation.

220



Type of Wastewater	Methane	Nitrous oxide
Domestic wastewater	Included in this supplement (section 6.2) with provision of methane correction factors (MCFs)	Included in this supplement (Section 6.3) with provision of default emission factors
Industrial wastewater including wastewater from processing factories of agricultural products and dairy farm *	Included in this supplement (Section 6.2) with provision of MCFs	Included in this supplement (Section 6.3) with provision of default emission factors
Collected runoff from agricultural land	Emissions can be calculated using same methodology as industrial wastewater and are covered in this supplement (Section 6.2)	Emissions can be calculated using same methodology as industrial wastewater and are covered in this supplement (Section 6.3)  Note: Indirect N <sub>2</sub> O emissions from N leaching and runoff from agricultural land are considered in Chapter 11, Volume 4 of the 2006 IPCC Guidelines. If agricultural runoff is collected and treated by CWs or SNTWs, the amount of N flows into CWs or SNTWs must be subtracted to avoid double counting.
Leachate from landfill	The amount DOC leached from the solid waste disposal site is not considered in the estimation of DOC <sub>f</sub> . Generally the amount of DOC lost with the leachate are less than 1 percent and can be neglected in the calculations (Chapter 3, Volume 5, 2006 IPCC Guidelines) and not considered in this supplement	Emissions can be calculated using same methodology as industrial wastewater and are covered in this supplement (Section 6.3)

221 \*Dairy farm wastewater does not cover manure itself but comes from other activities in the farm.

## 222 6.2 METHANE EMISSIONS FROM CONSTRUCTED 223 WETLANDS

### 224 6.2.1 Methodological issues

225 Methane emissions are a function of the organic materials loaded into CWs and an emission factor.

226 Three tiers of methods for estimation of CH<sub>4</sub> from CWs are summarized below.

227 The Tier 1 method applies default values for the emission factor and activity parameters. This method is  
228 considered *good practice* for countries with limited data.

229 The Tier 2 method follows the same method as Tier 1 but allows for incorporation of country-specific emission  
230 factor and country-specific activity data. For example, a specific emission factor based on field measurements  
231 can be incorporated under this method.

232 The Tier 3 method is used by countries with good data and advanced methodologies. A more advanced country-  
233 specific method could be based on treatment system-specific data such as plant species and composition of  
234 wastewater.

235 In general anaerobic conditions occur in CWs. However, CH<sub>4</sub> generated by CWs is not usually recovered and  
236 combusted in a flare or energy device, and so CH<sub>4</sub> recovery is not considered here.

237 The amount of vegetation harvested from CWs is generally very small and its impact on total emissions from  
238 CWs is considered insignificant. Moreover, the harvesting is usually not performed on regular basis and the  
239 quantity of harvested biomass is commonly not recorded so it is not considered in this supplement.

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240 **6.2.1.1 CHOICE OF METHOD**

241 A decision tree for domestic or industrial wastewater is shown in Figure 6.3.

242 The general equation to estimate CH<sub>4</sub> emissions from CWs treating domestic or industrial wastewater is given in  
243 Equation 6.1.

244

245 **EQUATION 6.1**246 **CH<sub>4</sub> EMISSIONS FROM CONSTRUCTED WETLANDS**

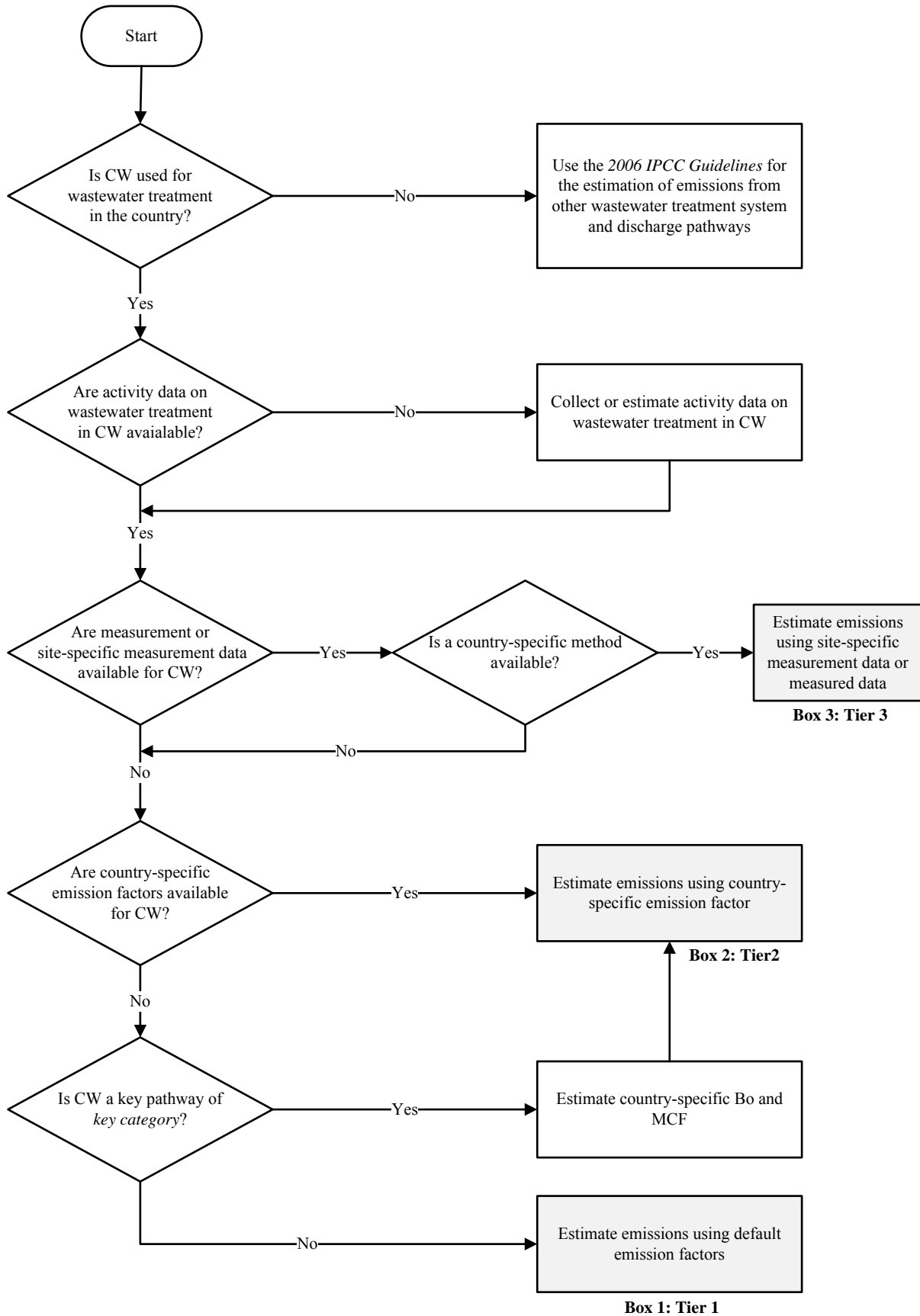
$$CH_4 Emissions = \sum_j (TOW_j \cdot EF_j) + \sum_{i,j} (TOW_{i,j} \cdot EF_j)$$

247 Where:

248 CH<sub>4</sub> emissions = CH<sub>4</sub> emissions in inventory year, kg CH<sub>4</sub>/yr249 TOW<sub>j</sub> = total organics in wastewater entering CW in inventory year, kg BOD/yr or kg  
250 COD/yr251 EF<sub>j</sub> = emission factor, kg CH<sub>4</sub>/kg BOD (for domestic wastewater only) or kg  
252 CH<sub>4</sub>/kg COD (for both domestic and industrial wastewater)253 If more than one type of CW is used in an industrial sector this factor would  
254 need to be a TOW<sub>i,j</sub>-weighted average.255 *i* = industrial sector256 *j* = type of CW

257

258 **Figure 6.3 Decision tree for CH<sub>4</sub> emissions from constructed wetlands**



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### 261 6.2.1.2 CHOICE OF EMISSION FACTORS

262 The emission factor for wastewater treatment using CWs is a function of maximum CH<sub>4</sub> producing potential (B<sub>o</sub>)  
263 and the methane correction factor (MCF).  
264

265 **EQUATION 6.2**  
266 **CH<sub>4</sub> EMISSION FACTOR FOR CONSTRUCTED WETLANDS**  
$$EF_j = B_o \cdot MCF_j$$

267 Where:

- 268 EF<sub>j</sub> = emission factor, kg CH<sub>4</sub>/kg BOD or kg CH<sub>4</sub>/ kg COD  
269 j = type of CWs  
270 B<sub>o</sub> = maximum CH<sub>4</sub> producing capacity, kg CH<sub>4</sub>/kg BOD or kg CH<sub>4</sub>/ kg COD  
271 MCF<sub>j</sub> = methane correction factor (fraction), See Table 6.4

272  
273 *Good practice* is to use country-specific data for B<sub>o</sub>, where available, expressed in terms of kg CH<sub>4</sub>/kg BOD  
274 removed for domestic wastewater or kg CH<sub>4</sub>/kg COD removed for industrial wastewater to be consistent with  
275 the activity data. If country-specific data are not available, the following default values can be used.  
276

277 The *2006 IPCC Guidelines* provide default B<sub>o</sub> values for domestic and industrial wastewater: 0.6 kg CH<sub>4</sub>/kg  
278 BOD and 0.25 kg CH<sub>4</sub>/kg COD.  
279

280 The MCF indicates the extent to which B<sub>o</sub> is realized in each type of CWs. It is an indication of the degree to  
281 which the system is anaerobic. The proposed MCFs for SF, HSSF and VSSF are provided in Table 6.4 and  
282 derived from literature-based analysis of CH<sub>4</sub> conversion rates. Each MCF in Table 6.4 is calculated from the  
283 relation of initial TOC loading to CH<sub>4</sub> emission flux derived from references provided in Table 6.2.  
284

CW type	MCF
Surface flow (SF)	0.4
Horizontal subsurface flow (HSSF)	0.1
Vertical subsurface flow (VSSF)	0.01

285  
286 These MCF values are derived based on actual measurement data and thus the operating and environmental  
287 conditions such as vegetation types and temperature effect have been taken into account. Based on the reported  
288 scientific data, there was insufficient information to distinguish the MCF values by vegetation types and  
289 operating temperatures. Nevertheless, these influencing factors can be considered for the estimation using higher  
290 tier approach. There was insufficient actual measurement data of hybrid systems to derive default MCF values. If  
291 the area fractions of SF, VSSF and HSSF for hybrid systems can be determined, the MCF values of the hybrid  
292 systems can be estimated as the area-weighted average of the MCFs for SF, VSSF and HSSF. Most commonly,  
293 SNTWs are surface flow type (Kadlec and Wallace, 2008), therefore, the default MCF of 0.4 can be used. If the  
294 type of CW cannot be recognized, the MCF of surface flow can be used in order to be conservative. Otherwise  
295 country-specific data should be used in higher tier method.

### 296 6.2.1.3 CHOICE OF ACTIVITY DATA

297 The activity data for this source category is the amount of organic materials (TOW) in the wastewater treated by  
298 CW. This parameter is a function of the population served by the CW system, and the biochemical oxygen  
299 demand (BOD) generation per person per day. BOD default values for selected countries are provided in the  
300 *2006 IPCC Guidelines* (Table 6.4, Chapter 6 of Volume 5 of the *2006 IPCC Guidelines*). In the case of industrial  
301 wastewater, COD loading to the CW system per day (kg COD/day) can be used. Examples of industrial  
302 wastewater data from various industries are provided in Table 6.9, Chapter 6, Volume 5 of the *2006 IPCC*  
303 *Guidelines*.

304 If industrial wastewater is released into domestic sewers, it is estimated together with domestic wastewater.  
 305 The equations for TOW are:  
 306

307 **EQUATION 6.3**  
 308 **TOTAL ORGANICALLY DEGRADABLE MATERIAL IN DOMESTIC WASTEWATER**  

$$TOW_j = P_j \cdot BOD \cdot I \cdot 0.001 \cdot 365$$

309  
 310

311 **EQUATION 6.4**  
 312 **TOTAL ORGANICALLY DEGRADABLE MATERIAL IN INDUSTRIAL WASTEWATER**  

$$TOW_{i,j} = COD_i \cdot W_{i,j} \cdot 365$$

313 Where:

- 314  $TOW_j$  = total organics in domestic wastewater treated in the CW in inventory year (kg  
 315 BOD/year)
- 316  $TOW_{i,j}$  = total organics in wastewater from industry  $i$  treated in the CW in inventory year (kg  
 317 COD/year)
- 318  $i$  = industrial sector
- 319  $P_j$  = population whose wastewater treated in CW\*
- 320 BOD = per capita BOD generation in inventory year (g BOD/person/day)
- 321  $I$  = correction factor for additional industrial BOD discharged into sewers (for  
 322 collected the default is 1.25, for uncollected the default is 1.00 as given in the 2006  
 323 *IPCC Guidelines*)
- 324  $COD_i$  = COD concentration in wastewater from industry  $i$  entering CW in the inventory year  
 325 (kg COD/m<sup>3</sup>)
- 326  $W_{i,j}$  = daily flow rate of industrial wastewater treated by CW, m<sup>3</sup>/day

327 \* Population should be subtracted from total population used in an Equation 6.3 in Chapter 6, Volume 5 in the 2006 *IPCC*  
 328 *Guidelines* to avoid double-counting.

## 329 6.2.2 Time series consistency

330 The same method and data sets should be used for estimating CH<sub>4</sub> emissions from CWs treating wastewater for  
 331 each year. The MCF for different treatment systems should not change from year to year, unless such a change is  
 332 justifiable and documented. If the share of wastewater treated in different treatment systems changes over the  
 333 time period, the reasons for these changes should be documented.  
 334

335 For activity data that are derived from population data, countries must determine the fraction of the population  
 336 served by CW systems. If data on the share of wastewater treated are missing for one or more years, the splicing  
 337 techniques such as surrogate data and extrapolation/interpolation described in Chapter 5, Time Series  
 338 Consistency, Volume 1 of the 2006 *IPCC Guidelines* can be used to estimate emissions. Emissions from  
 339 wastewater treated in CWs typically do not fluctuate significantly from year to year.

## 340 6.2.3 Uncertainties

341 Chapter 3 in Volume 1 of the 2006 *IPCC Guidelines* provides guidance on quantifying uncertainties in practice.  
 342 It includes guidance on eliciting and using expert judgments which in combination with empirical data can  
 343 provide overall uncertainty estimates. Table 6.5 provides default uncertainty ranges for emission factors and  
 344 activity data for domestic and industrial wastewater. The following parameters are believed to be very uncertain:

- 345 • The quantity of wastewater that is treated in CWs or SNTWs.
- 346 • The fraction of organics that is converted anaerobically to CH<sub>4</sub> during wastewater collection. This will  
 347 depend on hydraulic retention time and temperature in the wastewater collection pipeline, and on other

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- 348 factors including the presence of anaerobic condition in the wastewater collection pipeline and possibly  
349 components that are toxic to anaerobic bacteria in some industrial wastewater.
- 350 • The amount of industrial TOW from small or medium-scale industries and rural domestic wastewater that is  
351 discharged into CWs in developing countries.
  - 352 • Different plant species applied in CWs that are involved in gas exchange.

353

Parameter	Uncertainty range*
<b>Emission factor</b>	
Maximum CH <sub>4</sub> producing capacity (B <sub>0</sub> )	± 30%
Methane correction factor (MCF)	SF: ± 79% HSSF: ± 31% VSSF: ± 56%
<b>Activity data</b>	
Human population	± 5%
BOD per person	± 30%
Correction factor for additional industrial BOD discharged into sewers (I)	For uncollected, the uncertainty is zero %. For collected the uncertainty is ± 20%
COD loading from industrial wastewater	-55%, +103%

- 354 \* Uncertainty of MCF calculated as 95% confidence interval is shown in Table 1 in Annex. Uncertainty of COD loading  
355 from industrial wastewater is calculated based on Table 6.10 in Chapter 6 in Volume 5 of the *2006 IPCC Guidelines*.  
356 Others are the same to Tables 6.7 in Chapter 6 in Volume 5 of the *2006 IPCC Guidelines*.

## 357 6.2.4 QA/QC, Completeness and Reporting

358 It is *good practice* to conduct quality control (QC) checks and quality assurance (QA) procedures as outlined in  
359 Chapter 6, QA/QC and Verification, Volume 1 of the *2006 IPCC Guidelines*. Some fundamental QA/QC  
360 procedures include:

### 361 *Activity Data*

- 362 • Make sure that the sum of wastewater flows of all types of wastewater treatment processes including CWs  
363 equal 100 percent of wastewater collected and treated in the country.
- 364 • Inventory compilers should compare country-specific data on BOD in domestic wastewater to IPCC default  
365 values. If inventory compilers use country-specific values they should provide documented justification why  
366 their country-specific values are more appropriate for their national circumstances.

### 368 *Emission Factors*

- 369 • For domestic wastewater, inventory compilers can compare country-specific values for B<sub>0</sub> with the IPCC  
370 default value (0.25 kg CH<sub>4</sub>/kg COD or 0.6 kg CH<sub>4</sub>/kg BOD). As there are no IPCC default values for the  
371 fraction of wastewater treated anaerobically, inventory compilers are encouraged to compare values for  
372 MCFs against those from other countries with similar wastewater handling practices.
- 373 • Inventory compilers should confirm the agreement between the units used for organically degradable  
374 material in wastewater (TOW) with the units for B<sub>0</sub>. Both parameters should be based on the same units  
375 (either BOD or COD) in order to calculate emissions. This same consideration should be taken into account  
376 when comparing the emissions.
- 377 • For countries that use country-specific parameters or higher-tier methods, inventory compilers should  
378 crosscheck the national estimates with emissions estimated using the IPCC default method and parameters.
- 379 • For industrial wastewater, inventory compilers should cross-check values for MCFs against those from other  
380 national inventories with similar CW types.

## 382 **COMPLETENESS**

383 Completeness can be verified on the basis of the degree of utilization of a treatment or discharge system or  
384 pathway (T) for all wastewater treatment system used. The sum of T should equal 100 percent. It is a *good*  
385 *practice* to draw a diagram for the country to consider all potential anaerobic treatment and discharge systems  
386 and pathways, including collected and uncollected, as well as treated and untreated. Constructed wetlands and  
387 SNTWs are under treated and collected pathway. In general, the amount of vegetation harvested from CWs is  
388 very small. If vegetation biomass is removed for the purpose of composting, incineration and burning, disposal  
389 in landfills or as fertilizer on agricultural lands, the amount of biomass should be consistent with data used in the  
390 relevant sectors.

391 Completeness for estimating emissions from industrial wastewater depends on an accurate characterization of  
392 industrial sectors that produce organic wastewater and the organic loading applied to CW systems. So inventory  
393 compilers should ensure that these sectors are covered. Periodically, the inventory compilers should re-survey  
394 industrial sources, particularly if some industries are growing rapidly. This category should only cover industrial  
395 wastewater treated onsite. Emissions from industrial wastewater released into domestic sewer systems should be  
396 addressed and included with domestic wastewater.

## 397 **REPORTING**

398 Methane emission from CWs for wastewater treatment is reported in waste sector under the categories of  
399 domestic or industrial wastewater. Methane emission from CWs treating collected runoff from agricultural land  
400 is to be reported under the category of industrial wastewater.

# 401 **6.3 NITROUS OXIDE EMISSIONS FROM** 402 **CONSTRUCTED WETLANDS**

## 403 **6.3.1 Methodological issues**

404 Nitrous oxide (N<sub>2</sub>O) emissions can occur as direct emissions from wastewater treatment in CWs through  
405 nitrification and denitrification. Emissions are calculated based on the total nitrogen loaded into CWs and  
406 emission factor.

407 Three tier methods for N<sub>2</sub>O from this category are summarized below.

408 The Tier 1 method applies default values for the emission factor and activity parameters. This method is  
409 considered *good practice* for countries with no country-specific data.

410 The Tier 2 method follows the same method as Tier 1 but allows for incorporation of country-specific emission  
411 factors and country-specific activity data.

412 The Tier 3 method is used by countries with good data and advanced methodologies. A more advanced country-  
413 specific method is based on treatment system-specific data such as plant species and composition of wastewater.

414 The methodology provided assumes typical vegetation harvesting practices. However, the amount of vegetation  
415 harvested from CWs (studied until now) is generally very small and the harvested plant biomass is commonly  
416 not recorded so the harvesting practice is not considered as an influencing factor in the estimation of emissions.

417 Emissions from SNTWs treating uncollected wastewater are estimated using the same methodology. Indirect  
418 N<sub>2</sub>O emissions from domestic wastewater treatment effluent that is discharged into aquatic environments has  
419 already been covered in the *2006 IPCC Guidelines*.

### 420 **6.3.1.1 CHOICE OF METHOD**

421 A decision tree for domestic or industrial wastewater is shown in the Figure 6.4.

422 The general equation to estimate N<sub>2</sub>O emissions from CWs treating domestic or industrial wastewater is shown  
423 in Equation 6.5.

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426

**EQUATION 6.5**  
**N<sub>2</sub>O EMISSIONS FROM CONSTRUCTED WETLANDS**

$$N_2O \text{ Emissions} = \sum_j (N_j \cdot EF_j \cdot 44/28) + \sum_{i,j} (N_{i,j} \cdot EF_j \cdot 44/28)$$

427  
428

Where:

N<sub>2</sub>O emissions = N<sub>2</sub>O emissions in inventory year, kg N<sub>2</sub>O/yr429  
430N<sub>j</sub> = total nitrogen in domestic wastewater entering CWs in the inventory year, kg N/year431  
432N<sub>i,j</sub> = total nitrogen in industrial wastewater entering CW in the inventory year, kg N/year

433

EF<sub>j</sub> = emission factor, kg N<sub>2</sub>O-N/kg N434  
435If more than one type of CW is used in an industrial sector this factor would need to be a N<sub>i,j</sub>-weighted average.

436

*i* = industrial sector

437

*j* = type of CWs

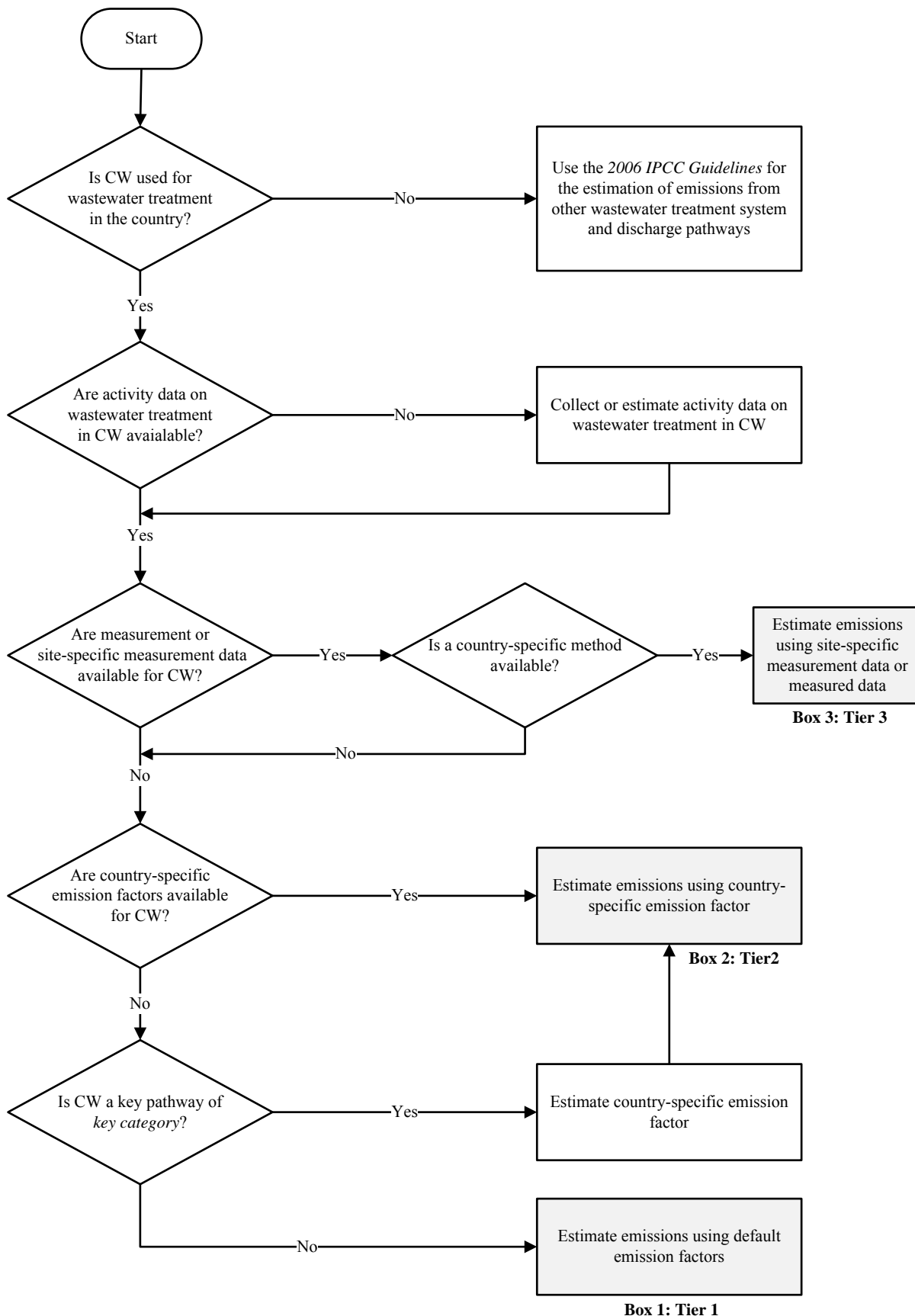
438

The factor 44/28 is the conversion of kg N<sub>2</sub>O-N into kg N<sub>2</sub>O.

439



440 **Figure 6.4** Decision tree for N<sub>2</sub>O emission from constructed wetland



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### 442 6.3.1.2 CHOICE OF EMISSION FACTORS

443 The default emission factors for N<sub>2</sub>O emitted from domestic and industrial wastewater treated by CWs are  
 444 0.0013 kg N<sub>2</sub>O-N/kg N for SF, 0.0079 kgN<sub>2</sub>O-N/kg N for HSSF and 0.00023 kgN<sub>2</sub>O-N/kg N for VSSF. These  
 445 values are based on data provided in the literatures and influenced by the extent of nitrification and  
 446 denitrification taking place in CWs, the coverage of vegetation in CWs and climatic conditions. There was  
 447 insufficient actual measurement data of hybrid systems to derive emission factors. If the area fractions of SF,  
 448 VSSF and HSSF for hybrid systems can be determined, the emission factors of the hybrid systems can be  
 449 estimated as the area-weighted average of the emission factors for SF, VSSF and HSSF CWs. *Good practice* is  
 450 to use country-specific data for emission factor, where available, expressed in term of kg N<sub>2</sub>O-N/kg N loaded for  
 451 domestic and industrial wastewater to be consistent with the activity data. The amount of N associated with N<sub>2</sub>O  
 452 emissions from CWs must be back calculated and subtracted from the N<sub>EFFLUENT</sub> (Equation 6.7 in Chapter 6,  
 453 Volume 5 of the *2006 IPCC Guidelines*).

### 454 6.3.1.3 CHOICE OF ACTIVITY DATA

455 The activity data for this source category are the amount of nitrogen in the wastewater entering CWs (TN). This  
 456 parameter is a function of the population served by the CW system, annual per capita protein consumption  
 457 (protein) and a factor for non-consumed nitrogen added to the wastewater for domestic wastewater. In case of  
 458 industrial wastewater, TN loading to the constructed wetland system in the inventory year (kg N) can be used  
 459 directly. The equations for determining TN for domestic and industrial wastewater are:

460

461  
462

$$\begin{aligned} & \text{EQUATION 6.6} \\ & \text{TOTAL NITROGEN IN DOMESTIC WASTEWATER} \\ & N_j = P_j \cdot \text{Protein} \cdot F_{NPR} \cdot F_{NON-CON} \cdot F_{IND-COM} \end{aligned}$$

463

464  
465

$$\begin{aligned} & \text{EQUATION 6.7} \\ & \text{TOTAL NITROGEN IN INDUSTRIAL WASTEWATER} \\ & N_{i,j} = TN_i \cdot W_{i,j} \end{aligned}$$

466 Where:

- 467  $N_j$  = total nitrogen in domestic wastewater entering CW in inventory year (kg N/year)
- 468  $N_i$  = total nitrogen in wastewater from industry  $i$  entering CW in inventory year (kg N/year)
- 469  $i$  = industrial sector
- 470  $P_j$  = human population whose wastewater entering CWs
- 471 Protein = annual per capita protein consumption, kg/person/yr
- 472  $F_{NPR}$  = fraction of nitrogen in protein (default is 0.16 kg N/ kg protein as given in the *2006*  
 473 *IPCC Guidelines*)
- 474  $F_{NON-CON}$  = factor for non-consumed nitrogen added to the wastewater (default is 1.1 for countries  
 475 with no garbage disposals, 1.4 for countries with garbage disposals as given in the  
 476 *2006 IPCC Guidelines*)
- 477  $F_{IND-COM}$  = factor for industrial and commercial co-discharged protein into sewer system (default  
 478 is 1.25 as given in *2006 IPCC Guidelines*)
- 479  $TN_i$  = total nitrogen concentration in wastewater from industry  $i$  entering CWs in inventory  
 480 year (kg N/m<sup>3</sup>)
- 481  $W_{i,j}$  = flow rate of industrial wastewater entering CW, m<sup>3</sup>/yr

482

483  $N_i$  is a function of total N concentration and flow rate which can be estimated by multiplying industrial product P  
 484 (tons/yr), wastewater generation (m<sup>3</sup>/ton) (Table 6.9, Chapter 6, Volume 5 in *2006 IPCC Guidelines*) and N  
 485 content in Table 6.6 of this supplement.

**TABLE 6.6**  
**EXAMPLE OF N CONTENT IN SOME NITROGEN-RICH INDUSTRIAL WASTEWATER**

Industry type	Wastewater generation W (m <sup>3</sup> /ton)	N content (kg/m <sup>3</sup> )
Alcohol refining	24 (16-32) <sup>1</sup>	2.40 (0.94-3.86) <sup>2</sup>
Fish processing industry	5 (2-8) <sup>2</sup>	0.60 (0.21-0.98) <sup>3</sup>
Seasoning source industry	NA	0.60 (0.22-1.00) <sup>3</sup>
Meat & poultry	13 (8-18) <sup>1</sup>	0.19 (0.17-0.20) <sup>3</sup>
Starch production	9 (4-18) <sup>1</sup>	0.90 (0.80-1.10) <sup>4</sup>
Nitrogen fertilizer plant	2.89 (0.46-8.3) <sup>2</sup>	0.50 (0.10-0.80) <sup>2</sup>
Landfill leachate	15-20% of annual precipitation in well compacted landfill site. 25-50% of annual precipitation for not well compacted landfill site <sup>6</sup> .	0.74 (0.01-2.50) <sup>5</sup>

Note: Average value and range (in brackets) are presented

Sources: <sup>1</sup> IPCC 2006; <sup>2</sup> Samokhin (1986); <sup>3</sup> Pilot Plant Development and Training Institute (1994); <sup>4</sup> Hulle *et al.* (2010); <sup>5</sup> Kjeldsen *et al.* (2002); <sup>6</sup> Ehrig (1983)

### 486 6.3.2 Time series consistency

487 The same method and data sets should be used for estimating N<sub>2</sub>O emissions from CWs for each year. If a  
488 country decides to change the estimation method from the default methodology (Tier 1) to country-specific (Tier  
489 2), this change must be made for the entire time series.

### 490 6.3.3 Uncertainties

491 Large uncertainties are associated with the default emission factors for N<sub>2</sub>O emissions from CWs due to limited  
492 available data (Table 6.7).

493

**TABLE 6.7**  
**NITROUS OXIDE METHODOLOGY DEFAULT UNCERTAINTIES**

Parameter	Default value	Range
<b>Emission factor</b> (kg N <sub>2</sub> O-N/kg N)	0.0013 for SF 0.0079 for HSSF 0.00023 for VSSF	± 90% for SF ± 79% for HSSF ± 70% for VSSF
<b>Activity data</b>		
Human population	Country-specific	± 10%
Annual per capita protein consumption	Country-specific	± 10%
Fraction of nitrogen in protein	0.16	0.15-0.17
Factor for non-consumed nitrogen	1.1 for countries with no garbage disposals, 1.4 for countries with garbage disposals	1.0-1.5
TN loading from industrial wastewater	Country-specific	-55%, +103%

494 \* Uncertainties of emission factors calculated as 95% confidence interval is shown in Table 6A1.1 in Annex. Uncertainty of  
495 TN loading from industrial wastewater is the same to that of COD loading from industrial wastewater (Expert judgement  
496 by Authors of this chapter). Others are derived from Tables 6.11 in Chapter 6 in Volume 5 of the 2006 IPCC Guidelines.

### 497 6.3.4 QA/QC, Completeness and Reporting

498 This method makes use of several default parameters. It is recommended to solicit experts' advice in evaluating  
499 the appropriateness of the proposed default factors. The methodology for estimating emissions is based on N  
500 associated with domestic and industrial discharge either collected into the collection system and treated in CWs

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501 or uncollected and discharged into SNTWs. This estimate can be seen as conservative and covers the entire  
502 source associated with domestic and industrial wastewater discharge.

503 **REPORTING**

504 Nitrous oxide emission from CWs for wastewater treatment is reported in waste sector under the categories of  
505 domestic or industrial wastewater. Nitrous oxide emissions from CWs treating collected runoff from agricultural  
506 land and landfill leachate are to be reported under the category of industrial wastewater. If agricultural runoff is  
507 collected and treated by CWs or SNTWs, the amount of nitrogen flows into CWs/SNTWs must be subtracted to  
508 avoid double counting.

509

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Final Draft

## 646 **Annex 6A.1 Estimation of default emission factors for CH<sub>4</sub> and** 647 **N<sub>2</sub>O in constructed wetlands for wastewater treatment**

648 We reviewed about 150 papers published in international peer-reviewed journals indexed by the Thomson  
 649 Reuters Web of Knowledge from 1994 to 2013. The terms “free water surface”, “surface flow”, constructed  
 650 wetland(s)”, “artificial wetland(s)”, “treatment wetland(s)”, “subsurface flow wetland(s)”, “vertical flow” and  
 651 “horizontal flow” in combination with the terms “carbon dioxide”, “CO<sub>2</sub>”, “methane”, “CH<sub>4</sub>”, “nitrous oxide”  
 652 and “N<sub>2</sub>O” were searched.

653 We found a total of 14 publications that provided information on emissions of either CH<sub>4</sub>, N<sub>2</sub>O or both gases in  
 654 surface flow (SF) constructed wetlands (CWs). These publications presented information on 17 different SF CW  
 655 systems, whereas for CH<sub>4</sub> and N<sub>2</sub>O, there were 24 and 25 subsystems/measuring events respectively. Six SF  
 656 CWs (Nykvärn, Lakeus, Ruka, Skjønhaug, Hässleholm, and Ibaraki) treated domestic wastewater (Johansson et  
 657 al., 2003, 2004; Søvik et al., 2006, Ström et al., 2006; Gui et al., 2007; Søvik and Kløve, 2007; Liu et al., 2009),  
 658 six CWs (mesocosms in Xue et al. (1999) paper, Donaumoos, Genarp, Görarp, Ormastorp, and Hovi) treated  
 659 waters of agricultural non-point pollution (Xue et al., 1999; Wild et al., 2001; Stadmark and Leonardson, 2005;  
 660 Søvik et al., 2006), two systems (Ngatea and Truro) were used for dairy farm wastewater treatment (Tanner et al.,  
 661 1997; Van der Zaag et al., 2010), the Kompsasuo CW treated wastewater from a peat extraction area (Søvik et al.,  
 662 2006), the Jiaonan CW (Tai et al., 2002) purified raw municipal wastewater, and synthetic wastewater is used in  
 663 the Jinan laboratory mesocosms (Wu et al., 2009).

664 Regarding the vertical subsurface flow (VSSF) CWs, there were only 4 measurement periods presented for 3  
 665 CWs from which CH<sub>4</sub> emission data and ratios could be calculated: Kõo in Estonia (Teiter and Mander 2005;  
 666 Søvik et al., 2006), Ski in Norway (Søvik et al., 2006), and Miho/Ibaraki, Japan (Gui et al., 2007; Liu et al.,  
 667 2009). For N<sub>2</sub>O emission, additionally laboratory microcosm experiments with different plant species from  
 668 Ibaraki, Japan (Inamori et al., 2008; Wang et al., 2008) were included.

669 For CH<sub>4</sub> fluxes from horizontal subsurface flow (HSSF) CWs we could use data from two system in Estonia  
 670 treating domestic wastewater, Kodijärve and Kõo (Mander et al., 2003, 2008; Teiter and Mander, 2005; Søvik et  
 671 al., 2006), four CWs treating domestic wastewater in Ski, Norway (Søvik et al., 2006), Barcelona, Spain (Garcia  
 672 et al., 2007), Miho/Ibaraki, Japan (Gui et al., 2007; Liu et al., 2007) and Slavosovice, Czech Republic (Picek et  
 673 al., 2007), a HSSF treating wastewater from a peat extraction area in Kompsasuo, Finland (Liikanen et al., 2006),  
 674 a HSSF treating landfill leachate in Bangkok, Thailand (Chiemchaisri et al., 2009), and a dairy farm wastewater  
 675 treatment HSSF in Truro, Nova Scotia, Canada (Van der Zaag et al., 2010). For N<sub>2</sub>O emissions from HSSFs,  
 676 also a CW for dairy farm wastewater treatment in Friedelhausen, Germany (Fey et al., 1998) has been included.

677 Tanner et al., (1997) presented estimated values for inflow total organic carbon (TOC<sub>in</sub>), Xue et al., (1999) for  
 678 inflow total nitrogen (TN<sub>in</sub>), and Søvik et al., (2006) for both TOC<sub>in</sub> and TN<sub>in</sub>. For most of the systems, TOC<sub>in</sub>  
 679 and TN<sub>in</sub> values were calculated based on area, hydraulic load and inflow TOC and TN concentration data. For  
 680 some systems only biological oxygen demand (BOD) values were usable, and for them the following  
 681 approximation based on domestic wastewater data was used: TOC = 0.5 BOD (Garcia et al., 2007). For the  
 682 calculations of emission factors, we used data series from one year or at least a vegetation period.

	Emission factor CH <sub>4</sub> -C/TOC (%)					Emission factor N <sub>2</sub> O-N/TN (%)				
	Average	Standard Error	Median	2.5%	97.5%	Average	Standard Error	Median	2.5%	97.5%
SF	42.2	20.4	18	4	446	0.13	0.024	0.11	0	0.47
HSSF	12.0	7.56	4.15	0.03	79	0.79	0.38	0.34	0.04	3.01
VSSF	1.17	0.33	1.28	0.38	1.73	0.023	0.005	0.018	0.001	0.096

683

684 Table 1 presents values of emission factors calculated based on literature sources described above.

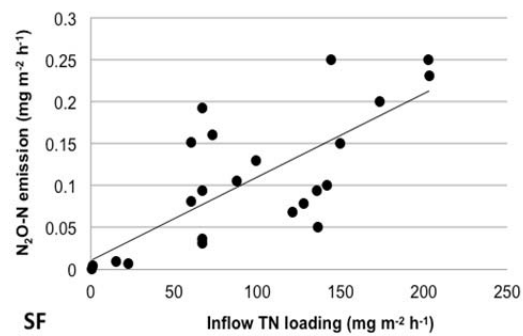
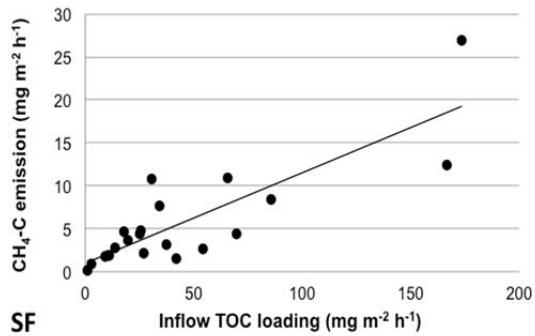
685 In Figure 1, correlation between the inflow TOC loading and CH<sub>4</sub>-C emission and between the inflow TN  
 686 loading and N<sub>2</sub>O emission in SF, HSSF and VSSF CWs is presented.

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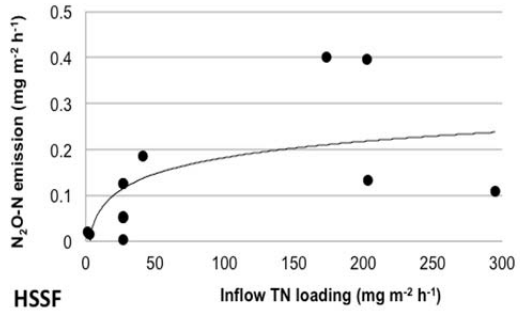
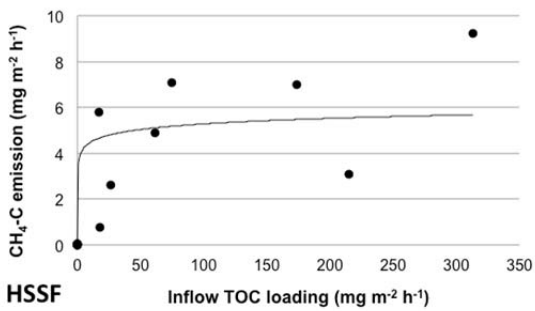


688 **Figure 6A1.1** The relationship between inflow TOC loading and CH<sub>4</sub>-C emission (left  
 689 column) and between inflow TN loading and N<sub>2</sub>O-N emission (right columns)  
 690 in SF, HSSF, and VSSF CWs. In all cases,  $p < 0.05$ .

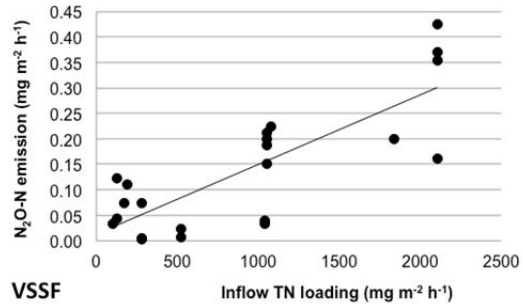
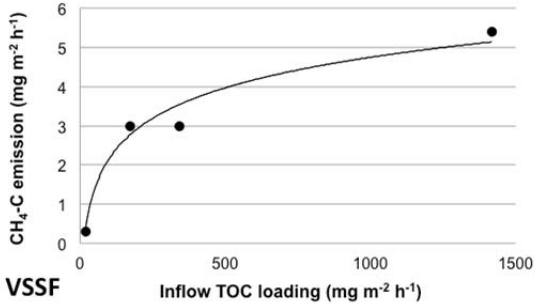
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