

1 **Supplementary material to:**
2 **Global anthropogenic methane emissions 2005-2030:**
3 **technical mitigation potentials and costs**

4 **Detailed descriptions of estimations by sector**

5

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9

10 **1 Introduction**

11 This is a description by sector of the estimations of global anthropogenic emissions of
12 methane (CH₄) presented in the paper “Global anthropogenic methane emissions
13 2005-2030: technical mitigation potentials and costs”. It provides further insights into
14 the details of the estimations of emissions, mitigation potentials and associated costs
15 as well as a discussion of the most important sources for uncertainty in the sector
16 estimates.

17

18 **2 CH₄ emission estimations by sector**

19 **2.1 Crude oil and natural gas production**

20 Extraction of crude oil and natural gas gives rise to CH₄ emissions, partly as a result
21 of intended flaring or venting of associated gas for security reasons and partly due to
22 unintended leakage of fugitive emissions, which occur along the whole production
23 process from well head to upgrading and storage (IPCC, 2006, Vol.2, Section 4.2).
24 Associated gas is a gas compound mainly consisting of CH₄, which is released as oil
25 or natural gas is pumped to the surface. For security reasons, the associated gas needs
26 to be released and is therefore flared off or simply vented. Alternatively, the

1 associated gas can be recovered and utilized for energy purposes provided there is an
 2 infrastructure present to transport the recovered gas to consumers.

3 Emissions from venting and flaring of associated gas are calculated separately for
 4 fugitive emissions and unintended leakage. Total emissions from oil and gas
 5 production are the sum of venting, flaring and leakage emissions:

$$6 \quad E_{its} = E_{its}^{venting} + E_{its}^{flaring} + E_{its}^{leakage} \quad (1)$$

7 Venting emissions from production of oil and gas, respectively, are calculated as:

$$8 \quad E_{it;oil}^{venting} = A_{it}^{oil} * 20 * (1 - r_i) * (c_i a_i^{conv} v_i^{conv} + (1 - c_i) a_i^{heavy} v_i^{heavy}) \quad (2)$$

$$9 \quad E_{it;gas}^{venting} = A_{it}^{gas} * 20 * a_i^{gas} (1 - r_i) v_i^{gas} \quad (3)$$

10
 11 where $A_{it}^{oil;gas}$ is energy content of marketable oil or gas produced in country i
 12 in year t ,

13 20 20 kt CH₄/PJ conversion of energy content to amount of CH₄,

14 c_i fraction of conventional oil (as opposed to heavy oil) produced,

15 a_i^{conv} , a_i^{heavy} , a_i^{gas} are associated gas as fractions of the total energy content of
 16 conventional, heavy oil, or gas produced in year 2005,

17 r_i is the fraction of associated gas recovered for utilization or
 18 reinjection in year 2005, and

19 v_i^{conv} , v_i^{heavy} , v_i^{gas} are fractions of unrecovered associated gas that is vented (as
 20 opposed to flared).

21

22 Amounts of associated gas flared are calculated as the residual when the sum of the
 23 amounts of associated gas recovered/reinjected (rec) and vented ($vented$) are
 24 subtracted from the total amount of associated gas generated ($total$), i.e.,

$$25 \quad E_{it}^{flaring} = 0.02 * (total - rec - vented) \quad (4)$$

26

27 Emissions are derived assuming two percent incomplete combustion of CH₄ from
 28 flares (Johnson and Kostiuik, 2002).

1 Country-specific information for year 2005 on amounts of associated gas and the
2 fraction of associated gas reinjected or recovered are taken from EIA (2010a)
3 International Energy Statistics and used as starting point for emission estimates. From
4 EIA (2010a) Country Analysis Briefs, types of hydrocarbon produced are identified,
5 i.e. conventional crude oil, heavy crude oil (API gravity <22.3°), oil sands and natural
6 gas, as well as the fraction of offshore production. For gas production only two
7 measurement results have been found for the fraction of associated gas to the energy
8 content of gas produced. These are 0.03 percent for Canada (Johnson and Coderre,
9 2011), which is applied to developed countries and 0.3 percent for Russia (PFC
10 Energy, 2007), which is applied to developing and transitional countries.

11 Measurement data published by Johnson and Coderre (2011) for Canadian oil and gas
12 production is used to derive default fractions for the amounts of associated gas vented
13 as opposed to flared for different types of hydrocarbons. Finally, the derived total
14 amount of associated gas flared from both oil and gas production in a country is
15 verified against country-specific satellite images of flares (NOAA, 2011). For a few
16 countries (Mexico, Denmark, China, Columbia) is it necessary to make slight
17 adjustments to the amounts of associated gas reported to EIA (2010a) or to the
18 reported amounts of gas recovered or reinjected in order to match satellite images of
19 flares reasonably well. A summary of the assumptions made for major oil and gas
20 producing countries is presented in Table 1.

21

1 Table 1: Assumptions for deriving CH₄ venting and flaring emission factors from oil and gas production in major source countries.

Country	Types of crude oil produced as fraction of total production (EIA, 2010a)		Fraction offshore production (EIA, 2010a)	Assumptions venting from oil production				Assumptions venting from gas production		Derived total estimates of associated gas for 2005			Satellite image flaring 2005 (NOAA, 2010)
	Conventional oil	Heavy oil ^a		Ass gas as % of crude oil produced ^b	Flaring or venting/ Recovery or reinjection	Venting from conventional oil prod	Venting from heavy oil prod	Ass gas as % of gas produced ^b	Venting as % ass gas flared or vented	Recov/Reinj	Flared	Vented	
				% of oil prod	% of ass gas	venting as % of ass gas flared or vented		% of gas production	% vented				bcm
Canada	40%	10% ^c	11%	14.5% ^{efg}	10/90 ^{efg}	29% ^f	88% ^f	0.03% ^f	40% ^f	26.6	2.1	1.0	1.3
Mexico	33%	67%	80%	6.7% ^j	25/75 ^j	29% ^f	88% ^f	0.3% ^h	40% ^f	13.4	1.9	2.7	1.9
USA	100%	0%	7%	35.9% ^e	3/97 ^e	29% ^f	88% ^f	0.03% ^f	40% ^f	164.9	3.9	1.6	2.8
Brazil	10%	90%	90%	8.1% ^e	45/55 ^e	29% ^f	88% ^f	0.3% ^h	40% ^f	6.2	2.0	3.2	1.6
Columbia	5%	95%	0%	22.3% ^e	35/65 ^j	29% ^f	88% ^f	0.3% ^h	40% ^f	5.8	0.5	2.6	0.5
Venezuela	100%	0%	0%	20.6% ^e	25/75 ^e	29% ^f	88% ^f	0.3% ^h	40% ^f	32.4	1.4	9.6	2.2
Denmark	100%	0%	100%	20% ^j	2/98 ^j	10% ^j	88% ^f	0.03% ^f	40% ^f	5.2	0.1	0.0	0.2
Norway	100%	0%	100%	25.7% ^e	1/99 ^e	29% ^f	88% ^f	0.03% ^f	40% ^f	51.6	0.6	0.2	0.5
Azerbaijan	100%	0%	80%	19.7% ^m	50/50 ^k	29% ^f	88% ^f	0.3% ^h	40% ^f	3.2	2.3	0.9	0.2
Kazakhstan	100%	0%	20%	19.7% ^m	50/50 ^k	29% ^f	88% ^f	0.3% ^h	40% ^f	9.6	6.8	2.8	6.2
Russia	100%	0%	0%	20% ^h	50/50 ^{hi}	29% ^f	88% ^f	0.3% ^h	40% ^f	68.7	50.1	21.0	58.3
Iran	100%	0%	33%	18.1% ^e	28/72 ^e	29% ^f	88% ^f	0.3% ^h	40% ^f	40.5	11.2	4.7	11.7
Iraq	100%	0%	0%	8.1% ^e	91/9 ^e	29% ^f	88% ^f	0.3% ^h	40% ^f	1.0	7.1	2.9	7.0
Kuwait	100%	0%	0%	8.1% ⁿ	70/30 ^j	29% ^f	88% ^f	0.3% ^h	40% ^f	4.7	2.8	8.2	2.3
Oman	100%	0%	0%	6.3% ^e	80/20 ^j	29% ^f	88% ^f	0.3% ^h	40% ^f	0.7	2.1	0.9	2.6
Qatar	100%	0%	0%	12.4% ^e	66/34 ^e	29% ^f	88% ^f	0.3% ^h	40% ^f	3.1	4.3	1.8	2.3
S. Arabia	85%	15%	20%	1.1% ^e	56/44 ^e	29% ^f	88% ^f	0.3% ^h	40% ^f	5.9	3.5	4.2	3.5
UAE	100%	0%	10%	12.2% ^e	5/95 ^e	29% ^f	88% ^f	0.3% ^h	40% ^f	23.2	1.0	0.5	0.9
Nigeria	100%	0%	20%	18.8% ^e	77/23 ^e	29% ^f	88% ^f	0.3% ^h	40% ^f	8.4	19.7	8.1	21.3
China	100%	0%	15%	16% ^q	10/90 ^j	29% ^f	88% ^f	0.3% ^h	40% ^f	38.0	3.1	1.3	3.0
Indonesia	100%	0%	0%	25.9% ^e	27/73 ^e	29% ^f	88% ^f	0.3% ^h	40% ^f	15.1	4.1	1.7	3.0
Malaysia	100%	0%	100%	40.6% ^e	11/89 ^e	29% ^f	88% ^f	0.3% ^h	40% ^f	18.5	1.7	0.7	1.8
Rest of World										61.5	37.8	18.9	37.1
World total										608	170	100	172

^a API gravity <22.3°; ^b Energy content of associated gas as fraction of PJ crude oil or gas produced (EIA, 2010a) or if data is missing defaults are 5% for heavy crude oil and 35% for conventional oil derived from Johnson and Coderre (2011). CH₄ content of associated gas assumed 86% on average (ERCB, 2010) and 35.9 MJ/m³ CH₄.; ^c In addition Canada produces 50% of oil from oil sands.; ^d Information derived from OME (2001) and from Table 4 on existing gas pipelines.; ^e Source: EIA (2011); ^f Source: derived from Johnson and Coderre (2011); ^g Source: ERCB(2010); ^h Source: derived from PFC Energy (2007); ⁱ Source: Hulbak Røland (2010); ^j Reported value adjusted by author to fit with satellite image of flaring (NOAA, 2011); ^k Assumed same as Russia; ^l Assumed same as Germany; ^m Assumed same as Kazakhstan; ⁿ Assumed same as Iraq; ^o Assumed same as Qatar; ^p Assumed same as N.Zeeland; ^q Assumed default for Asia when EIA (2010a) data missing; ^r Assumed default for Africa when EIA (2010a) data is missing.

1 Fugitive CH₄ emissions are usually unintended with irregular occurrence and
 2 therefore highly uncertain. In GAINS, fugitive emissions from oil and gas production
 3 are calculated as follows:

$$4 \quad E_{it;oil}^{leakage} = A_{it}^{oil} * (\gamma_i ef_{offshore}^{oil} + p_i ef_{oilsand} + (1 - \gamma_i - p_i) ((1 - c_i) ef_{onshore}^{heavy} + c_i ef_{onshore}^{conventional})) \quad (5)$$

$$6 \quad E_{it;gas}^{leakage} = A_{it}^{gas} * (\gamma_i ef_{offshore}^{gas} + (1 - \gamma_i) ef_{onshore}^{gas}) \quad (6)$$

7
 8 where A_{it} is the energy content of marketable oil (or gas)
 9 produced in country i in year t ,
 10 γ_i is the fraction of oil (or gas) produced offshore,
 11 $ef_{offshore}^{oil}$ is the IPCC default emission factor for offshore oil production,
 12 p_i is the fraction of oil produced from oilsands,
 13 $ef_{oilsand}$ is the IPCC default emission factor for oil production from
 14 oilsands,
 15 c_i is the fraction of conventional oil produced,
 16 $ef_{onshore}^{heavy}$ is the IPCC default emission factor for heavy oil production,
 17 $ef_{onshore}^{conv}$ is the IPCC default emission factor for conventional oil
 18 production.
 19 $ef_{offshore}^{gas}$ is the IPCC default emission factor for offshore gas production,
 20 $ef_{onshore}^{gas}$ is the IPCC default emission factor for onshore gas production.

21
 22 GAINS uses IPCC (2006, Vol.2, Tables 4.2.4 and 4.2.5) default emission factors as
 23 specified separately for developed countries and developing/transitional countries. For
 24 developed countries, the median of the emission factor range given by IPCC is used,
 25 while for developing/transitional countries the range is usually wide and therefore a
 26 general assumption is made about double factors compared with developed countries.
 27 Adopted fugitive emission factors and IPCC ranges for default emission factors are

1 presented in Table 2 by type of hydrocarbon produced and for on-shore/off-shore
 2 production.

3 Table 2: Default emission factors for unintended fugitive emissions from oil and gas
 4 production used in GAINS and in comparison with IPCC (2006, Vol.2, Tables 4.2.4
 5 and 4.2.5).

		Developed countries		Developing/transitional countries	
		GAINS	IPCC (2006) range	GAINS	IPCC (2006) range
		kt CH ₄ /PJ	kt CH ₄ /PJ	kt CH ₄ /PJ	kt CH ₄ /PJ
Oil production	Conventional oil on-shore	0.06	0.00004-0.094	0.12	0.00004-1.5
	Heavy oil on-shore	0.1863	0-0.3726	0.3726	0.1863-3.066
	Conventional and heavy oil off-shore	0.000015	0-0.00003	0.000015	0.000013-0.00013
	Oilsands	0.0542	0.0135-0.095	0.0542	0.018-0.135
Gas production	Natural gas on-shore	0.06	0-0.12	0.12	0.1-2.15
	Natural gas off-shore	0.00974	0-0.0195	0.00974	0.0058-0.034

6

7
 8 Maximum technically feasible reduction of CH₄ emissions from oil and gas
 9 production is defined such that all countries are assumed able to recover and utilize at
 10 least 95 percent of the associated gas. Extended recovery and utilization requires
 11 infrastructure capacity to transfer the gas from the oil and gas fields to the consumers.
 12 As existing pipelines are usually operated close to capacity maximum (EIA, 2010b),
 13 additional recovery and utilization of associated gas from major producing regions in
 14 the world requires new infrastructure capacity. OME (2001) has analysed the
 15 technical possibilities and costs of extending gas supply to the European market from
 16 different world regions. Supply costs include production, transportation and transit
 17 fees to the EU-15 border and are calculated both for extended gas pipeline capacity
 18 and LNG production capacity. In GAINS, transport and transit fee costs from the
 19 OME study are used with the exception of the costs for setting up gas processing
 20 plants (GPPs) and upgrading the gas. The investment cost of GPPs is set to 0.2
 21 M€/M³ associated gas throughput, which with a 86 percent CH₄ content of
 22 associated gas being upgraded to 97 percent, corresponds to a cost of about 3 M€/PJ
 23 gas throughput, which falls within the cost range 2.4-3.7 M€/PJ estimated by GGFR
 24 (2007) and Persson (2003).

25 There are several cost-effective and low cost options available for reducing
 26 unintended leakage during extraction of oil and natural gas. For an exhaustive list, see

1 USEPA (2011a). In GAINS, this option is defined as extending leakage control in
2 developing countries to the standard currently observed in developed countries. This
3 means cutting emissions in developing countries by 50 percent, i.e., from 0.6 to 0.3 PJ
4 gas per PJ conventional oil produced on-shore, and no further mitigation in developed
5 countries. Given the large number of cost-effective measures available (USEPA,
6 2011a), it is assumed that a 50 percent cut in developing countries is possible at zero
7 mitigation cost and current gas price levels. This corresponds to an investment cost of
8 6.5 M Euro/PJ gas or oil produced annually or an annualized cost of 0.5 MEuro/PJ
9 production when equipment lifetime is 20 years.

10 As noted by Buzcu-Guven et al. (2010), the data on flaring and venting volumes of
11 associated gas are highly speculative and “order of magnitude errors are within the
12 realm of possibility”. The uncertainty in estimates is both found in the total amount of
13 associated gas generated as well as in the fractions of the gas being recovered, flared
14 or vented. There is a general lack of measurement data to verify assumptions on these
15 fractions. The rare exceptions are the satellite images of flares from NOAA (2010)
16 and the detailed measurement data published by Johnson and Coderre (2011) on
17 amounts of associated gas flared or vented from different types of oil and gas wells in
18 Alberta, Canada. Applying the information from Canadian oil and gas sources to other
19 parts of the world is of course a crude approximation, however, when the derived
20 amounts of flaring are verified against the satellite flaring images there is a reasonably
21 close match for most countries (see Table 1). IPCC (2006, Vol.2, Tables 4.2.4 and
22 4.2.5) suggests an uncertainty range of ± 75 percent for Tier 1 default emission factors
23 for flaring and venting of associated gas. The estimates used here are, however, scaled
24 with a number of country-specific factors and should therefore be lower. When
25 varying the country-specific assumptions about the fractions of associated gas
26 recovered/reinjected or vented as opposed to flared, within reasonable ranges, the
27 resulting uncertainty range falls within ± 30 percent on a global scale.

28 The uncertainty range specified above assumes that reported country-specific data on
29 associated gas is reasonably accurate, in particular for large oil and gas producing
30 countries. If this is not the case, the uncertainty range is higher. As an example, the
31 amount of associated gas as fraction of crude oil produced reported to EIA (2010a) for
32 Saudi Arabia is about 1 percent in 2005. If this were incorrect and the country instead
33 flares at levels comparable with what its neighbours report, e.g., Iraq (8 percent),

1 Oman (6 percent), Qatar (12 percent), and UAE (12 percent), then Saudi Arabian
2 venting emissions would increase five to ten times, which translates into global
3 anthropogenic CH₄ emissions being four to eight percent higher in 2005.

4 IPCC (2006, Vol. 2, Tables 4.2.4 and 4.2.5) gives an uncertainty range for unintended
5 leakage from oil production to ± 100 percent for developed countries and -12.5 to
6 +800 percent for developing and transitional countries. Since GAINS weighs in
7 country-specific information about type of oil produced in emission estimates and the
8 emission factor for heavy oil is about three times higher than for conventional oil, the
9 uncertainty range is scaled down three times to ± 30 percent for developed countries
10 and -4 to +240 percent for developing and transitional countries. The corresponding
11 IPCC uncertainty ranges for unintended leakage from gas production are ± 100
12 percent for developed countries and -40 to +250 percent for developing and
13 transitional countries. These ranges are applied here.

14 A recent study by Howarth et al. (2011) suggests that CH₄ emissions from shale gas is
15 3.6 to 7.9 percent of the gas produced. This means emissions are at least ten times
16 higher per unit of gas produced than for natural gas production. As both leakage rates
17 as well as the extent of future shale gas extraction are very uncertain, a higher
18 emission factor for shale gas is not assumed in emission estimations. It is, however,
19 still possible to explore what it could mean for future emissions. In the US shale gas
20 production is expected to increase from 5 to 75 percent of total gas production
21 between 2005 and 2035 (Howarth et al., 2011). Using a higher emission factor of five
22 percent for shale gas adds about 1200 kt CH₄ in 2005 and 16641 kt CH₄ in 2030 only
23 for the US. Also Australia, Canada, China, India, Austria, Bulgaria, Germany,
24 Hungary, Ireland and Poland have expressed interests in expanding extraction of shale
25 gas. If these countries extract 10-50 percent of their expected natural gas production
26 in 2030 from shale gas then when added to the emissions from US shale gas
27 extraction this source would add about five percent to global anthropogenic CH₄
28 emissions.

29 **2.2 Crude oil transportation and refining**

30 CH₄ emissions from oil refinery and transport are fugitive emissions related to
31 evaporation losses from storage, filling and unloading activities as well as fugitive
32 leaks (IPCC, 2006, Vol.2, p. 4.34). The IPCC (2006, Vol.2, pp.4.52-4.61) guidelines

1 provide emission factors for oil transportation based on the amount of oil transported,
 2 while emission factors for refining and storage are based on the amount of oil refined
 3 (IEA-WEO, 2009). Since it is not possible to find systematic data on the amount of oil
 4 transported by tanker, trucks or rails by region, GAINS assumes that the amount
 5 transported corresponds to the amount of oil refined. Thus, to calculate emissions
 6 from this source the activity data used is amount of oil refined combined with IPCC
 7 default emission factors for oil refined and transported summed up:

$$8 \quad E_{it} = \sum_m [A_{it} * (ef^{refined} + ef^{transported}) * (1 - remeff_m) * Appl_{im}], \quad (7)$$

9 where A_{it} is amount of oil refined in country i in year t ,

10 $ef^{refined}$ is the IPCC default emission factor for oil refined,

11 $ef^{transported}$ is the IPCC default emission factor for oil transported,

12 $remeff_m$ is the removal efficiency of technology m , and

13 $Appl_{im}$ is the application rate of technology m to emissions in country i
 14 in year t .

15 IPCC default emission factors for this sector are presented in Table 3 together with
 16 GAINS assumptions for no control and controlled emission factors.

17 The maximum technically feasible reduction is defined by the sum of the lower range
 18 IPCC default emission factor for oil refined and a fifty percent reduction in leakage
 19 emissions from oil transportation. This corresponds to the relative reduction in
 20 leakage emissions considered technically feasible for oil and gas production sectors
 21 (see Section 2.1). In 2005, developed countries are assumed to have control installed
 22 corresponding to between 2/3 of the level considered technically feasible to control
 23 (UNFCCC, 2010). Costs for these measures are taken from AEAT (1998).

24

1 Table 3: Default emission factors for CH₄ emissions from oil refinery and transport. Source:
 2 Derived from IPCC (2006, Vol.2, pp.4.52-4.61)

Emission source	GAINS		IPCC (2006)	unit
	No control	Control		
Fugitive emissions at oil refinery	0.0455	0.0029	0.0029 to 0.0455	kt CH ₄ /Mt oil refined
Fugitive emissions from transport by pipeline	0.0049	0.00245	0.0049	kt CH ₄ /Mt oil transported by pipeline
Fugitive emissions from transport by tanker, truck and rail cars	0.0225	0.0166	0.0225	kt CH ₄ /Mt oil transported by tanker/truck
All sources	0.0729	0.0166	0.0303 to 0.0729	kt CH ₄ /Mt oil refined

3

4 **2.3 Natural gas transportation**

5 Loss of natural gas during long-distance transmission is an important source of CH₄
 6 emissions. IPCC guidelines (2006, Vol. 2, p.4.48-4.62) recommend for Tier 1
 7 estimations the use of default emission factors per million m³ of marketable gas. I find
 8 it problematic to use the IPCC default factors here as emissions from gas transmission
 9 are likely to be influenced by both the volume of gas transported and the distance the
 10 gas is transported. Instead of using the volume-based IPCC default factors, typical
 11 emission factors are derived for a number of countries for which detailed data is
 12 available on reported emissions, amounts of gas transported, and km long-distance on-
 13 shore gas pipelines. Emissions are measured per unit of a product of the two factors
 14 distance (in km) and volume (in bcm). Emissions are estimated as follows:

$$15 \quad E_{it} = \sum_m [l_{it} * v_{it} * ef_i * (1 - remeff_{im}) * Appl_{im}], \quad (8)$$

16 where l_{it} is the length of long-distance on-shore pipelines (in km) in
 17 country i in year t ,

18 v_{it} is the volume of gas transported (in bcm) in country i in year t ,

19 ef_i is the derived default emission factor in kg CH₄/(km*bcm),

1 rem_{im} is the removal efficiency of technology m , which is defined as
2 the quota between the maximum feasible reduction emission
3 factor and the current emission factor ef_i , and

4 $Appl_{im}$ is the application rate of technology m to emissions in country i
5 in year t .

6 The reason for choosing this particular specification including the chosen scaling of
7 the variables with distance in km and volume in bcm, rests on the observation that for
8 a few number of countries presented in Table 4 and for which there exists reported
9 data on emissions, volume transported and length of onshore gas pipelines, the
10 resulting derived leakage rates fit fairly well for comparable countries and also match
11 measurement results when such are available. E.g., the derived estimate of about 1
12 percent leakage from Russian gas pipelines is verified by measurement data from
13 Lelieveld et al., 2005 and Lechtenböhmer et al., 2007, who measure ranges of 0.4 to
14 1.6 percent and 0.5 to 1.5 percent for overall leakages from gas pipelines in Russia.
15 For the US, the derived leakage rate of 0.41 percent is comparable in magnitude to
16 0.66 percent measured by Kirschgessner et al., 1997 and 0.40 percent reported in
17 USEPA (2006a). Although an approach which regards both volume and distance
18 transported in the emission estimations can be viewed as an improvement over IPCC
19 Tier 1 emission factors, it must still be considered a crude approximation. E.g., most
20 likely the weights given to pipeline length and volume transported in emission
21 estimations differ between pipeline systems. If measurement data on emissions, on-
22 shore pipeline lengths and volume transported were available for a larger number of
23 countries or pipeline systems, it would have been possible and desirable to use
24 statistical methods to empirically determine the functional form for the relationship
25 between distance, volume transported and emissions. With current data availability
26 this is unfortunately not possible.

27 Data sources for pipeline length and amounts of gas transported are UNFCCC (2010)
28 CRF tables complemented with information from IEA (2010) and CIA World
29 Factbook (2010) as well as country-specific sources (SPP 2007; TAG 2007; EIA
30 2007; Wuppertal Institute 2005). The derived country-specific factors are presented in
31 Table 4. The factors show close values within groups of similar countries, i.e., a range
32 of 8.9 to 14.7 kg CH₄/(km*bcm) for Western developed countries (Austria, Australia,
33 Canada, Japan and the US) and a range of 37.7 to 97.2 kg CH₄/(km*bcm) for Eastern

1 European and Central Asian countries (Russia, Slovakia, Ukraine and Kazakhstan).
2 Averages of the emission factors derived for the reference countries are then applied
3 to other countries for which reported emissions are not as complete, see Table 4. For
4 Annex 1 countries, results are verified against reported emissions to UNFCCC (2010).
5 Unless mitigation options apply, current leakage rates remain constant in the future
6 and growth in future emissions is assumed proportional to growth in domestic gas
7 consumption.

8 Leakage of CH₄ emissions from long-distance gas pipelines arise for several reasons,
9 e.g., untight compressor seals and valves or because pipelines are flushed with gas
10 during start-ups. In the maximum technically feasible reduction case is assumed that
11 all countries are able to reduce emissions from gas pipelines to at least the level
12 currently observed in Western Europe, Japan and North America. On the basis of the
13 derived emission factors in developed countries, it is considered technically possible
14 to reduce emissions in all countries to 10 kg CH₄/(bcm*km). Costs are taken from
15 Ecofys (1998) and reflect costs for a set of measures including electrical start-up with
16 no flushing of natural gas during start-up and inspection and maintenance programs to
17 secure compressor seals and valves.

18 As mentioned, growth in future emissions assumes no further expansion in pipeline
19 lengths, but a future growth proportional to growth in transported volumes. This
20 assumption is rather crude as future emissions may not grow proportionately with
21 volumes transported but rather result from a combination of extended pipeline
22 capacity and transported volumes. Because of difficulties finding systematic
23 information about unutilized capacity in existing pipelines and the uncertainty
24 surrounding the likelihood that currently planned pipeline extensions will be realized,
25 the transportation capacity assumed reflects the capacity as of 2012 and remains
26 constant throughout future baseline emission estimates. For the uncertainty analysis,
27 the IPCC (2006, Vol.2, Tables 4.2.4 and 4.2.5) default range of ±30 percent for Tier 1
28 emission factors is adopted for this source.

1 Table 4: Derivation of reference emission factors for long-distance gas pipelines. Data
 2 sources: UNFCCC (2010), IEA (2010), EIA (2010b), Wuppertal Institute (2005), TAG
 3 (2007), SPP (2007), CIA World Factbook (2010).

Reference cases in Annex-1 countries	Length of onshore gas pipelines	Gas transported 2005	Energy content of gas transported 2005	Emissions reported for 2005 to UNFCCC (2009)	Derived factors		
					Emission factors		Leakage rate
					kg CH ₄ / (bcm*km)	kt CH ₄ / PJ transport	% of gas transport
	km	bcm	PJ	kt CH ₄			
Canada	83195	366	13980	270	8.9	0.019	0.10%
Japan	2720	33	1318	1	10.9	0.001	0.00%
USA	278089	589	22591	1862	11.4	0.082	0.41%
Austria	6290	33	1300	3	13.0	0.002	0.01%
Australia	27105	23	915	9	14.7	0.010	0.05%
Slovakia	2270	77	2919	7	37.7	0.002	0.01%
Russia	170000	622	23382	5048	47.7	0.216	1.08%
Kazakhstan	5808	128	4979	63	85.6	0.013	0.06%
Ukraine	37820	60	2349	221	97.2	0.094	0.47%
Assumed emission factors for other country groups	Non-Annex 1				39.0		
	EU-15				13.0		
	EU-10, Turkey, Croatia				37.7		
	N. Zealand				10.9		
	Former Soviet Union				76.9		
Max technically feasible reduction					10.0		
World^a	933993	2692	108374	7805	24.6	0.072	0.68%

4 ^aWorld sums when using production quantities reported to UNFCCC, which do not always correspond exactly to quantities in IEA-WEO 2009.

5

6 2.4 Gas distribution networks

7 CH₄ emissions from gas use come from leakage in consumer distribution networks
 8 and during end-use by consumers. The activity data is amount of gas consumed by
 9 sectors from IEA-WEO (2009) Reference scenario. IPCC (2006, Vol.2, Tables 4.2.4
 10 and 4.2.5) provide Tier 1 default emission factors for developed and
 11 developing/transitional countries. These correspond to default leakage rates of 0.15
 12 and 0.35 percent, respectively, with uncertainty ranges up to ± 500 percent. Because
 13 of the wide uncertainty range of the default factors, country-specific leakage rates are
 14 used as reported for year 2005 to UNFCCC (2009) for Annex 1 countries. To split
 15 total losses into residential and non-residential users, measurement results from the
 16 UK are used (Dennett and Vallender 2011). These measure total gas losses from gas
 17 distribution grids in UK to 0.57 percent of throughput, with 80 percent from low
 18 pressure mains systems supplying small consumers. This translates into losses from
 19 residential and non-residential sectors of 1 and 0.23 percent of gas consumption,
 20 respectively. The same proportional split between losses in residential and non-
 21 residential sectors is assumed in all countries. For Non-Annex 1 countries the UK
 22 leakage rates are used as default except for countries in the Former Soviet Union,

1 where losses are assumed comparable to levels reported by Russia to UNFCCC (i.e.
2 about three times the UK leakage rate). Uncontrolled emission estimates are the
3 product of the reported emission factors and the amount of gas consumed.

4 CH₄ emissions from consumer distribution networks can be reduced by replacing old
5 town gas distribution networks made from grey cast iron by polyethylene (PE) or
6 polyvinylchloride (PVC) networks. This typically reduces almost all fugitive
7 emissions from this source (AEAT, 1998). In baseline, all grey cast iron networks are
8 assumed successively replaced in developed countries, reaching 80 percent
9 replacement in 2030 (from the 1990 level). An alternative option is to increase the
10 control frequency. AEAT assumes a doubling of the control frequency from every
11 fourth (baseline assumption) to every second year removes 50 percent of emissions. In
12 the maximum technically feasible reduction scenario, all grey cast iron networks are
13 replaced by 2030. Costs for these options are taken from AEAT (1998).

14 As country-specific emission factors reported to UNFCCC are used and a distinction
15 is made between residential and non-residential sectors, the uncertainty range in the
16 GAINS estimate ought to be considerably lower than the range of -2 to +500 percent
17 for IPCC (2006, Vol.2, Tables 4.2.4 and 4.2.5) Tier 1 emission factors. The assumed
18 uncertainty range is 0 to +30 percent on a global scale.

19 **2.5 Fuel combustion for energy purposes in stationary and mobile**
20 **sources**

21 Apart from leakage in consumer gas distribution networks, CH₄ emissions from
22 energy distribution and use also derive from incomplete combustion of any type of
23 fuel. Emissions are calculated as activity data times a fuel and sector specific emission
24 factor. Activity data is PJ of energy used by fuel and sector as reported and projected
25 by IEA-WEO (2009).

26 Emissions from incomplete combustion in non-residential stationary sources are
27 estimated using default IPCC emission factors (2006, Vol. 2, pp. 2.16-2.23, p. 3.24).
28 For the residential sector, emission factors specified for different types of boilers and
29 fuels are used as reported by Johansson et al. (2004), Kjällstrand and Olsson (2004),
30 Olsson and Kjällstrand (2006) and Delmas (1994). These emission factors are
31 considerably higher than IPCC default factors in particular for biomass combustion.

1 Emissions from combustion in mobile sources are estimated using IPCC (2006, p.
2 3.24) default emission factors. As these are specified per km travelled, they are
3 converted to emissions per energy unit consumed using vehicle specific conversion
4 factors from IEA-WEO (2009). For passenger cars and light duty vehicles, emission
5 factors are specified by fuel and vehicle type and by the emission control standard of
6 the vehicles. For other means of transportation, emission factors are specified only by
7 types of fuel and vehicle, while no default factors by emission control standard are
8 available.

9 No options for mitigating CH₄ emissions are identified for combustion activities.

10 If IPCC default emissions factors (2006, Vol. 2, pp. 2.16-2.23, p. 3.24) are applied
11 also to the residential sector, global emissions from incomplete combustion of fuel for
12 energy purposes in stationary and mobile sources would amount to about 3400 kt CH₄
13 in 2005. This is considerably lower than the estimate of 10800 kt CH₄ when using the
14 Swedish emission factors used in this study. In general, uncertainty is likely to be high
15 in particular for emissions from the residential sector where the variability in emission
16 factors is high between different types of fuels and boilers.

17 **2.6 Coal mining**

18 Formation of coal produces CH₄, which is released to the atmosphere during mining.
19 IPCC (2006, Vol.2, p.4.9) identifies three sources of CH₄ emissions from coal mining:
20 liberation of CH₄ during breakage of coal in the coal mine, post-mining emissions
21 during handling, processing and transportation of mined coal, and emissions from
22 abandoned coal mines. Due to difficulties with obtaining systematic information about
23 the number and size of abandoned coal mines, the emission source is excluded in this
24 study. Hence, only emissions from coal mines in operation are regarded.

25 Following the split of the activity data into brown and hard coal, emissions are
26 calculated separately for the two coal types. All brown coal is assumed to be surface
27 mined, while hard coal is assumed being partly surface mined and partly mined
28 underground.

29 Emissions from brown and hard coal production are calculated as follows:

$$30 \quad E_{BC;it} = A_{BC;it} \times \left[ef_{m;i}^{surface} + ef_{postm;i}^{surface} \right] \quad (9)$$

$$\begin{aligned}
1 \quad E_{HC;it} &= A_{HC;it} \times \left[\gamma_{HC;i}^{undergr} \times \left(ef_{m;i}^{undergr} + ef_{postm;i}^{undergr} \right) \right] + \\
2 \quad &+ A_{HC;it} \times \left[\left(1 - \gamma_{HC;i}^{undergr} \right) \times \left(ef_{m;i}^{surface} + ef_{postm;i}^{surface} \right) \right] \quad (10)
\end{aligned}$$

3 where

$$4 \quad ef_{m;i}^{surface} = ef_{m;i;NOC}^{surface} \times \left(1 - r_{dgas} \right) \times CLEappl_{dgas;i} \quad (11)$$

$$5 \quad ef_{m;i}^{undergr} = ef_{m;i;NOC}^{undergr} \times \left[\alpha_{VAM;i} \times \left(1 - r_{VAM} \right) \times CLEappl_{VAM;i} \right] \quad (12)$$

$$6 \quad + ef_{m;i;NOC}^{undergr} \times \left[\left(1 - \alpha_{VAM;i} \right) \times \left(1 - r_{dgas} \right) \times CLEappl_{dgas;i} \right]$$

7 and where

8 $A_{BC;it}$ and $A_{HC;it}$ are amounts of brown and hard coal produced in country i in
9 year t ,

10 $ef_{m;i;NOC}^{surface}$ is a country-specific no control emission factor for surface
11 mining emissions,

12 $ef_{postm;i}^{surface}$ is the average world IPCC default emission factor for post-
13 mining emissions from surface mines,

14 $ef_{m;i;NOC}^{undergr}$ is a country-specific no control emission factor for underground
15 mining emissions,

16 $ef_{postm;i}^{undergr}$ is the average world IPCC default emission factor for post-
17 mining emissions from underground mines,

18 $\gamma_{HC;i}^{undergr}$ is a country-specific fraction of hard coal being mined
19 underground as opposed to surface mining,

20 $\alpha_{VAM;i}$ is a country-specific fraction of underground mining emissions
21 being released through the ventilation air as opposed to pre-
22 mine degasification emissions,

23 r_{dgas} and r_{VAM} are the removal efficiencies of technologies removing
24 degasification and ventilation air methane, respectively,

1 $CLEappl_{VAM;i}$ is the application of technology removing ventilation air
2 methane, and

3 $CLEappl_{dgas;i}$ is the application of technology removing degasification
4 methane.

5 For hard coal, country-specific fractions for surface vs underground mining are
6 derived from UNFCCC (2010) CRF tables for Annex-1 countries, while all hard coal
7 is assumed mined underground in non-Annex-1 countries. Emissions from surface and
8 underground mining are split into mining and post-mining emissions. Mining
9 emission factors are derived from UNFCCC (2010) CRF tables, from a national
10 source for China (China University of Petroleum, 2008) and using IPCC (2006, Vol.2,
11 p.4.12) world low end default factors whenever country-specific information is
12 unavailable. For post-mining emission factors, default IPCC (2006, Vol.2, p.4.12)
13 factors are used specified separately for surface and underground mines. Underground
14 mining emissions derive from emissions during pre-mine degasification (degas) and
15 from CH₄ being released through the ventilation air system (VAM) during mining.
16 Country-specific fractions of VAM emissions to degasification emissions are taken
17 from USEPA (2003) for a selection of countries (see Table 5). For all other countries,
18 the US fraction of 0.6 VAM to 0.4 degasification emissions is assumed.

19 In 2005, China was mining 38 percent of global coal production and the share is
20 expected to increase in the future. The release of coalbed CH₄ in China is on average
21 assumed to 11 m³/ton coal. This factor is derived from China University of Petroleum
22 (2008) assuming 41 percent of CH₄ is emitted from coal mines with average release of
23 15 m³/t coal, 28 percent from mines releasing 9 m³/t coal, 18 percent from mines
24 releasing 7 m³/t coal, and 13 percent from mines releasing 5 m³/t coal.

25 As shown in Table 6, the weighted average release of CH₄ from global coal mining in
26 2005 is 7.8 m³/t coal produced in GAINS, with on average 1.4 m³/t coal for brown
27 coal and 9.9 m³/t coal for hard coal. This can be compared with the weighted global
28 emission factor of 11.9 m³/t coal when using average default IPCC (2006) emission
29 factors for surface and underground mining in all countries (and applying the same
30 surface/underground split of 2005) and the considerably lower estimates of 6.7 m³/t
31 by USEPA (2003) and 4.3 m³/t by GMI (2009).

1 Emissions from surface mining can be reduced if CH₄ is recovered through pre-mine
2 drainage up to ten years before the surface mining starts (USEPA, 2008). Currently
3 there is only one project known to be recovering and utilizing CH₄ from pre-mine
4 drainage at a surface mine in the US and details about the removal efficiency of this
5 option are uncertain (Sino-US New Energy Sci-Tech Forum, 2009). The CH₄ content
6 of pre-mine drainage gas from surface mines is assumed to 60 percent and it is
7 considered technically possible to recover 90 percent of the drainage gas (USEPA,
8 2010). Drainage gas is assumed to make up 40 percent of surface mining emissions,
9 while the residual mining emissions are released during mining operations.

10 Utilization of drained gas requires upgrading to grid quality (from 60 to 97 percent
11 CH₄) and extended transport capacity of gas either by pipeline or liquefied and
12 transported by ship. The same cost assumptions as applied for recovery and utilization
13 of associated gas from oil and gas production are applied here (see Section 2.1).

14 No mitigation potential is assumed from post-mining emissions.

15 Currently in the US, at least 90 percent of degasification emissions from underground
16 coal mines are recovered and utilized (USEPA, 2010) and this is assumed technically
17 possible in other countries as well. Costs for degasification are taken from Thakur
18 (2006) and include costs for in-mine drilling, underground pipeline costs, and
19 hydraulic fraction of vertical wells and other gob wells. Costs correspond to 0.023
20 Euro/ton coal for mildly gassy coal seams containing less than 2.8 m³ CH₄/ton coal,
21 0.39 Euro/ton coal for medium gassy coal seams containing 2.8-8.5 m³ CH₄/ton coal,
22 and 2.74 Euro/ton coal for highly gassy coal seams containing more than 8.5 m³
23 CH₄/ton coal. It is assumed that 90 percent of degasification emissions can be
24 recovered and utilized from underground mines. For coal seams exceeding 2.8 m³
25 CH₄/ton coal the recovered gas is assumed of a quality high enough (>93 percent) to
26 be injected into natural gas pipelines without further upgrading. Gas recovered from
27 coal seams containing less than 2.8 m³ CH₄/ton coal is assumed to need upgrading
28 from 60 to 97 percent CH₄. Note that extended pre-mine degasification is also
29 expected to reduce the risk of explosions, which is a major health benefit of this
30 measure. Due to difficulties with assessing the value of this benefit, it has not been
31 regarded in the cost assessment.

1 Installation of Ventilation Air Methane (VAM) oxidizers is still in a starting phase,
2 with a handful of installations worldwide, e.g., in Australia, China and the UK
3 (Mattus and Källstrand, 2010), however, the technology used is well known and has
4 been applied frequently for controlling odor and VOC emissions worldwide. The
5 technology oxidizes at least 95 percent of VAM when applied to a ventilation shaft. It
6 uses the energy released during the oxidation to keep the process running, which
7 keeps fuel costs limited to the initial start-up phase. A prerequisite for the oxidation
8 process to run without interruptions is that the CH₄ concentration in the ventilation air
9 is at least 0.3 percent. Securing this concentration level without increasing explosion
10 risks (i.e. CH₄ concentrations in the air should never be in the explosive range
11 between 5 and 15 percent), may in some mines require investments in more efficient
12 ventilation systems.

13 A general assumption is made that it is technically possible to keep CH₄ concentration
14 levels at a steady rate of at least 0.3 percent, and therefore to install self-sustained
15 VAM oxidizers (Mattus and Källstrand, 2010), on 50 percent of the ventilation air
16 emitted from underground coal mines in all countries, with the exceptions of South
17 Africa and India, where current VAM concentration rates are on average 0.1 percent
18 USEPA (2003). With better information about the distribution of VAM concentration
19 rates in different countries, this assumption may need to be revised in the future. It is
20 not assumed possible to install VAM oxidizers on surface mines.

1 Table 5: Assumptions for estimating CH₄ emissions from coal mining in the GAINS model
 2 and resulting implied emission factors (for major coal producing countries).

Region	Country	Fraction of coal mined underground (as opposed to surface)	Fraction of mine gas exiting through ventilation shafts as opposed to pre-mining degasification	Average VAM concentr. rate	Implied emission factor coal mining (total surface and underground mining)		
					No control	With current recovery rate maintained	With maximum technically feasible reduction
					kt CH ₄ /Mt coal produced		
Africa	South Africa	88% ^{a)}	90% ^{f)}	0.1% ^{f)}	1.9	1.9	1.8
	Other Africa	100% ^{b)}	60% ^{b)}	0.3% ^{b)}	8.4	8.4	3.3
Asia	Indonesia	100% ^{b)}	60% ^{b)}	0.3% ^{b)}	8.4	8.4	3.3
	Mongolia	1% ^{d)}	60% ^{b)}	0.3% ^{b)}	1.0	1.0	0.6
	North Korea	100% ^{b)}	60% ^{b)}	0.3% ^{b)}	8.4	8.4	3.3
	Thailand	0% ^{d)}	60% ^{b)}	0.3% ^{b)}	0.9	0.9	0.6
	Vietnam	100% ^{b)}	60% ^{b)}	0.3% ^{b)}	8.4	8.4	3.3
Australia and N. Zealand	Australia	19% ^{c)}	60% ^{b)}	0.4% ^{f)}	3.1	2.5	1.5
	New Zealand	14% ^{c)}	60% ^{b)}	0.3% ^{b)}	2.8	2.8	1.1
Canada	Canada	1% ^{c)}	60% ^{b)}	0.3% ^{b)}	0.6	0.6	0.4
Central Asia	Kazakhstan	13% ^{c)}	60% ^{b)}	0.29% ^{f)}	6.6	6.5	3.2
China	China	100% ^{d)}	60% ^{g)}	0.46% ^{f)}	7.9	7.7	3.6
EU-27	Bulgaria	7% ^{c)}	60% ^{b)}	0.3% ^{b)}	0.9	0.9	0.6
	Czech Rep.	23% ^{c)}	73% ^{f)}	0.26% ^{f)}	3.2	2.2	1.4
	Estonia	0% ^{c)}	60% ^{b)}	0.3% ^{b)}	0.9	0.9	0.6
	Germany	12% ^{c)}	61% ^{f)}	0.3% ^{f)}	1.5	1.0	0.5
	Greece	0% ^{c)}	60% ^{b)}	0.3% ^{b)}	1.1	1.1	0.8
	Hungary	0% ^{c)}	60% ^{b)}	0.3% ^{b)}	0.1	0.1	0.1
	Poland	60% ^{c)}	72% ^{f)}	0.26% ^{f)}	3.4	2.9	1.6
	Romania	10% ^{c)}	60% ^{b)}	0.3% ^{b)}	0.8	0.8	0.6
	Spain	44% ^{c)}	60% ^{b)}	0.3% ^{b)}	3.1	3.1	1.6
	United Kingdom	48% ^{c)}	70% ^{f)}	0.3% ^{b)}	12.4	9.9	4.0
Europe -east	Serbia-Monten.	0% ^{b)}	60% ^{b)}	0.3% ^{b)}	0.9	0.9	0.6
	Turkey	4% ^{c)}	60% ^{b)}	0.3% ^{b)}	2.7	2.7	1.5
	Ukraine	99% ^{c)}	60% ^{b)}	0.3% ^{f)}	24.6	23.1	7.6
India	India	93% ^{e)}	60% ^{b)}	0.1% ^{f)}	3.0	3.0	2.3
Latin and Central	Brazil	60% ^{d)}	60% ^{b)}	0.3% ^{b)}	5.4	5.4	2.2
	Mexico	100% ^{b)}	60% ^{b)}	0.5% ^{f)}	8.4	8.4	3.3
	Other L. America	100% ^{b)}	60% ^{b)}	0.3% ^{b)}	8.4	8.4	3.3
Russia	Russia	35% ^{c)}	70% ^{f)}	0.38% ^{f)}	8.3	8.2	4.1
United States	United States	33% ^{c)}	60% ^{f)}	0.39% ^{f)}	3.5	2.8	1.5

Sources: a) UNFCCC (2000), b) Author's assumption, c) UNFCCC (2010), d) IEA-WEO (2009) fraction of hard coal mined, e) UNFCCC (2004), f) USEPA (2003), g) Current VAM fraction reported for China is 88 percent (USEPA, 2003), however, due to an expected increase in degasification (GMI,2011) it is assumed 60 percent also for China.

3
4
5
6

1 Table 6: Comparison of implied CH₄ emissions (mining and post-mining) released before
 2 recovery per ton coal produced in GAINS and other sources, unit: m³ CH₄/ton coal produced.

Region	Country	GAINS model	GAINS source	USEPA (2003)	GMI (2009)
Africa	South Africa	2.6	UNFCCC -NIR (2000), IPCC (2006)	2.0	2.1
	Other Africa	11.7	IPCC (2006) low end default factor		
Asia	Indonesia	11.7	IPCC (2006) low end default factor		0.2
	Mongolia	1.4	IPCC (2006) low end default factor		0.5
	North Korea	11.7	IPCC (2006) low end default factor		
	Thailand	1.2	IPCC (2006) low end default factor		
	Vietnam	11.7	IPCC (2006) low end default factor		2.6
Australia and N. Zealand	Australia	4.3	UNFCCC-CRF (2010)	4.1	4.1
	New Zealand	3.9	UNFCCC-CRF (2010)		5.4
Canada	Canada	0.9	UNFCCC-CRF (2010)		1.0
Central Asia	Kazakhstan	9.3	UNFCCC-CRF (2010)	5.9	5.5
China	China	11.0	China University of Petroleum (2008)	9.4	4.7
EU-27	Bulgaria	1.1	UNFCCC-CRF (2010)		3.7
	Czech Rep.	4.4	UNFCCC-CRF (2010)	5.5	5.5
	Estonia	1.2	UNFCCC-CRF (2010)		
	Germany	2.0	UNFCCC-CRF (2010)	5.3	2.9
	Greece	1.6	UNFCCC-CRF (2010)		
	Hungary	0.1	UNFCCC-CRF (2010)		3.6
	Poland	4.8	UNFCCC-CRF (2010)	6.2	5.1
	Romania	0.9	UNFCCC-CRF (2010)		71.1
	Spain	4.3	UNFCCC-CRF (2010)		4.4
	United Kingdom	17.2	UNFCCC-CRF (2010)	11.8	23.8
Europe -east	Serbia Montenegro	1.2	IPCC (2006) low end default factor		
	Turkey	3.7	UNFCCC-CRF (2010)		2.2
	Ukraine	34.2	UNFCCC-CRF (2010)	29.0	30.9
India	India	4.2	UNFCCC -NIR (2004), IPCC (2006)	1.9	3.2
Latin and Cenral America	Brazil	7.5	IPCC (2006) low end default factor		
	Mexico	11.7	IPCC (2006) low end default factor	7.7	16.5
Other Latin America	Other Latin America	11.7	IPCC (2006) low end default factor		
	Russia	11.6	UNFCCC-CRF (2010)	10.2	6.7
United States	United States	4.9	UNFCCC-CRF (2010)	5.1	3.8
World -weighted		7.8		6.7	4.3
World-IPCC (2006)		11.9	IPCC <i>average</i> default mining+post-mining (surface: 1.2 m ³ /t, undergr: 19.1 m ³ /t)		

3

4

5 USEPA (2003, p.30) estimate costs for installing VAM oxidizers without energy
 6 recovery on the basis of measured concentrations and flow rates on 58 ventilation
 7 shafts at underground coal mines in the US. The resulting curve is adapted to estimate
 8 costs in non-US countries. Table 7 shows a comparison between the costs presented
 9 by USEPA for different VAM concentration rates and the projected costs as estimated
 10 by GMI (2008) for installing 11 sets of VAM oxidizers without energy recovery on
 11 the Xiaodongshan Shaft in the Jincheng mining area in China. As shown, cost
 12 estimates by USEPA (2003) are considerably lower. The higher costs projected for the

1 Chinese coal mine are applied here, but adapted to different VAM concentration
 2 levels using a scaling proportional to the curve reported by USEPA (2003).
 3 Table 7: Basis for calculation of VAM oxidizer costs, Euro/t CO₂eq (2005 prices).

VAM concentr ation rate	GAINS model			USEPA (2003, p.30)	
	VAM ox installations on the Xiaodongshan Jingshen Mine (GMI, 2010) scaled to VAM concentr rates as in USEPA (2003)			Total cost	Total cost
	Investment cost	O&M cost	Total cost		
Lifetime=16 yrs, discount rate=4%	Lifetime=16 yrs, discount rate=4%	Lifetime=16 yrs, discount rate=4%	Lifetime=16 yrs, discount rate=15%	Lifetime=16 yrs, discount rate=4% ^a	
<0.2%	VAM oxidation not possible				
0.20%	2.67	2.52	5.19	2.85	2.16
0.30%	2.53	2.39	4.92	2.7	2.04
0.40%	2.39	2.25	4.64	2.55	1.93
0.50%	2.25	2.12	4.37	2.4	1.81
1.00%	1.54	1.46	3.00	1.65	1.25

4 ^a Author assumes 50% investment cost and 50% O&M costs.

5 If VAM oxidizer technology is combined with improved ventilation systems, it may
 6 be possible to extend the installation of oxidizers as it then becomes possible to better
 7 control a steady rate of at least 0.3 percent CH₄ in the ventilation air without
 8 jeopardizing security. An improved ventilation system is assumed to double the
 9 ventilation capacity of the mine compared with a conventional system, which would
 10 double the amount of electricity used for ventilation. Unruh (2002) estimates the total
 11 energy consumption by the US coal mining industry in 1998 to 25.5 billion kWh.
 12 According to Papar et al. (1999), the energy use of the entire US mining industry is 44
 13 billion kWh in 1998 whereof 12 billion kWh (27 percent) is used for mine ventilation.
 14 If this fraction is applied to the coal mining industry, the energy consumption for coal
 15 mine ventilation in the US is 6.9 billion kWh in 1998. With a total production of
 16 1117.5 Mt coal in 1998 (whereof 63 percent surface and 37 percent underground
 17 mined), then current use of electricity for mine ventilation per ton coal produced is 6.2
 18 kWh (EIA, 2010c). The total cost for this option is the sum of the cost for improved
 19 ventilation and the installation of VAM oxidizers. Just like the degasification options,
 20 upgraded ventilation in coal mines is likely to reduce the risk of explosions, which is a
 21 major health benefit of this measure. Due to difficulties assessing the value of this
 22 benefit, it has not been regarded in the analysis. It is assumed that with improved
 23 ventilation it is possible to extend the application of VAM oxidizers to 70 percent of
 24 VAM emitted from underground mines in all countries (except S Africa and India).
 25 IPCC (2006, Vol. 2, p. 4.12) define an uncertainty range for underground coal mining
 26 emission factors of ±40 percent with the low-end estimate for mining depths < 200 m

1 and the high-end estimate for depths > 400 m. Taking several country-specific factors
 2 into account, including effects of mine depths in major producer countries, the
 3 uncertainty range for the GAINS estimate is evaluated at ±20 percent on a global
 4 scale. For surface mining GAINS adopts the uncertainty range of a factor two
 5 suggested by IPCC (2006, Vol.2, Table 4.1.4). For post-mining emission factors,
 6 IPCC (2006, Vol.2, pp. 4.12, 4.19) provide uncertainty ranges of ±100 percent for
 7 surface mining and ±60 percent for underground mining.

8 Uncertainty is also present in the fixed assumptions about removal efficiencies. E.g.,
 9 country- and site-specific obstacles for implementation of VAM oxidizers have not
 10 been fully explored due to lack of site-specific information.

11 Including emissions from abandoned coal mines in the analysis would increase CH₄
 12 emissions from this sector. Currently, CH₄ emissions from abandoned coal mines in
 13 the US correspond to about 9 percent of VAM emissions (UNFCCC, 2011). If this
 14 factor is applied worldwide, the global release is in 2005 is 1400 kt and growing to
 15 2400 kt CH₄ in 2030.

16 **2.7 Livestock**

17 CH₄ emissions from livestock emerge from enteric fermentation during the digestive
 18 process in the stomachs of ruminants as well as from manure management. GAINS
 19 estimate CH₄ emissions separately for the animal types dairy cows, non-dairy cattle,
 20 pigs, sheep and goats, camels, buffalo, and horses. For dairy cows, non-dairy cattle
 21 and pigs, animal types are further split by whether animals are subject to liquid or
 22 solid manure management. CH₄ emissions from livestock are estimated as the sum of
 23 the two emission types *n* (enteric fermentation and manure management) for a certain
 24 animal type *s* in country *i* and year *t*:

25
$$E_{its} = \sum_{lmns} [A_{itls} * ef_{ilns}^{NOC} * (1 - remeff_{mns} * h_{itms}) * Appl_{itlsm}], \quad (13)$$

26 where A_{itls} is the number of animals of type *s* in country *i* and year *t*, with
 27 manure management *l* (solid or liquid),
 28 ef_{is}^{NOC} is the no control emission factor for emission type *n*, animal
 29 type *s* in country *i* and subject to manure management *l*,

1 $remeff_{mns}$ is the removal efficiency of technology m when applied to
2 emissions of type n and animal type s ,

3 h_{itms} is a factor correcting for application limitations of technology
4 m , e.g., indoor housing rates for feed options or large farm rate
5 for farm-scale anaerobic digestion,

6 $Appl_{itlms}$ is the application rate of technology m to animal type s with
7 manure management l , in country i and year t .

8 Activity data is number of animals by type. Sources for historical and current animal
9 numbers are EUROSTAT (2009) for EU countries, UNFCCC (2010) CRF tables for
10 other Annex 1 countries, and FAOSTAT (2010) for non-Annex1 countries.
11 Projections for EU are taken from the CAPRI Model (2009) to be consistent with the
12 EU effort sharing agreement (EC, 2009). For other world regions, future drivers are
13 based on forecasts by major world regions as developed by FAO (2003).

14 Implied emission factors reported by countries to UNFCCC (2010) are used when
15 available, complemented with regional specific IPCC (2006, Vol.4, Tables 10.10,
16 10.11, 10.14 and 10.15) default emission factors.

17 For dairy cows, enteric fermentation emissions per animal are affected by the milk
18 productivity of the cow. This effect is particularly accentuated for highly productive
19 milk cows. To capture this, the no control emission factor for dairy cows is specified
20 as the sum of a fixed emission factor per animal for cows producing up to 3000 kg per
21 head per year and an additional term describing the emission factor per milk yield for
22 milk production exceeding the productivity level 3000 kg per animal per year, i.e.,

23 $ef_{it;cow}^{NOC} = ef_i^{animal} + ef_i^{milk} * (x_{it} - 3000)$ (14)

24 where ef_i^{animal} is the default emission factor for cows in country i producing
25 3000 kg milk per year,

26 ef_i^{milk} is the emission factor per kt milk produced above the threshold
27 level 3000 kg milk per animal per year, and

28 x_{it} is the average milk yield per animal in country i and year t .

29

1 CH₄ emissions from enteric fermentation in cows and non-dairy cattle can be reduced
2 by changing animal diets, e.g., by replacing roughage with concentrates or introducing
3 more fat in the diet. As feed changes require control over what animals eat, they are
4 only assumed applicable to animals when these are kept indoor. Removal efficiency
5 for this option is five to ten percent when applied (Ecofys 2009, ECCP 2003, Lovett et
6 al. 2006, Boadi et al. 2004). Costs for feed changes refer to the cost of replacing 6
7 percent of the daily dry matter intake (DMI) by soy oil (Ecofys, 2009). Indoor
8 housing rates are taken from Klimont and Brink (2003) for European countries and
9 extended to other countries by applying European rates in countries with similar
10 livestock structure. No indoor housing of cows and cattle is assumed for Africa,
11 Central Asia and Latin America.

12 CH₄ emissions from liquid manure management from cows, non-dairy cattle and pigs
13 can be reduced by treating the manure in anaerobic digesters (AD) (Ecofys 2009,
14 AEAT 2001). AD plants produce biogas, which can be recovered and utilized for
15 energy purposes. Three scales of AD installations for treatment of manure are
16 considered in GAINS: farm-, household-, and community- scale digesters. Farm scale
17 AD is assumed economically feasible for handling manure from all large farms, i.e.,
18 farms with a minimum size of 100 dairy cows, 200 beef cattle or 1000 pigs. Large
19 farm fractions are taken from EUROSTAT (2008) and European rates are applied to
20 other countries with similar agricultural structures. Removal efficiency for farm-scale
21 AD is assumed 80 percent (Ecofys 2009). For developing countries, GAINS also
22 include the option of treating manure in household-scale ADs (An et al. 1997;
23 Dhingra et al. 2011). These are common in some developing countries and digest
24 manure and other organic waste material from farm households. The generated biogas
25 is recovered and utilized by the households, e.g., to fuel cooking stoves. The digesters
26 are cheap to install but labour intensive to operate effectively (An et al. 1997), which
27 make application limited to developing countries with low average wage rates in
28 agriculture¹. Household size digesters are currently assumed to treat 5 to 10 percent of
29 cattle manure in India (MNRE, 2006) with an assumed possible extension to 30

¹ Afghanistan, Armenia, Azerbaijan, Bangladesh, Bhutan, Cambodia, China, Egypt, Turkmenistan, Tadjikistan, Uzbekistan, Indonesia, Kazakhstan, Georgia, North Korea, Kyrgyzstan, Laos, Malaysia, Mongolia, Myanmar, Nepal, Pakistan, Philippines, Sri Lanka, Thailand and all African countries except South Africa.

1 percent of livestock manure in all developing countries. Community-scale AD plants
2 receive manure from several farms in the vicinity of the plant. Transportation of
3 manure for long distances is costly and increase emissions of both CH₄ and carbon
4 dioxide. This option is therefore only applied in GAINS to countries with intensive
5 pig farming (more than 200 pigs per m²), which is Belgium, Denmark, the
6 Netherlands and Malta, and in India to the extent that current use of village size
7 digesters has been documented (MNRE, 2006). Costs for farm-scale and community-
8 scale AD installations are taken from Ecofys (2009). Cost for household-scale ADs is
9 taken from An et al. (1997).

10 Uncertainty in baseline emission estimates is primarily linked to difficulties with
11 measuring emissions from a large number of point sources (animals) and to convert
12 highly variable measurement results to default emission factors. Comparability of
13 results also suffers from a large variation in the measurement techniques used (Iqbal
14 et al. 2008). GAINS adopts the IPCC (2006, Vol.4, p.10.33) uncertainty range of ± 30
15 percent for enteric fermentation emissions and ± 25 percent for manure emissions
16 when assessed on a global scale.

17 Uncertainty about the future mitigation potential is particularly present for emissions
18 from enteric fermentation and linked to both the selection of mitigation options
19 included in the model and to highly uncertain removal efficiencies. This is an area of
20 intensive research and several options show promising results, however, widespread
21 adoption appears slow. These include the use of vaccination against methanogenic
22 bacteria in the rumen (Wright et al., 2004; Ecofys 2009) and the use of propionate
23 precursors as feed additive or through genetic engineering of feed plants for grazing
24 animals (Newbold et al. 2005, Ecofys 2009). As these options are not yet in
25 commercial use, they have not been considered feasible as mitigation options in the
26 2030 timeframe in GAINS.

27 General productivity increases in milk and meat production through genetic
28 engineering is sometimes mentioned as a possible CH₄ mitigation option with the
29 assumption that a cut in animal stocks can outweigh increases in emissions per
30 animal. Lovett et al (2006) show that general productivity increases tend to increase
31 system emissions because of negative effects on animal health, which increase the
32 fraction of non-productive (young) animals in the stock. General productivity
33 increases is therefore not regarded as a viable CH₄ mitigation option in this study.

1 Although the current technical mitigation potential for enteric fermentation in GAINS
 2 is limited to a few percent on a global scale, it may grow suddenly in response to new
 3 developments in mitigation options and their commercial availability.

4 **2.8 Rice cultivation**

5 CH₄ emissions from rice cultivation result from anaerobic decomposition of organic
 6 material in flooded rice fields. Emissions depend among other factors on the season,
 7 soil characteristics, soil texture, use of organic matter and fertilizer, climate, as well as
 8 agricultural practices (IPCC, 2006, Vol.4, p. 5.45). The emission calculation
 9 methodology used in GAINS follows the IPCC guidelines (2006, p. 5.49) and adopts
 10 IPCC default emission factors unless country-specific factors have been reported to
 11 UNFCCC (2009). The IPCC method is based on the annual harvested area with
 12 scaling factors for different water regimes. In GAINS, these are translated into three
 13 cultivation activities:

- 14 • *Continuously flooded cultivation area*: fields have standing water throughout
 15 the growing season and only drying out for harvest.
- 16 • *Intermittently flooded cultivation area*: fields have at least one aeration period
 17 of more than three days during the growing season. Compared with
 18 continuously flooded rice fields, IPCC suggests that intermittently flooded rice
 19 fields emit 27 to 78 percent of continuously flooded fields, where the range
 20 depends on if the fields are rainfed or irrigated. GAINS uses the assumption
 21 50 percent emissions per hectare from intermittently flooded compared with
 22 continuously flooded fields.
- 23 • *Upland rice cultivation area*: fields are never flooded for a significant period
 24 of time and are not assumed to emit CH₄.

25 CH₄ emissions from rice cultivation in country *i* in year *t* are calculated as follows:

$$26 \quad E_{it} = \sum_{sm} A_{it} * ef_{i,flood}^{IPCC} * h_i * \beta_s * V_{is} * (1 - remeff_{sm}) * Appl_{it,sm}, \quad (15)$$

27 where A_{it} is the rice cultivation area in country *i* in year *t*,

28 $ef_{i,flood}^{IPCC}$ is the IPCC default emission factor for CH₄ emissions

29 from flooded rice fields (1.3 kg CH₄ ha⁻¹ day⁻¹),

1 h_i is the duration of the growing season expressed in days
 2 per year (=185 days per year),
 3 β_s is an emission scaling factor for water regime s (=1 for
 4 continuously flooded, =0.5 for intermittently flooded, and =0
 5 for upland rice).
 6 V_{is} is the fraction of rice cultivated land under water regime s ,
 7 $remeff_{sm}$ is the removal efficiency of technology m when applied to
 8 water regime s , and
 9 $Appl_{itsm}$ is the application rate of technology m when applied to water
 10 regime s .

11 Activity data for rice cultivation is measured in million hectares of land and is taken
 12 from FAOSTAT (2008) with projections based on FAO (2003). Data on country-
 13 specific water regimes is taken from IRRI (2007).

14 Introducing intermittent aeration of continuously flooded rice fields reduces CH₄
 15 emissions, while saving water, but is also likely to increase weed growth in the fields
 16 (Barrett et al. 2004, Ferrero and Nguyen 2004). This increases labour costs by an
 17 estimated 20 percent (Barrett et al. 2004), which is equivalent to about 60 additional
 18 work hours annually per hectare in developing countries (Heytens, 1991) and 12
 19 additional work hours annually per hectare in developed countries, where herbicides
 20 are used for controlling weeds (Shibayama, 2001). According to IRRI (2007),
 21 intermittent aeration of continuously flooded rice fields may reduce water use by 16 to
 22 24 percent. The mitigation potential of this option is assumed 22 percent, based on the
 23 IPCC default emission factor for intermittent aeration of continuously flooded rice
 24 fields. Assuming that continuously flooded rice fields need 1000 mm water input per
 25 year (Bouman, 2001) and the average cost of irrigated water is 0.02 US\$ per m³
 26 (FAO, 2004), then saving 22 percent of water corresponds to a cost-saving of about
 27 35 Euro per ha.

28 Certain rice hybrids may affect CH₄ emissions. By careful selection of low CH₄
 29 producing hybrids, emissions can be ten percent lower (ADB 1998). ADB (1998)
 30 estimates that Chinese rice yields may increase by as much as 10 to 20 percent from
 31 switching to low CH₄ rice hybrids. In other parts of the world, where high yield rice

1 hybrids are already in extensive use, potentials for additional yield increases are likely
2 to be lower. In GAINS is assumed that the potential reduction in CH₄ emissions from
3 switching to alternative rice hybrids is 10 percent with a 3 percent increase in crop
4 yield, when applied as the sole option. When applied in combination with other
5 options, like intermittent aeration of continuously flooded fields, the removal
6 efficiency of this option is set to 5 percent.

7 Application of sulphate-containing substrates to rice fields reduce CH₄ emissions
8 because CH₄ producing bacteria compete for the same substrate as the sulphate
9 reducing bacteria (van der Gon et al. 2001). The associated costs are the costs of
10 acquiring sulphate containing fertilizers like e.g., ammonium sulphate and spreading
11 them on the fields. In GAINS, this option is assumed to remove 20 percent of
12 emissions when applied as a sole option and 10 percent when applied in combination
13 with other options.

14 Estimation of emissions from soils belong to one of the most uncertain estimates in
15 emission inventories in general (Winiwarter and Rypdal 2001, Winiwarter and Muik
16 2010). In GAINS, baseline emission estimates are scaled for different water regimes,
17 however, IPCC (2006, Vol. 4, p. 5.44) recommends in addition to scale with the
18 application of organic amendments. As it was not possible to collect country-specific
19 data for organic amendments, variation in this factor is not regarded in estimations.
20 IPCC (p. 5.52) provide a default value for the uncertainty in scaling fraction estimates
21 of ± 20 percent, which is adopted in this study.

22 **2.9 Open burning of agricultural waste residues**

23 This sector refers to open burning of agricultural waste, e.g., burning of plant residues
24 on fields. Activity data is amount of agricultural waste burned, which is taken from
25 UNFCCC (2009) for Annex-1 countries and from Niemi (2006) for countries without
26 reporting to UNFCCC. A few countries (France, Germany, Ireland, Luxembourg,
27 Netherlands, Sweden and the UK) report no generation of agricultural waste for open
28 burning and hence these countries have no emissions at all from this source. The
29 uncontrolled emission factor used is the IPCC default emission factor for open
30 burning of waste (IPCC, 2006, Vol.5, p.5.20) which corresponds to 6.5 kt CH₄ per Mt
31 waste burned.

1 The maximum technically feasible reduction of emissions from introducing a ban on
2 agricultural waste burning is set to 3 kt CH₄ per Mt waste. Due to enforcement
3 problems, it is not assumed that all of emissions from this source can be controlled
4 with a ban. The cost of mitigating methane through this option is set to zero, because
5 it is assumed that this is a measure that would be implemented primarily to control
6 other types of airborne emissions like VOCs and black carbon. CH₄ control would
7 come as a co-benefit.

8 IPCC (2006, Vol. 5, p.5.23) estimate uncertainty in default emission factors for
9 combustion of solid waste to an order of magnitude.

10 It should be noted that due to lack of systematic information, GAINS does not cover
11 emissions from pre-scribed burning of savannahs and human-induced forest fires.
12 Estimates by USEPA (2011b) suggest that a more complete coverage of emissions
13 from open biomass burning would add considerably to emissions. USEPA estimates
14 global emissions from all three sources to about 20 000 kt CH₄ per year, to be
15 compared with about 3000 kt CH₄ per year estimated in GAINS for burning of
16 agricultural waste residues only.

17 **2.10 Biodegradable solid waste**

18 CH₄ from municipal and industrial solid waste is generated when biodegradable
19 matter is digested under anaerobic conditions in landfills or during temporary storage
20 of waste aimed for different types of treatment. CH₄ may also be released during
21 loading or emptying of the reactor when organic waste is treated in anaerobic
22 digesters to produce biogas or energy. The activity data used in GAINS is the total
23 amount of waste generated before it is disposed of to landfills or other types of
24 treatment. Amounts of waste generated are first split into municipal or industrial solid
25 waste and then by waste composition for municipal solid waste and by manufacturing
26 industry sub-sector for industrial solid waste.

27 CH₄ from waste deposited on landfills is formed and released with a time delay of up
28 to several decades. IPCC (2006, Vol. 5, Ch. 3) recommends the use of a First-order-
29 decay model taking up to fifty years disposal into account. The GAINS model
30 structure does not allow for implementation of a full First-order-decay model. Instead,
31 a simplified structure is used, where the delay between waste disposal and CH₄
32 release is accounted for as a lag in the activity data of 10 years for fast degrading

1 organic waste like food and garden waste and 20 years for more slowly degrading
 2 waste like paper, wood and textile waste. The lags correspond to approximate average
 3 half-life values for the respective waste types (IPCC, 2006, Vol.5, Tables 3.3 and 3.4).
 4 CH₄ emissions from municipal (or industrial) solid waste in country *i* in year *t* are
 5 estimated as the sum of emissions from a certain waste type *s* (or industry sector)
 6 summed over emissions from waste diverted to waste treatment option *m* :

$$7 \quad E_{it} = \sum_s \sum_m A_{i;(t-y_s);s} * ef_s * (1 - remeff_{sm}) * Appl_{it sm} \quad (16)$$

8 where $A_{i;(t-y_s);s}$ is amount of waste type (or industry sector) *s* generated in year
 9 $t-y_s$, where y_s is the average lag in CH₄ release assumed for
 10 waste type (or industry sector) *s*,

11 $remeff_{sm}$ is the removal efficiency of waste treatment option *m*, and

12 $Appl_{it sm}$ is the application of waste treatment option *m* to waste type (or
 13 industry sector) *s* in country *i* in year *t*, and

14 ef_s is the IPCC default emission factor for waste type (or industry
 15 sector) *s* deposited in a landfill without recovery of landfill gas.

16

17 From IPCC (1997, Vol.3, Ch.6, Equation 1) the following expression for ef_s is
 18 obtained:

$$19 \quad ef_s = DOCm_s * DOCf * MCF * F * 16/12 * (1 - OX), \quad (17)$$

where

$DOCm_j$ is the fraction of Decomposable Organic Carbon (DOC) in waste
 type/sector *s*,

$DOCf$ is the fraction of DOC that can decompose (default used is 0.5),

MCF_i is the Methane Correction Factor correcting for aerobic decomposition
 and vary with the management standard of the landfills,

F is the fraction of CH₄ in generated landfill gas (default used is 0.5),

$16/12$ is the molecular weight ratio CH₄/C,

OX is the oxidation factor correcting for increased oxidation from covering of
 landfills (default used is 0.1).

1 IPCC (2006, Vol.5, Tables 2.4 and 2.5) default factors are used for the content of
2 decomposable organic carbon (*DOC_m*) in different types of biodegradable waste as
3 well as for the Methane Correction Factor (MCF) applied to different management
4 standards of the landfills.

5 Activity data used is country-specific amounts of waste generated from 1980
6 onwards. Historical reported waste generation rates are taken from various sources as
7 presented in Table 8 together with approximations made for countries where data is
8 missing. While data on MSW generation is available for most parts of the world,
9 generation rates for industrial solid waste are rare. It has only been possible to find
10 European data (EUROSTAT, 2005) on industrial waste generation. The European
11 rates per value added are used to extend the data to other world regions as shown in
12 Table 8. This is of course a very crude assumption, as both differences in production
13 technologies used as well as barriers in global markets are likely to affect amounts of
14 waste generated and the relative value of the product in the market, which in turn
15 affects the amount of waste generated per value added. It would have been desirable
16 to relate industrial waste amounts to physical rather than monetary units in order to at
17 least control for differences in product market values, however, data on physical
18 production units coupled with associated waste amounts is not readily available.

19 The historical waste generation rates are used as basis for future projections. Waste
20 generation elasticity estimates are produced using a detailed data set from
21 EUROSTAT (2005) covering 31 European countries with annual data from 1985 to
22 2003. For municipal solid waste, the data set comprise 236 observations in an
23 unbalanced panel. Generation of municipal solid waste (MSW) per capita is estimated
24 as a function of GDP per capita (IMF, 2006) and urbanization rate (UNstat, 2009):

$$25 \log(MSWcap)_{it} = \alpha + \beta_1 * \log(GDPcap)_{it} + \beta_2 * \log(urbrate)_{it} + \varepsilon_{it}, \quad (18)$$

26 where

27 *MSWcap* is kg MSW per capita per year,

28 *GDPcap* is the average annual Gross Domestic Product in Euro per capita,

29 *urbrate* is the fraction of the total population living in urban areas,

30 $\varepsilon_{it}=u_i+v_{it}$ is an error term which is separated into an individual effects term and a
31 residual omitted variables term, and

1 $\varepsilon_{it} \sim IID(0, \sigma_{\varepsilon}^2)$ is an error term which is independent and identically distributed.

2 Estimations are conducted in LIMDEP 8.0 (Greene, 2005) using panel data methods,
3 i.e. estimating OLS, fixed effect and random effect models². A Lagrange multiplier
4 (LM) test of poolability show that the fixed or random effect models are preferred to
5 the OLS model. A Hausman specification test shows that the fixed effect model is
6 preferred to the random effect model. Results are presented in Table 9 and show that
7 income per capita affects MSW generation per capita with an elasticity of 0.48
8 (significant at a 1 percent level) on average for the whole sample³. The elasticity for
9 the urbanization rate has an expected negative sign (significant at a 5 percent level).

10 The elasticity for generation of industrial solid waste on a sub-sector level is estimated
11 in response to changes in value added for the industry sub-sector (UNIDO 2006):

$$12 \log(INW)_{it} = \alpha + \beta * \log(VA)_{it} + \varepsilon_{it}, \quad (19)$$

13 where

14 INW is Mt industrial solid waste per year,

15 VA is the value added at factor cost in M Euro per year,

16 $\varepsilon_{it} = u_i + v_{it}$ is an error term which is separated into an individual effects term and a
17 residual omitted variables term, and

18 $\varepsilon_{it} \sim IID(0, \sigma_{\varepsilon}^2)$ is an error term which is independent and identically distributed.

19 Again panel data methods are used. For industrial solid waste, sample sizes are rather
20 small, less than eighty observations, however as shown in Table 9, in the preferred
21 random effect model the resulting parameter estimates are significant at a 1 percent
22 level and with the expected positive signs. Least elastic to changes in value added to

² In a fixed effect model the variance within each country is separated out and the regression is performed only on the within variance, while the variance between countries is captured in country-specific constants. This has the advantage that the variance in waste amounts per capita that depends exclusively on country-specific differences is controlled for. In the random effects model estimates are based on a weighted average of the within and between country variances. In the OLS model, the within and between country variances are bluntly added up without using weights (Hsiao, 1986).

³ Separate models for Western Europe (EU-15 Norway and Switzerland) and Eastern Europe (EU-12, Croatia and Macedonia) were also run, however, with no large differences in parameter estimates compared with the full sample runs.

1 amounts of waste generated is wood and wood products industry (0.3), while pulp and
 2 paper has the highest elasticity (1.0).

3 The elasticity estimates for waste generation are used to calculate predicted future
 4 values for generation of solid waste.

5 Table 8: Baseline assumptions for solid waste amounts generated.

Sector	World region	Basis for future projections	Annual waste generated	Source		
Municipal solid waste (MSW)						
	EU-27, Iceland, Norway, Switzerland, Turkey	MSW in kg/person 2005	300 to 793	EUROSTAT 2009		
	USA	MSW in kg/person 1996	453	IPCC (1997, Vol.2, p.6.6-6.7); Eawag (2008)		
	Canada		493			
	Australia		460			
	New Zealand		485			
	Russia		318			
	Japan		157			
	India		73			
	China		307			
	Indonesia		186			
	Brazil		168			
	Chile		537			
	Peru		360			
	Guatemala		215			
	Egypt		146			
	Nigeria	146				
	Rest of Africa	MSW in	146	Author's assumptions		
	Rest of Former Soviet Union and Eastern Europe	kg/person 1996	318			
	Rest of Latin America		360			
	Rest of South Asia		186			
Industrial solid waste						
Food, beverages and tobacco	EU-15	industrial solid waste per value added in 2000, ton/M Euro value added	390	EUROSTAT 2005		
	EU-10 (New Member States)		724			
Pulp and paper	EU-15		271			
	EU-10 (New Member States)		542			
Textile, footwear and leather	EU-15		105			
	EU-10 (New Member States)		154			
Wood and wood products	EU-15		963			
	EU-10 (New Member States)		1544			
Rubber and plastics	EU-15		86			
	EU-10 (New Member States)		106			
Other manufacturing	EU-15		244			
	EU-10 (New Member States)		1331			
	Other industrialized countries				like EU-15	Author's assumptions
	Other transitional or developing countries				like EU-10 (New Member States)	

6

7

1

2 Table 9: Results from estimations of elasticity for generation of municipal and
3 industrial solid waste. Values in brackets are t-values. Preferred models in italics.

Dependent variable	Unit	No. of obs.	Explanatory variable	OLS	Fixed effect	Random effect	LM-test	Hausman-test
Municipal solid waste	Mt per capita	236	Constant	4.60 (27.6)	<i>n.a.</i>	2.67 (10.5)	235.2	43.6
			GDP per capita	0.17 (10.6)	<i>0.48 (15.9)</i>	0.37 (14.6)		
			Urbanization rate	0.05 (0.53)	<i>-0.17 (1.87)</i>	0.37 (14.6)		
			R-square	0.39	<i>0.85</i>	<i>n.a.</i>		
Food, beverages and tobacco industry waste	Mt	70	Constant	0.77 (0.90)	<i>n.a.</i>	<i>0.78 (0.72)</i>	18.7	0.01
			Value added	0.83 (7.73)	0.83 (3.27)	<i>0.81 (5.82)</i>		
			R-square	0.47	0.71	<i>n.a.</i>		
Pulp and paper industry waste	Mt	70	Constant	-0.12 (0.20)	<i>n.a.</i>	<i>-1.35 (1.72)</i>	54.4	0.24
			Value added	0.85 (9.85)	1.07 (7.84)	<i>1.03 (9.56)</i>		
			R-square	0.59	0.95	<i>n.a.</i>		
Textile, leather and footwear industry waste	Mt	71	Constant	-1.38 (1.99)	<i>n.a.</i>	<i>-0.53 (0.48)</i>	67.8	6.4
			Value added	0.88 (9.26)	-0.32 (0.72)	<i>0.74 (4.74)</i>		
			R-square	0.55	0.89	<i>n.a.</i>		
Wood and wood products industry waste	Mt	71	Constant	3.12 (4.79)	<i>n.a.</i>	<i>3.64 (2.66)</i>	173.3	0.78
			Value added	0.47 (4.44)	0.23 (1.39)	<i>0.33 (4.58)</i>		
			R-square	0.22	0.83	<i>n.a.</i>		

4

5

6 CH₄ emissions from biodegradable solid waste can be controlled by separating out
7 different types of waste treatment for recycling, composting, anaerobic digestion or
8 incineration. Following the EU waste legislation, i.e., the Waste Directive (EC, 2006)
9 and the Landfill Directive (EC, 1999), separation and treatment of biodegradable
10 waste should be preferred to landfill disposal. Landfill disposal of biodegradable
11 waste must be reduced by at least 65 percent between 1995 and 2016 in all EU
12 member states and all landfill sites must have gas recovery facilities installed by 2009.

13 In GAINS, the maximum feasible reduction of CH₄ emissions in the waste sector is
14 modelled as an “optimal” waste treatment path as defined by the current EU
15 legislation. Source separation of waste for recycling or energy recovery purposes is
16 preferred to landfill disposal with gas recovery. In the optimal case, all biodegradable
17 waste is source separated from the waste stream and none is disposed of to landfills.
18 Some EU member states (Denmark, Germany and Sweden) are close to this optimum
19 and have since 2005 introduced a complete ban on landfilling of biodegradable waste.

20 A list of waste treatment options considered in GAINS are presented in Table 10
21 together with a definition of the optimal control. In baseline emissions, the effects of

1 already implemented waste treatment options as well as future effects of adopted
2 legislation are taken into account in applied control strategies. Information on current
3 adoption of waste treatment are taken from UNFCCC (2009) CRF and NIR and
4 complemented with other sources (IPCC 2006, Vol.5, ch.2; Sakai, 1997; USEPA,
5 2006b). Data on collection and treatment rates of MSW in developing countries is
6 difficult to obtain or verify. IPCC (2006, Vol.5, Ch.2) and Eawag (2008) provide
7 guiding information for a few developing countries, which is used to make general
8 assumptions for developing countries. It is assumed that about half of household food
9 waste is centrally collected and deposited to an unmanaged landfill, while the other
10 half ends up in household composts. Household paper waste ends up in an unmanaged
11 landfills and household wood waste is to fifty percent burnt and to fifty percent
12 deposited to an unmanaged landfill. These assumptions are of course very crude and
13 the picture may change with better information about the situation in developing
14 countries.

15 Costs for source separation and treatment of municipal solid waste are taken from
16 various sources. The additional cost of collecting source separated waste compared to
17 mixed waste is derived from Tanskanen (2000) and assumes 33 Euro/t waste for paper
18 waste and 62 Euro/t waste for food and garden waste. The net cost-saving of reducing
19 the amount of waste disposed of and treated in landfills is estimated at 20 Euro/t waste
20 (AEAT, 1998). The cost for upgrading recovered biogas from 60 to 97 percent is
21 taken from Persson (2003). Cost and potential for energy recovery from incineration
22 of waste are taken from IPCC (2006).

23 Costs for treating household and industry food waste in anaerobic digesters for biogas
24 recovery are taken from AEAT (1998, 2001). The digestion process is assumed to
25 convert 60 percent of the original waste amount to biogas. Half of the rest product is
26 assumed to be compost material which can be given away for free, a quarter is a
27 liquor which needs further treatment at 12 Euro/t liquor, and the last quarter (i.e. 10
28 percent of the primary waste amount) is a residual which is landfilled at a cost of 20
29 Euro/t residual (AEAT, 2001).

30 The cost of recycling wood industry waste into chipboards is taken from Wilson
31 (2003).

1 The net cost of household paper recycling is defined as the sum of the additional cost
2 of collecting source separated waste compared to mixed waste (Tanskanen, 2000) and
3 of converting paper waste to recycled pulp (AEAT, 1998) minus the cost-savings of
4 depositing less waste to landfills (AEAT, 1998) and income from selling recycled
5 pulp at a market value (FAOSTAT, 2010). When summing up the different cost items
6 the net unit cost turns out negative, i.e., with these assumptions there appears to be a
7 relatively large net profit from recycling household paper waste. I find two possible
8 reasons for the negative cost estimates for this mitigation measure. First, the market
9 value of recycled pulp may be distorted upwards due to close integration between
10 primary and secondary markets for pulp, i.e. between virgin and recycled pulp, as
11 such integration has been documented several times (OECD, 2007; Ackerman and
12 Gallagher, 2002). Being close substitutes, it is likely that virgin pulp producers have
13 an interest in keeping the price of recycled pulp at a level where the costs for virgin
14 pulp production are covered. Second, costs for separate collection may be higher than
15 assumed in particular in countries currently lacking infrastructure and institutions for
16 mixed collection of household waste. Initial costs for setting these up and educating
17 the public on how to use them are likely to be considerable. Because of difficulties
18 with guessing the non-distorted market value of recycled pulp and lack of information
19 about costs for setting up basic infrastructure for waste collection in developing
20 countries, the cost estimates for the municipal solid waste sector should be treated
21 with caution until further verification is possible.

22 Crude approximations of the amounts of waste generated and the application of
23 different treatment options, in particular for developing countries, contribute to
24 uncertainty in estimated emissions. The effect on emissions from the combined
25 uncertainty for activity data and treatment rates is estimated at ± 50 percent on a global
26 scale.

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1 Table 10: CH₄ mitigation options for solid waste in GAINS. Sources: IPCC (2006); AEAT
 2 (1998); IPCC (2006); Tanskanen (2000); Persson (2003); Wilson (2003).

Sector	Control options in GAINS	Definition of optimal control
MSW - food and garden	Anaerobic digestion w gas recov. and utiliz.	Biodegradable waste currently landfilled is source separated and treated in anaerobic digesters with gas recovery and utilization. Current capacity for composting and mixed incineration remain but without further extensions in the future.
	Household composting	
	Large-scale composting	
	Incineration	
	Landfill with gas recovery and flaring	
	Landfill with gas recovery and utilization	
MSW - paper	Landfill without gas recovery	90 percent of paper waste source separated for recycling and the rest is incinerated. Current capacity for mixed incineration remains but without further extension in the future.
	Paper recycling	
	Incineration	
	Landfill with gas recovery and flaring	
	Landfill with gas recovery and utilization	
MSW - wood	Landfill without gas recovery	Biodegradable waste currently landfilled is source separated and incinerated for energy recovery.
	Incineration	
	Landfill with gas recovery and flaring	
	Landfill with gas recovery and utilization	
Food industry	Landfill without gas recovery	Biodegradable waste is treated in anaerobic digesters with gas recovery and utilization.
	Anaerobic digestion w gas recov. and utiliz.	
	Composting	
	Incineration	
	Landfill with gas recovery and flaring	
	Landfill with gas recovery and utilization	
Pulp and paper industry	Landfill without gas recovery	All waste (black liquor) recovered and incinerated for energy purposes.
	Incineration	
	Landfill with gas recovery and flaring	
	Landfill with gas recovery and utilization	
Textile industry	Landfill without gas recovery	All waste recovered and incinerated for energy purposes.
	Incineration	
	Landfill with gas recovery and flaring	
	Landfill with gas recovery and utilization	
Wood industry	Landfill without gas recovery	All waste max recovered and recycled for wood board production, residuals incinerated for energy purposes.
	Recycling for board production	
	Incineration	
	Landfill with gas recovery and flaring	
	Landfill with gas recovery and utilization	

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1 2.11 Wastewater

2 Wastewater treatment plants serve to decompose compounds containing nitrogen and
3 phosphor as well as carbon from the wastewater before discharge. Main gaseous
4 products are CO₂ and molecular nitrogen, but during the process also CH₄ is released.
5 CH₄ is formed whenever wastewater with high organic content is handled under
6 anaerobic conditions.

7 In the GAINS model, domestic and industrial wastewater are accounted for
8 separately. Domestic wastewater is split into centralized and decentralized collection,
9 basically referring to wastewater from urban and rural population, except for
10 industrialized countries where wastewater collection services often include some rural
11 areas as well. Fractions of wastewater collected centrally are taken from UNFCCC
12 (2010), FAO (2009) and UNstat (2010).

13 Uncontrolled emissions are defined as emissions when wastewater is emitted directly
14 to a water body without prior collection and treatment. As anaerobic conditions are
15 formed when large quantities of wastewater are collected and stored, CH₄ formation in
16 the uncontrolled case are likely to be very limited and to increase for any form of
17 organized wastewater collection. Collection is however a prerequisite for treatment,
18 which is important for combating water pollution from excessive nitrogen and
19 phosphor. Uncontrolled CH₄ emission factors are derived following the IPCC
20 guidelines (2006, Vol.5, Equations 6.1 to 6.3):

$$21 \quad E_{it} = \sum_s A_{it} * h_s * ef_i * (1 - remeff_m) * Appl_{itm} \quad (20)$$

22 where A_{it} is total population in country i and year t ,

23 h_s is fraction of total population connected to treatment system s
24 (centralized or decentralized treatment)

25 $remeff_m$ is the removal efficiency of technology m ,

26 $Appl_{itm}$ is the application of technology m in country i and year t , and

$$27 \quad ef_i = BOD_i * B_0 * MCF_0, \quad (21)$$

28 where BOD is amount of biochemical oxygen demand per person in country
29 i ,

30 B_0 is maximum CH₄ producing capacity,

1 MCF_0 is the methane correction factor, i.e. the fraction of BOD
 2 converted to CH₄.

3 Country-specific IPCC (2006, Vol.5, Table 6.4) default factors for the amount of
 4 biochemical oxygen demand per person (BOD) are used with a range from 12.4 to
 5 31.0 kt CH₄/M people. A IPCC (2006, Vol.5, Table 6.2) default factor of 0.6 kt
 6 CH₄/kt BOD is used for the maximum CH₄ producing capacity (B_0). Methane
 7 correction factors (MCF_0) of 0.1 for uncontrolled decentralized collection and 0.5 for
 8 uncontrolled centralized collection apply.

9 Industry sectors identified by IPCC (2006, Vol.5, p.6.19) as potential sources for
 10 wastewater CH₄ emissions are food, organic chemical, and pulp- and paper industry.
 11 Emissions are calculated as:

$$12 \quad E_{it} = \sum_{sm} A_{its} * COD_i * ef_i * (1 - remeff_m) * Appl_{itm} \quad (22)$$

13 where A_{its} is the amount of product A produced in country i in year t ,
 14 COD_i is the chemical oxygen demand in untreated wastewater
 15 generated per tonne product produced in country i ,
 16 $remeff_m$ is the removal efficiency of technology m ,
 17 $Appl_{itm}$ is the application of technology m in country i and year t , and
 18 where

$$19 \quad ef_i = B_0^{COD} * MCF_0, \quad (23)$$

20 where B_0^{COD} is maximum CH₄ producing capacity,
 21 MCF_0 is the methane correction factor, i.e., the fraction of CH₄
 22 generated which is not oxidised but released as CH₄.

23
 24 A IPCC (2006, Vol.5, Table 6.2) default factor of 0.25 kt CH₄/kt COD is applied for
 25 the maximum CH₄ producing capacity (B_0^{COD}). A methane correction factor (MCF_0)
 26 of 0.5 is applied for the uncontrolled case.

27 Activity data is the amount of COD in untreated industrial wastewater. These amounts
 28 are derived from production volumes combined with COD generation factors as

1 specified in Table 11. Production volumes in tonne product are taken from FAOSTAT
2 (2011), USDA (2011), and EC (2003). Future production volumes are projected
3 proportional to growth in value added in each industry sector (food, organic chemical,
4 and pulp- and paper). For pulp- and paper industry, wastewater and COD generation
5 rates reported in literature differ considerably between processes and between
6 developed and developing countries. By comparing reported values from different
7 sources, process specific generation rates are derived as presented in Table 11. It
8 should be noted that when using process specific generation rates, the estimated
9 amounts of COD and CH₄ generated from this industry come out several times lower
10 than if using the IPCC default factor (2006, Vol.5, Table 6.9) for pulp- and paper
11 industry, which amounts to 162 m³ wastewater per ton product and 9 kg COD per m³
12 wastewater.

13 There are no wastewater options available that primarily target CH₄ emissions. There
14 are, however, several different ways of treating wastewater, which have different
15 implications for CH₄ emissions (Pohkrel and Viraraghavan, 2004 and Thompson et
16 al., 2001). When domestic wastewater is centrally collected and emitted to a water
17 body with only mechanical treatment to remove larger solids, plenty of opportunities
18 for anaerobic conditions and CH₄ formation are created. With well managed aerobic
19 or anaerobic treatment, the CH₄ formation is effectively mitigated and CH₄ emissions
20 can be kept on a negligible level. With less well managed systems the occurrence of
21 anaerobic conditions increase as well as CH₄ formation (IPCC 2006, Vol.5, Tables 6.3
22 and 6.8). Anaerobic treatment has advantages over aerobic treatment like lower costs,
23 smaller volumes of excess sludge produced, and the possibility of recovering useful
24 biogas, which can be upgraded to gas grid quality (Lettinga 1995, Thompson et al.
25 2001). For industrial wastewater, it is assumed that the most effective way to reduce
26 CH₄ emissions is to apply a two-stage process where the water is treated anaerobically
27 with recovery of the biogas in a first stage, which is then followed by an aerobic
28 treatment in a second stage (Latorre et al., 2007). In rural areas, domestic wastewater
29 can be collected and treated in latrines, septic tanks or similar anaerobic treatment
30 (USEPA, 1999).

31 Current applications of different treatment practices for domestic and industrial
32 wastewater are taken from UNFCCC (2009) CRF tables and NIRs for Annex 1 and
33 non-Annex 1 countries complemented with information from FAO (2009) wastewater

1 database, IPCC (2006, Vol.5, Table 6.5), USDC (2005), and Pokhrel and
2 Viraraghavan (2004). Investment costs for sewage treatment are taken from EEA
3 (2005) and operation and maintenance costs from Hernandez-Sancho and Sala-
4 Garrido (2008). Rural wastewater treatment costs are from USEPA (1999).

5 IPCC (2006, Vol. 5, p.6.17) estimates the uncertainty range for both maximum CH₄
6 producing capacity (B₀) and BOD per person to ± 30 percent.

7 A major uncertainty source in the estimation of CH₄ emissions from wastewater are
8 amounts and COD content in industry wastewater. Reported data is rare and anecdotal
9 and the water efficiency in industry is likely to fluctuate widely with the chosen
10 production technology as well as access to quality water and local water price. IPCC
11 (2006, Vol. 5, p.6.23) provides a general uncertainty range for amount of COD per ton
12 product of -50 to +100 percent, which is applied here.

1 Table 11: Assumptions for estimation of amounts of pre-treatment COD (chemical oxygen demand) in industry wastewater.

Industry	Product/production process	Wastewater generation		COD content pre-treatment		COD concentration in untreated wastewater		Sources
		developed countries	developing/transitional	developed countries	developing/transitional	developed countries	developing/transitional	
		m3/t dry product		kg COD/ton dry product		kg COD/m3 wwater		
Food	Beer of barley	6.3	6.3	18.3	18.3	2.9	2.9	IPCC (2006, Vol.5, Table 6.9)
	Vegetable oils	3.1	3.1	3.1	3.1	1	1	
	Wine	23	23	34.5	34.5	1.5	1.5	
	Sugar (centrifugal)	9	9	28.8	28.8	3.2	3.2	
	Meat (total)	13	13	53.3	53.3	4.1	4.1	
	Milk (total)	7	7	18.9	18.9	2.7	2.7	
Organic chemical	Organic chemicals ^a	9 m3/ Euro value added	9 m3/ Euro value added	27 kg COD/ Euro value added	27 kg COD/ Euro value added	3	3	EUROSTAT (2005), IPCC (2006, Vol.5, Table 6.9)
Pulp	Bleached sulphate pulp	70	200	150	500 ^b	2.14	2.50	Judd and Jefferson (2003), Pokhrel and Viraraghavan (2004), Khansorthong and Hunsom (2009), Buzzini and Pires (2002), Thompson et al. (2001), Latorre et al. (2007)
	Unbleached sulphate pulp	50	100	60	200 ^b	1.20	2.00	
	Bleached sulphite pulp	150	200	160	500 ^b	1.07	2.50	
	Unbleached sulphite pulp	70	200	100	300 ^b	1.43	1.50	
	Mechanical wood pulp	20	100	60	300 ^b	3.00	3.00	
	Semi-chemical wood pulp	30	100	140	600 ^b	4.67	6.00	
	Other fibre pulp	20	100	60	300 ^b	3.00	3.00	
	Recovered pulp	20	100	60	300 ^b	3.00	3.00	
Paper	Newsprint	25	50	1.5	5	0.06	0.10	
	Printing&writing paper	29	60	1.5	5	0.05	0.08	
	Recovered paper	20	40	1	5	0.05	0.13	
	Household/sanitary/tissue	15	30	1.5	5	0.10	0.17	
	Wrapping papers	20	40	1.5	5	0.08	0.13	
	Paper&paperboard-other	8	20	1.5	5	0.19	0.25	

^a Production volumes not available. Amounts refer to average wastewater generation rates for organic chemical industry in Europe (EUROSTAT, 2005)

^bCOD content in effluent wastewater when recovered black liquor has been removed, i.e. COD generated in the digester house not included.

1 References

- 2 Ackerman, F. and K. Gallagher: Mixed signals: market incentives, recycling, and the price spike of 1995,
3 Resources, Conservation and Recycling, Vol.35, pp.275-295, 2002.
- 4 ADB: ALGAS -Asia least cost greenhouse gas abatement strategy -People's Republic of China, Asia
5 Development Bank, Manila, 1998.
- 6 AEAT: Options to reduce methane emissions –A Report produced for DGXI, European Commission, AEA
7 Technology, UK, 1998.
- 8 AEAT: Economic evaluation of emission reductions of nitrous oxide and methane in agriculture in the EU,
9 Report prepared for DG Environment, European Commission, AEA Technology, UK, 2001.
- 10 An, B. X., T. R. Preston and F. Dolberg: The introduction of low-cost polyethylene tube biodigesters on small
11 scale farms in Vietnam, Livestock Research for Rural Development, Vol.9 (2), 1997.
- 12 Barrett, C. B., C. M. Moser, O. V. McHugh and J. Barison: Better technology, better plots, better farmers?
13 Identifying changes in productivity and risk among Malagasy rice farmers, American Journal of Agricultural
14 Economics, Vol.86 (4), pp.869-888, 2004.
- 15 Boadi, D., C. Benchaar, J. Chiquette and D. Massé: Mitigation strategies to reduce enteric methane emissions
16 from dairy cows: Update review, Canadian Journal of Animal Science, Vol.84, pp.319-335, 2004.
- 17 Bouman, B.A.M.: Water-efficient management strategies in rice production”, IRRI Mini Review 26.2,
18 International Rice Research Institute, Los Banos, Philippines, 2001.
- 19 Buzzini, A.P. and E.C. Pires: Cellulose pulp mill effluent treatment in an upflow anaerobic sludge blanket
20 reactor, Process Biochemistry, Vol.38, pp.707-713, 2002.
- 21 Buzcu-Guven, B., R. Harriss and D. Hertzmark: Gas flaring and venting: extent, impacts and remedies, Energy
22 Forum, The James A. Baker III Institute for Public Policy, Rice University, Houston, 2010.
- 23 CAPRI model: Common Agricultural Policy Regional Impact Analysis Model, Bonn University, Bonn, 2009.
- 24 China University of Petroleum: Feasibility Study of Coal Bed Methane Production in China, EU-China Energy
25 and Environment Programme, EuropeAid/120723/D/SV/CN, Beijing, March 2008.
- 26 CIA World Factbook: US Central Intelligence Agency, Washinton D.C., 2010.
- 27 Delmas, R.: An overview of present knowledge on methane emission from biomass burning, Fertilizer Research,
28 Vol.37, pp.181-190, 1994.
- 29 Dennet, J. and S. Vallender: Reducing fugitive emissions from gas distribution systems by the systematic
30 application of pressure profiling technology, National Grid, Warwick, UK, 2011.
- 31 Dhingra, R., E. R. Christensen, Y. Liu, B. Zhong, C. Wu, M. G. Yost and J. V. Remais: Greenhouse gas
32 emission reductions from domestic anaerobic digesters linked with sustainable sanitation in rural China”,
33 Environmental Science and Technology Vol.45, pp.2345-2352, 2011.
- 34 Eawag: Global waste challenge, Swiss Federal Institute for Aquatic and Science and Technology, Dübendorf,
35 Switzerland, 2008.
- 36 EC: Council Directive 1999/31/EC of 26 April 1999 on the landfill of waste, European Commission, Brussels,
37 1999.
- 38 EC: Sugar -international analysis, production structures within the EU, European Commission, Brussels, 2003.
- 39 EC: Directive 2006/12/EC of the European Parliament and of the Council of 5 April 2006
40 on waste, European Commission, Brussels, 2006.
- 41 EC: Decision No 406/2009/EC of the European Parliament and of the Council of 23 April 2009 on the effort of
42 Member States to reduce their greenhouse gas emissions to meet the Community’s greenhouse gas emission
43 reduction commitments up to 2020, European Commission, Brussels, 2009.
- 44 ECCP: Mitigation potential of greenhouse gases in the agricultural sector –Final report by working group 7,
45 European Climate Change Programme, European Commission, Brussels, 2003.

1 EEA: Effectiveness of urban wastewater treatment policies in selected countries: an EEA pilot study, European
2 Environment Agency, Copenhagen, 2005.

3 Ecofys: Emission reduction potential and costs for methane and nitrous oxide in the EU-15, Report for DGXI,
4 European Commission, Ecofys, Utrecht, June 1998.

5 Ecofys: Methane and nitrous oxide, Sectoral emission reduction potentials and economic costs for climate
6 change policies SERPEC-CC, Report for DG Environment, European Commission, Ecofys, Utrecht, 2009.

7 EIA: Natural Gas Compressor Stations on the Interstate Pipeline Network: Developments since 1996, US Energy
8 Information Administration, Office of Oil and Gas, Washington D. C., November 2007.

9 EIA: various internet pages, e.g., International Energy Statistics and Country Analysis Briefs,
10 <http://www.eia.doe.gov/>, US Energy Information Administration, Washington D. C., 2010a.

11 EIA: Natural Gas Pipeline Capacity & Utilization, <http://www.eia.doe.gov/>, US Energy Information
12 Administration, Washington D. C., 2010b.

13 EIA: Coal production, selected years 1949-2009, <http://www.eia.doe.gov/>, US Energy Information
14 Administration, Washington D. C., 2010c.

15 EIA: various internet pages, e.g., International Energy Statistics and Country Analysis Briefs,
16 <http://www.eia.doe.gov/>, US Energy Information Administration, Washington D. C., 2011.

17 ERCB: Upstream petroleum industry flaring and venting report, Energy Resources Conservation Board, Canada,
18 November 2010.

19 EUROSTAT: <http://epp.eurostat.ec.europa.eu/>, European Commission, Brussels, 2005.

20 EUROSTAT: <http://epp.eurostat.ec.europa.eu/>, European Commission, Brussels, 2008.

21 EUROSTAT: <http://epp.eurostat.ec.europa.eu/>, European Commission, Brussels, 2009.

22 FAO: World agriculture: towards 2015/2030 –Summary report, Food and Agriculture Organization of the United
23 Nations, Rome, 2003.

24 FAO: Water charging in irrigated agriculture –an analysis of international experience, FAO water reports 28,
25 Food and Agriculture Organization, Rome, 2004.

26 FAO: Wastewater database, Food and Agriculture Organization, Rome, 2009.

27 FAOSTAT: <http://faostat.fao.org/>, Food and Agriculture Organization, Rome, 2010.

28 FAOSTAT: <http://faostat.fao.org/>, Food and Agriculture Organization, Rome, 2011.

29 Ferrero, A. and N. V. Nguyen: Constraints and opportunities for the sustainable development of rice-based
30 production systems in Europe in . N. V. Nguyen (ed.) Proceedings of the FAO Rice Conference, Food and
31 Agriculture Organization, Rome, 2004.

32 GGFR: Using Russia’s associated gas, Report prepared by PFC Energy for the Global Gas Flaring Reduction
33 Partnership and the World Bank, 2007.

34 GMI: VAM Utilization Project at Xiaodongshan Shaft of Sihe Mine, Jincheng Anthracite Mining Group,
35 Jincheng Mining Area, Shanxi Province, China, Global Methane Initiative, Washington D. C., 2008.

36 GMI: Global overview of CMM opportunities 2009, Coalbed Methane Outreach Program of the Global Methane
37 Initiative, Washington D. C, 2009.

38 GMI: CMM Country Profiles: China, Global Methane Initiative, Washington D.C., 2011.

39 Greene, W.: LIMDEP Version 7.0, Econometric Software, Inc., New York, 2005.

40 Hernandez-Sancho, F. and R. Sala-Garrido: Cost modelling in waste water treatment processes: an empirical
41 analysis for Spain, in P. Hlavinek, O. Bonacci, J. Marsalek and I. Mahrikova (eds.) Dangerous Pollutants
42 (Xenobiotics) in urban water cycle, Springer, Dordrecht, 2008.

43 Heytens, P.: Chapter 6: Technical change in wetland rice agriculture, in S. Pearson, W.Falcon, P. Heytens, E.
44 Monke and R. Naylor (eds.) Rice Policy in Indonesia, Cornell University Press, Ithaca and London, 1991.

45 Howarth, R. W., R. Santoro and A. Ingraffea: Methane and the greenhouse-gas footprint of natural gas from
46 shale formations”, Climatic Change Letter, DOI 10.1007/s10584-011-0061-5, 2011.

- 1 Hsiao, C.: Analysis of Panel Data, Econometric Society Monographs No. 11, Cambridge University Press,
2 Cambridge, 1986.
- 3 Hulbak Røland T.: Associated petroleum gas in Russia –Reasons for non-utilization, FNI Report 13/2010,
4 Fridtjof Nansen Institute, Lysaker, 2010.
- 5 IEA-WEO: IEA World Energy Outlook 2009, International Energy Agency, Paris, 2009.
- 6 IEA: Natural Gas Information 2009, IEA Statistics, International Energy Agency, Paris, 2010.
- 7 IMF: International Financial Statistics, International Monetary Fund, Washington D.C, 2006.
- 8 IPCC: Revised 1996 IPCC Guidelines for National Greenhouse Gas Inventories, Intergovernmental Panel on
9 Climate Change, UK, 1997.
- 10 IPCC: 2006 IPCC Guidelines for National Greenhouse Gas Inventories, Intergovernmental Panel on Climate
11 Change, Japan, 2006.
- 12 IPPC: Reference document on the best available techniques for waste Incineration, Integrated Pollution
13 Prevention and Control, European Commission Joint Research Centre, Seville, August 2006.
- 14 IRRI: Distribution of rice crop area by environment 2004-2006, International Rice Research Institute, Los
15 Banos, the Philippines, 2007.
- 16 Iqbal, M. F., Y-F. Cheng, W-Y. Zhu, B. Zeshan: Mitigation of ruminant methane production: current strategies,
17 constraints and future options, World Journal of Microbiology and Biotechnology, Vol.24, pp.2747-2755, 2008.
- 18 Johansson, L. S., B. Leckner, L. Gustavsson, D. Cooper, C. Tullin, A. Potter: Emission characteristics of
19 modern and old-type residential boilers fired with wood logs and wood pellets, Atmospheric pollution, Vol. 38,
20 pp.4183-4195, 2004.
- 21 Johnson, M. R. and A. R. Coderre: An analysis of flaring and venting activity in the Alberta upstream oil and gas
22 industry, Journal of Air and Waste Management, Vol.61, pp.190-200, 2011.
- 23 Johnson, M. R. and L. W. Kostiuk: A parametric model for the efficiency of a flare in crosswind”, Proceedings
24 of the Combustion Institute, Vol.29, pp.1943-1950, 2002.
- 25 Judd, S. and B. Jefferson: Membranes for Industrial Wastewater Recovery and Re-use, Elsevier, UK, 2003.
- 26 Khansorthong, S. and M. Hunsom: Remediation of wastewater from pulp and paper mill industry by the
27 electrochemical technique, Chemical Engineering Journal, Vol.151, pp.228-234, 2009.
- 28 Kirschgessner, D. A., R. A. Lott, R. M. Cowgill, M. R. Harrison, T. M. Shires: Estimate of methane emissions
29 from the U.S. natural gas industry, Chemosphere, Vol. 35, pp. 1365-1390, 1997.
- 30 Kjällstrand, J. and M. Olsson: Chimney emissions from small-scale burning of pellets and fuelwood –examples
31 referring to different combustion appliances, Biomass&Bioenergy, Vol.27, pp.557-561, 2004.
- 32 Klimont, Z. and C. Brink: The RAINS model update of the ammonia module and introduction of agricultural
33 greenhouse gases –Final report to the Royal Ministry of Environment, Norway, 2003.
- 34 Latorre, A., A. Malmqvist, S. Lacorte, T. Welander, D. Barcelo: Evaluation of the treatment efficiencies of paper
35 mill whitewaters in terms of organic composition and toxicity, Environmental Pollution, Vol.147, pp.648-655,
36 2007.
- 37 Lechtenböhmer, S., C. Dienst, M. Fishedick, T. Hanke, R. Fernandez, D. Robinson, R. Kantamaneni, B. Gillis:
38 Tapping the leakages: methane losses, mitigation options and policy issues for Russian long distance gas
39 transmission pipelines, International Journal of Greenhouse Gas Control, Vol.1, pp.387-395, 2007.
- 40 Lelieveld, J., S. Lechtenböhmer, S.S. Assonov, C. A. M. Brenninkmeijer, C. Dienst, M. Fishedick and T.
41 Hanke: Low methane leakage from gas pipelines, Nature, Vol.434, pp.841-842, 2005.
- 42 Lettinga, G.: Anaerobic digestion and wastewater treatment systems, International Journal of General and
43 Molecular Microbiology, Vol.67, pp.3-28, 1995.
- 44 Lovett, D. K., L. Shalloo, P. Dillon and F. P. O'Mara: A systems approach to quantify greenhouse gas fluxes
45 from pastoral dairy production as affected by management regime, Agricultural Systems, Vol. 88, pp.156-179,
46 2006.

- 1 Mattus, R. and Å. Källstrand: Chapter 12: Fossil Energy and Ventilation Air Methane, in Reay, D. P. Smith and
2 A. van Amstel (Eds.) Methane and Climate Change, Earthscan, London, 2010.
- 3 MNRE: Annual report: 2005–2006, Ministry of New and Renewable Energy, Government of India, New Delhi,
4 2006.
- 5 Newbold, C. J., S. Lopez, N. Nelson, J. O. Ouda, R. J. Wallace, A. R. Moss: Propionate precursors and other
6 metabolic intermediates as possible alternative electron acceptors to methanogenesis in ruminal fermentation,
7 British Journal of Nutrition, Vol.94, pp.27-35, 2005.
- 8 Niemi, J.V.: Atmospheric emissions from open biomass burning –development of datasets for RAINS model,
9 IIASA Interim Report IR-05-007, International Institute for Applied Systems Analysis (IIASA), Laxenburg,
10 2006.
- 11 NOAA: Global Gas Flaring Estimates Database, National Geophysical Data Centre, Boulder, 2010.
- 12 OECD: Improving recycling markets, OECD publishing, Paris, 2006.
- 13 Olsson, M. and J. Kjällstrand: Low emissions from wood burning in an ecolabelled residential boiler”,
14 Atmospheric Environment, Vol.40, pp.1148-1158, 2006.
- 15 OME: Assessment of internal and external gas supply options for the EU, evaluation of the supply costs of new
16 natural gas supply projects to the EU and an investigation of related financial requirements”, Observatoire
17 Mediterranee de l’Energie, Nanterre, 2001.
- 18 Papar, R., A. Szady, W. D. Huffer, V. Martin, A. McKane: Increasing energy efficiency in mine ventilation
19 systems, Industrial Energy Analysis, Lawrence Berkeley National Laboratory, University of California, 1999.
- 20 Persson, M.: *Utvärdering av uppgraderingstekniker för biogas* (in Swedish), Evaluation of upgrading techniques
21 for biogas, SGC Rapport 142, Swedish Centre for Gas Technology, Malmö, 2003.
- 22 PFC Energy: Using Russia’s associated gas, Report prepared for the Global Gas Flaring Reduction Partnership
23 and the World Bank, Washington D.C, December 2007.
- 24 Pokhrel, D. and T. Viraraghavan: Treatment of pulp and paper mill wastewater –a review, Science of the Total
25 Environment, Vol.333, pp.37-58, 2004.
- 26 Sakai, S.: Municipal solid waste management in Japan, Waste Management, Vol. 16, pp.395-405, 1997.
- 27 Sino-US New Energy Sci-Tech Forum: Summary Report –Conference on Coalmine Methane Recovery and
28 Utilization, Jincheng, China, February 24-27, 2009.
- 29 Shibayama, H.: Weeds and weed management in rice production in Japan, Weed biology and management, Vol.
30 1, pp. 53-60, 2001.
- 31 SPP: Slovak Gas Works Annual Report 2002-2006, Slovenský plynárenský priemysel, a.s., Bratislava, 2007.
- 32 TAG: Nominal, committed, available and used transportation capacities, www.taggmbh.at, Trans Austria Gas
33 (TAG) GmbH, Austria, 2007.
- 34 Tanskanen, J.-H.: Strategic planning of municipal waste management, Resources, Conservation and Recycling,
35 Vol.30, pp.111-133, 2000.
- 36 Thakur, P. C.: Coal seam degasification, in Kissell, F.N. (ed.) Handbook for Methane Control in Mining,
37 Information Circular 9486, Department of Health and Human Services, National Institute for Occupational
38 Safety and Health, Pittsburgh, US, 2006.
- 39 Thompson G., J. Swain, M. Kay and C.F. Forster: The treatment of pulp and paper mill effluent: a review,
40 Bioresource Technology, Vol.77, pp.275-286.
- 41 UNstat: Population database, United Nations Department of Economic and Social Affairs, New York, 2009.
- 42 UNstat: Environmental indicators database –water, United Nations Department of Economic and Social Affairs,
43 New York, 2010.
- 44 UNFCCC: South Africa National Inventory Report to the United Nations Framework Convention on Climate
45 Change, Bonn, 2000.
- 46 UNFCCC: India National Inventory Report to the United Nations Framework Convention on Climate Change,
47 Bonn, 2004.

1 UNFCCC: Common Reporting Format (CRF) tables and National Inventory Reports (NIRs) (version
2 downloaded in July 2009), United Nations Framework Convention on Climate Change, Bonn, 2009.

3 UNFCCC: Common Reporting Format (CRF) tables and National Inventory Reports (NIRs) (version
4 downloaded in April 2010), United Nations Framework Convention on Climate Change, Bonn, 2010.

5 UNFCCC: United States Common Reporting Format to the United Nations Framework Convention on Climate
6 Change, Bonn, 2011.

7 UNIDO: Industrial Statistics Database, 3-digits level of ISIC Code, Revision 2., United Nations Industrial
8 Development Organization, Vienna, 2006.

9 Unruh, B.: Delivered energy consumption projections by industry in the Annual Energy Outlook 2002, US
10 Energy Information Administration, Washington D. C., 2002.

11 USDA: Sugar: world markets and trade, database, United States Department of Agriculture, Washington D. C.,
12 2011.

13 USDC: Water supply and wastewater treatment market in China, US Department of Commerce, International
14 Trade Administration, Washington D.C., 2005.

15 USEPA: Decentralized systems technology fact sheet –septic tank –soil absorption systems, EPA 932-F-99-075,
16 US Environmental Protection Agency, Washington D.C., 1999.

17 USEPA: Assessment of the worldwide market potential for oxidizing coal mine ventilation air methane, EPA
18 430-R-03-002, US Environmental Protection Agency, Washington D. C., July 2003.

19 USEPA: Inventory of U.S. Greenhouse gas emissions and sinks: 1990-2004, US Environmental Protection
20 Agency, Washington D.C, 2006a.

21 USEPA: Municipal solid waste generation, recycling, and disposal in the United States: Facts and figures 2006 ,
22 US Environmental Protection Agency, Washington D.C, 2006b.

23 USEPA: US surface coal mine methane recovery project opportunities”, EPA Publication 430R08001, US
24 Environmental Protection Agency, Washington D. C., July 2008.

25 USEPA: Coalbed methane outreach program, <http://www.epa.gov/cmop/>, US Environmental Protection Agency,
26 Washington D.C, 2010.

27 USEPA: Natural Gas STAR Program, <http://www.epa.gov/gasstar/>, US Environmental Protection Agency,
28 Washington D.C, 2011a.

29 USEPA: DRAFT: Global Anthropogenic Non-CO2 Greenhouse Gas Emissions: 1990-2030, EPA 430-D-11-003,
30 August 2011, US Environmental Protection Agency, Washington D.C., 2011b.

31 Van der Gon, H. A. D., P. M. Van Bodegom, R. Wassmann, R. S. Lantin and T. M. Metra-Corton: Sulphate-
32 containing amendments to reduce methane emissions from rice fields: mechanisms, effectiveness and costs,
33 Mitigation and Adaptation Strategies for Global Change, Vol.6, pp.71-89, 2001.

34 Wilson, B.: Using wood for chipboard, Proceedings from workshop on Timber Recycling Opportunities in the
35 Midlands, 28 April 2003, Aston University, Birmingham, UK, 2003.

36 Winiwarter, W. and K. Rypdal: Assessing the uncertainty associated with national greenhouse gas emission
37 inventories: A case study for Austria, Atmospheric Environment 35, 5425-5440, 2001.

38 Winiwarter, W. and B. Muik: Statistical dependences in input data of national greenhouse gas inventories:
39 Effects on the overall inventory uncertainty, Climatic Change 103 (1-2), 19-36, 2010.

40 Wright, A. D. G., P. Kennedy, C. J. O’Neill, A. F. Toovey, S. Popovski, S. M. Rea, C. L. Pimm, and L. Klein:
41 Reducing methane emissions in sheep by immunization against rumen methanogens, Vaccine, Vol.22, pp.3976-
42 3985, 2004.

43 Wuppertal Institute: Greenhouse gas emissions from Russian natural gas export pipeline system, Final Report,
44 Wuppertal Institute for Climate, Environment and Energy in cooperation with Max-Planck-Institute for
45 Chemistry, Wuppertal and Mainz, 2005.

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