

SNAP CODE:	112201
	112202
	112203
	112204
	112205

SOURCE ACTIVITY TITLE:	OTHER SOURCES AND SINKS:
	FOREST AND GRASSLAND CONVERSION
	<i>Tropical Forests</i>
	<i>Temperate Forests</i>
	<i>Boreal Forests</i>
	<i>Grassland</i>
	<i>Other</i>

NOSE CODE:	301.22.01
	301.22.02
	301.22.03
	301.22.04
	301.22.05

NFR CODE:	5 B
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A specific methodology for these activities has not been prepared yet as this is a new sub group.

The expert panel leaders for this sub group are listed below.

These activities are not believed to be a significant source of PM_{2.5} (as of December 2006).

Updated with particulate matter details by:

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December 2006

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SNAP CODE:	112301
	112302
	112303
	112304
	112305

SOURCE ACTIVITY TITLE:	OTHER SOURCES AND SINKS:
	ABANDONMENT OF MANAGED LANDS
	<i>Tropical Forests</i>
	<i>Temperate Forests</i>
	<i>Boreal Forests</i>
	<i>Grassland</i>
	<i>Other</i>

NOSE CODE:	301.23.01
	301.23.02
	301.23.03
	301.23.04
	301.23.05

NFR CODE:	N/A
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SNAP CODE: 112400

SOURCE ACTIVITY TITLE: OTHER SOURCES AND SINKS
CO₂ Emissions From/or Removal into Soils

NOSE CODE:

NFR CODE: N/A

A specific methodology for these activities has not been prepared yet as this is a new sub group.

The expert panel leaders for this sub group are listed below.

These activities are not believed to be a significant source of PM_{2.5} (as of December 2006).

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SNAP CODE: 112500

SOURCE ACTIVITY TITLE: OTHER SOURCES AND SINKS
Other

NOSE CODE:

NFR CODE: 5 E

A specific methodology for these activities has not been prepared yet as this is a new sub group.

The expert panel leaders for this sub group are listed below.

These activities are not believed to be a significant source of PM_{2,5} (as of December 2006).

Updated with particulate matter details by:

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SNAP CODE:	1101
	1102
	1111
	1112

SOURCE ACTIVITY TITLE:	OTHER SOURCES AND SINKS
	<i>Non-managed deciduous forests</i>
	Non-managed coniferous forests
	<i>Managed deciduous forests</i>
	<i>Managed coniferous forests</i>

NOSE CODE:	11.01.04, 11.01.05, 301.01.(06-11), 301.01.(15-17)
	301.02.(04-12), 301.02.(15-16)
	301.11.(04-11), 301.11.(15-17)
	301.12.(04-12), 301.12.(15-16)

NFR CODE:	5 E
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1 ACTIVITIES INCLUDED

All types of foliar forest emissions will be considered, non-managed and managed, deciduous and coniferous. Forest foliage is primarily a source of VOC, and we distinguish here between isoprene, monoterpenes, and 'other VOC'. Emissions from forest soils are covered in the chapter dealing with activities 110117, 110216, 111117 and 111216. Note that for methane the flux is believed to be from the atmosphere to the forest floor, so in any case a zero emission factor is recommended for this species. Emissions from forest fires are covered in 1103 "Forest and other vegetation fires". Emissions from the forest undergrowth and root system have not yet been included, although may be added at a later stage. Emissions from shrub-like vegetation, maquis, garrique, or other vegetation types are covered in SNAP 1104 "Natural grassland and other vegetation".

2 CONTRIBUTIONS TO TOTAL EMISSIONS

According to CORINAIR-90, forests (deciduous and coniferous) contributed 19% of total European NMVOC emissions, 4.4% of CH₄ emissions, 14.3% of N₂O emissions and 0.8% of NH₃ emissions.

However, as noted below the natural emission estimates for VOC as supplied for CORINAIR-1990 must now be regarded as outdated. Table 2.1 compares the more recent estimate of Simpson et al. (1995) and Guenther et al. (1995) with estimates of anthropogenic emissions.

Table 2.1: Comparison of estimated isoprene, OVOC, and monoterpene emissions from forests with man-made VOC. Units: ktonnes per year

Country	Isoprene	OVOC	Monoterp.	Man-made VOC**
Albania	6	9	16	30
Austria	32	78	30	418
Belgium	30	13	7	364
Bulgarian	135	44	41	178
Czechoslov.***	70	95	124	
Denmark	7	7	4	167
Finland	82	354	398	209
France	480	216	215	2393
Germany	121	190	249	3154
Greece	21	35	62	293
Hungary	82	16	23	205
Iceland	0	0	7	6
Ireland	2	6	11	102
Italy	53	89	142	2080
Luxembourg	2	1	0.4	19
Netherlands	8	6	5	424
Norway	29	104	143	266
Poland	63	176	113	802
Portugal	36	61	70	202
Romania	154	83	55	567
Spain	137	248	272	1050
Sweden	108	389	370	528
Switzerland	5	17	30	284
Turkey	213	460	175	
Russia	2006*	3197*	1060-3490[I]	3566
UK	53	27	39	2287
Sum	4000	6000	3700-6100	20000

Notes: All isoprene and OVOC emissions are from Simpson et al., 1995.

All monoterpene emissions (except Russia) are from Guenther et al., 1995, in ktonne carbon. * 1989 estimates were made for whole Soviet Union, however, Russia is expected to account for the majority of emissions.

Man-made emissions are unofficial estimates, generally derived by subtracting estimated Natural and Agricultural emissions from total emissions. * Former Czechoslovakia

Other references: I - Isidorov, 1992, sum of pine+fir emissions

These activities are not believed to be a significant source of PM_{2.5} (as of December 2006).

3 GENERAL

3.1 Description

The subject of emission inventories for emissions from vegetation is still very much in its infancy in Europe, and the design of an inventory procedure should reflect this. Indeed, NMVOC inventories prepared for the CORINAIR 1990 data-base have already been outdated by recent re-evaluations of the emission factors on which these have been based (Guenther et al., 1993, 1995, 1998, Simpson et al., 1995, Seufert et al., 1997). Generally, the mix of emissions varies greatly both in and between vegetation types, and knowledge of this mix is constantly being updated and in some cases completely revised.

*** It is more important at this stage to assemble the land-use information than to estimate the emissions directly. ***

This is especially true for NMVOC as models are almost completely reliant on good land-use databases for their biogenic emissions estimates. Hopefully the procedures suggested here will lead to a Europe-wide database, which will greatly improve emission estimates for model calculations and policy decisions.

Biogenic VOC is also a rather loose term for a wide range of compounds, of which only a few are generally of most interest. Isoprene is generally the compound of most importance for ozone modelling for example, and it is useful to inventory this compound specifically. Emissions of the various terpenes may also be important, although there are great uncertainties associated with their atmospheric behaviour. Similarly, the remaining VOC species ('other VOC', or OVOC) doubtless play some role in atmospheric chemistry problems, but little is known about the chemistry of many components or the quantitative emissions of individual species. Emissions may be large, however.

A review of the sources and chemistry of biogenic VOCs has recently been given by Fehsenfeld et al., 1992. A special-issue Atmospheric Environment dealing with a large number of European measurements has recently been published (Seufert et al., 1997).

Emissions vary greatly from one tree species to another. And as knowledge has increased some species previously classified as non-isoprene emitters have actually now been found to emit isoprene in significant quantities. Conversely, oaks were previously thought to be always high isoprene emitters, whereas now it is recognised that some evergreen oaks emit little isoprene but very high amounts of terpenes (Seufert et al., 1997).

These considerations have been reflected in the new SNAP codes adopted for this chapter, which assign codes to specific types of trees, rather than to "high-isoprene emitters" etc.

Light and temperature controls on emissions

For all types of vegetation, an appropriate system describing the emissions flux on an hourly basis is that of Guenther et al. (1996):

$$\text{Flux } (\mu\text{g m}^{-2} \text{ yr}^{-1}) = \int \varepsilon \cdot D \cdot \gamma \, dt \quad (1)$$

where ε is the average emission potential ($\mu\text{g g}^{-1} \text{ h}^{-1}$) for any particular species, "D" is the foliar biomass density ($\text{g dry weight foliage m}^{-2}$), and γ is a unit less environmental correction factor representing the effects of short-term (e.g. hourly) temperature and solar radiation changes on emissions.

Guenther et al. (1991, 1993) showed that, to a very good approximation, the short-term (hourly) variations in emissions of isoprene could be described by the product of a light dependent factor, C_L and a temperature dependant factor, C_T . Thus, the so-called ISOG algorithm:

$$\gamma_{\text{iso}} = C_L \cdot C_T \quad (2, \text{ISOG})$$

The light factor, C_L is given by:

$$C_{L_{iso}} = \frac{\alpha c_{L1} L}{\sqrt{1 + \alpha^2 L^2}} \quad (3)$$

where α (= 0.0027) and c_{L1} (= 1.066) are empirical constants, and L is the PAR flux ($\mu\text{mol photons (400-700nm) m}^{-2} \text{s}^{-1}$). Temperature dependence $C_{T_{iso}}$ is described by:

$$C_{T_{iso}} = \frac{\exp(C_{T1}(T - T_S)/RT_S T)}{1 + \exp(C_{T2}(T - T_M)/RT_S T)} \quad (4)$$

where R is the gas constant (= 8.314 J K⁻¹ mol⁻¹), and c_{T1} (= 95000 J mol⁻¹), c_{T2} (= 230000 J mol⁻¹), and T_M (= 314 K) are empirical coefficients based upon measurements of three plant species: eucalyptus, aspen, and velvet bean, but which seem to be valid for a variety of different plant species (Guenther et al. 1993, Guenther 1997). T_S (= 303 K) is the standard temperature.

The environmental correction factor for monoterpene emissions from most plants are parameterised using the following equation (Guenther et al. 1993):

$$\gamma_{mts} = \exp(\beta \bullet (T - T_S)) \quad (5a, \text{MTS})$$

where β (= 0.09 K⁻¹) is an empirical coefficient based on non-linear regression analysis of numerous measurements present in the literature. This type of emission is associated with vaporisation of terpenes from stores within the plant tissue, and this algorithm is referred to here as MTS.

Recently it was shown that some evergreen oaks, and also Norway spruce, show a light-dependency of monoterpene emissions. At least for *Q. ilex* this dependency seems to be well described by the Guenther isoprene algorithms (Kesselmeier et al., 1996, Seufert et al., 1997). Denoting this behaviour by MTL, we have:

$$\gamma_{mtl} = \gamma_{iso} \quad (5b, \text{MTL})$$

These emission algorithms represent our current knowledge of terpenoid emission by plants. These algorithms will likely need to be revised in the future, when a better biological understanding of the biosynthesis and emission of terpenoids is available, since there remains an uncertainty in the resulting emission estimates that is about a factor of 3 or more. This variation is mainly due to the (1) differences in the emissions from branch to branch and from tree to tree, (2) variation with season, (3) nutrient condition of the plant, (4) stress and (5) experimental errors. Suggestions for improved algorithms have been made by Schuh et al. (1997) and Schnitzler et al. (1997), but the generality of these suggestions need further testing before we can recommend a change from the basic Guenther algorithms.

The relationship between environmental conditions and emission of OVOC is even less understood than isoprene and monoterpenes. Emissions of some of these compounds, including a group of C₆ unsaturates, are strongly influenced by external factors other than light and temperature, such as plant wounding by microbes, insects or mechanical stress. Given the lack of other information regarding the factors controlling oxygenated hydrocarbon emission, the use of equation (5) for parameterisation of oxygenated hydrocarbon emission is recommended (Guenther et al, 1994, Geron et al., 1994, König et al. 1995). i.e.:

$$\gamma_{\text{ovoc}} = \gamma_{\text{mts}} \quad (6)$$

These light and temperature dependencies are illustrated in Figures 3.1 and 3.2.

Figure 3.1: Temperature dependency of isoprene (ISOG) and of terpene stores (MTS) emissions.

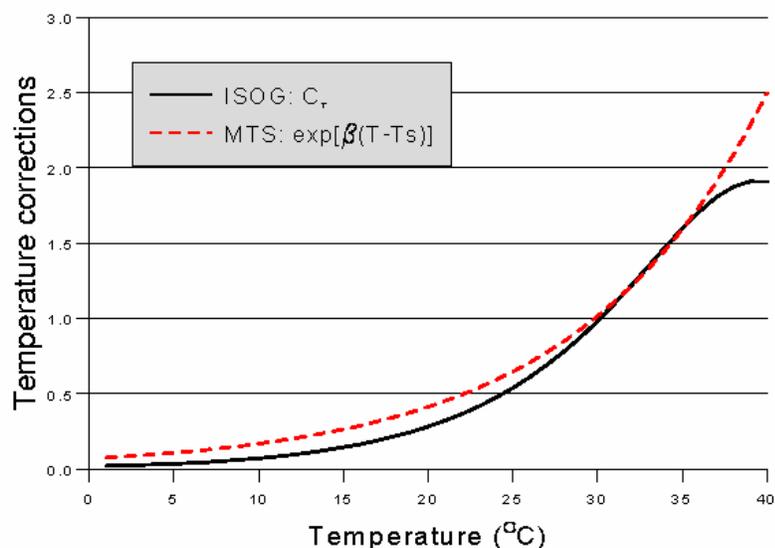
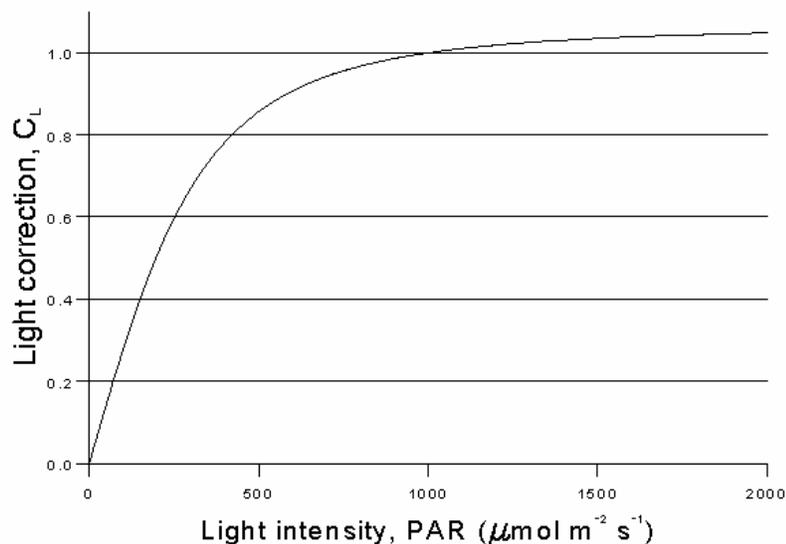


Figure 3.2: Light dependency for emissions of compounds, which are emitted as they are synthesised (e.g. isoprene)



Calculation of annual emissions then requires in principal both temperature and radiation data over the whole year with appropriate spatial resolution. However, many simplifications are possible and the simpler methodology (section 4) proposes a seasonal approach. The use of equations 1-6 above is covered in the detailed methodology (section 5).

3.2 Definitions

Some relevant terms are:

Forest - for the purposes of this guidebook the definition of forest should be as inclusive as possible. In theory all trees should be included, although in practice other definitions of forest may be included in statistical definitions, e.g.:

UNECE/FAO Forest means land with tree crown cover of more than about 20% of the area, with trees usually growing to more than about 7 m in height and able to produce wood. This includes both closed forest formations where trees of various storeys and undergrowth cover a high proportion of the ground and open forest formations with a continuous grass layer in which tree synusia cover at least 10% of the ground.

Branch-level - refers to emissions or measurements where the ambient radiation and temperature is an average over a whole branch, including both sun leaves and shade leaves.

Leaf-level - data refer to data appropriate to a single leaf. Leaf-level emission potentials are on average 1.75 times higher than branch-level rates because the latter are more shaded (Guenther et al., 1994). Emission potentials in this chapter are only given as branch level. (U.S. papers tend to give leaf-level, which requires modelling the shading within a forest canopy)

Coniferous - all trees classified botanically as *Gymnospermae*, generally referred to as softwoods or needle-leafed species.

Non-coniferous - all trees classified botanically as *Angiospermae*, generally referred to as hardwood or **broad-leaved** species. Note that such species can be other deciduous or evergreen.

Deciduous - all plants that shed leaves, usually in the autumn.

DW - dry weight of plants (used for emission rates), as opposed to fresh weight.

Foliar Biomass densities - as used here give the mass of foliage per unit projected ground area, and must not be confused with total biomass densities which have the same units (g m^{-2}) but include wood mass.

OVOC - Other volatile organic compound. Any non-methane VOC species other than isoprene or monoterpenes emitted by vegetation, including oxygenated VOC but also non-oxygenated.

PAR - photosynthetically active radiation, typically about 45-50% of total global radiation, covering the wavelength range 400-700nm.

3.3 Techniques

3.4 Emissions

Biogenic emissions consist of a wide variety of species. Attention has mainly focused on isoprene and the class of monoterpene compounds (alpha-pinene, beta-pinene, limonene, etc.). The remaining 'other' VOC (OVOC) species consist of a large number of species including hydrocarbons and oxygenated compounds (alcohols, aldehydes, etc.), and have proven difficult to quantify in atmospheric samples. See section 9.

3.5 Controls

'Control measures' is not usually an applicable concept for forest emissions. However, it can be mentioned that much of the current forest cover in Europe is artificial, in the sense that the selection of species has been decided by human intervention. Thus, Sitka forest plantations in the U.K. represent an emissions increase over the coniferous forest, which they replaced, so control in terms of species selection could be envisaged. Such action to reduce 'natural' emissions has so far only been undertaken in California as far we are aware!

4 SIMPLER METHODOLOGY

All methodologies for calculating biogenic emissions essentially involve multiplying an emissions factor for a type of vegetation by a statistic giving the amount of vegetation in the country or grid square. Two major alternatives for this are (1) to perform these calculations at a genera or preferably species specific level (requiring for example separate statistics for Norway spruce, Douglas fir, etc.), or (2) to perform the calculations for different ecosystem types. In this latter method, each ecosystem is assumed to consist of a number of species, and the assigned emission rates attempt to give the average emissions from this category.

The rest of this chapter follows a species orientated method (1) approach as far as possible. The main justification for this is that the recent European measurements have differed sufficiently from their American counterparts on an ecosystem basis that where possible detailed species measurements should form the basis of the database. Of course, data still does not exist for many vegetation types in Europe, in which case some ecosystem-assumptions are necessary anyway. These will be based as far as possible on knowledge of European species.

As noted in section 3, an appropriate system describing the emissions flux on an hourly basis is that of Guenther et al. (1996):

$$\text{Flux } (\mu\text{g m}^{-2}\text{yr}^{-1}) = \int \varepsilon \cdot D \cdot \gamma \text{ dt} \quad (1)$$

where ε is the average emission potential ($\mu\text{g g}^{-1}\text{h}^{-1}$) for any particular species, "D" is the foliar biomass density ($\text{g dry weight foliage m}^{-2}$), and γ is a unitless environmental correction factor representing the effects of short-term (e.g. hourly) temperature and solar radiation changes on emissions. For isoprene emissions, and light-activated terpene emissions (so far only quantified for two forest species, *Picea abies* and *Quercus ilex*), γ is a function of light and temperature, and is denoted γ -iso. Terpene and OVOC emissions from most vegetation types are simply dependent on temperature, in which case γ is temperature-only dependant, and denoted γ -mt.

The simplified methodology consists of modifying equation (1) to be a seasonal rather than an hourly calculation.

$$F = \varepsilon \cdot D \cdot \Gamma \quad (7)$$

Where Γ represents the integrated value of γ over the growing season of the vegetation concerned.

Using meteorological data from the EMEP MSC-W models the integrated values, Γ -iso and Γ -mts, have been calculated for both 6 monthly (May-October) and 12 monthly growing seasons, as averages over each country. These have been calculated from hourly γ values, and thus have units of hours. The Γ values are tabulated in Table 4.1. With this simplified methodology we could estimate for example the isoprene emissions from 1 km² of deciduous oak (e.g. *Q. robur*) as simply:

$$\begin{aligned} \text{Emission} &= \text{Area} \times \varepsilon \cdot D \cdot \Gamma\text{-iso} \\ &= 10^6 (\text{m}^2) \times 60 (\mu\text{g g}^{-1} \text{h}^{-1}) \times 320 (\text{g m}^{-2}) \times \Gamma\text{-iso (h)} \end{aligned}$$

For Austria, for example, Table 4.1 gives Γ -iso for 6-months as 452, therefore we have:

$$\text{Emission} = 10^6 (\text{m}^2) \times 60 (\mu\text{g g}^{-1} \text{h}^{-1}) \times 320 (\text{g m}^{-2}) \times 452 (\text{h}) = 8.67 \text{ tonne km}^{-2}$$

Table 4.1: Country average values of integrated environmental correction factors, Γ -iso and Γ -mts for 6 and 12 month growing seasons (unit= hours).

	Γ -mts = Γ -ovoc		Γ -iso	
	6-month	12-month	6-month	12-month
Albania	745	976	563	719
Austria	588	734	452	540
Belarus	753	895	581	684
Belgium	739	969	580	712
Bosnia Herz.	709	893	561	686
Bulgaria	824	1029	620	755
Croatia	883	1121	667	815
Czech Republic	712	885	533	633
Denmark	518	704	373	485
Estonia	565	669	422	491
Finland	458	523	339	379
France	840	1107	669	829
Germany	698	890	525	632
Greece	1076	1440	816	1057
Hungary	966	1188	730	874
Ireland	467	713	337	478
Italy	904	1208	711	902
Latvia	636	757	486	572
Lithuania	675	813	516	613
Luxembourg	786	1003	620	745
Macedonia,F.Y.R.	631	783	492	597
Moldova, Rep. of	858	1040	649	771
Netherlands	676	901	513	643
Norway	327	397	240	284
Poland	736	912	558	669
Portugal	1015	1388	853	1093
Romania	783	964	587	706
Russia, Fed.	808	917	637	717
Slovakia	797	977	607	724
Slovenia	745	940	562	682
Spain	982	1301	806	1004
Sweden	423	508	315	368
Switzerland	465	580	368	432
Turkey	976	1263	783	983
United Kingdom	493	720	358	492
Ukraine	856	1023	656	771
Yugoslavia	752	937	557	674

5 DETAILED STATE OF THE ART METHODOLOGY

The detailed methodology still relies on the basic equations (1-6) given above, but allows for the use of better input information and a more refined calculation if local meteorological data are available. We give details for calculations at either a monthly or hourly resolution.

5.1 Monthly calculation

For the monthly calculation we make the following assumptions for the integration of the ISOG-type emissions:

1. The light-intensity variation given by equation 2 can be replaced by a simple step-function, where $C_L = 1$ during most of the day and zero otherwise.
2. The calculation of the temperature correction (Eqns. 4,5) need not be done every hour, but instead may be approximated by a monthly average daytime temperature.
3. Ambient temperature and light-intensity provide a reasonable approximation to leaf-level light and temperature.

Approximation (1) is generally rather good, as light levels quickly reach $1000 \mu\text{mol m}^{-2} \text{s}^{-1}$ during the morning hours in most locations, even with moderate cloud cover. ($200 \mu\text{mol m}^{-2} \text{s}^{-1}$ is set as the cut-off for defining daylength as this corresponds to approx. $C_L = 0.5$). Approximation (2) introduces larger errors, but only of order 20% or so, which is much less than the uncertainties in the emission potentials. Approximation (3) has been tested by Simpson et al.(1995) and shown to introduce only moderate uncertainties for European conditions, again much less than those of the emission potentials.

The number of light-hours per day corresponding to the above definition can be calculated as a simple function of latitude and month:

Table 5.1: Number of light-hours* per day (N_L) as a function of latitude and month.

Lat	Jan	Feb	Mar	Apr	May	Jun	Jul	Aug	Sep	Oct	Nov	Dec
80	0.0	0.0	0.0	13.1	24.0	24.0	24.0	15.2	6.1	0.0	0.0	0.0
78	0.0	0.0	4.4	12.9	20.5	24.0	24.0	14.6	7.2	0.0	0.0	0.0
76	0.0	0.0	5.8	12.7	18.6	24.0	20.2	14.1	7.9	0.0	0.0	0.0
74	0.0	0.0	6.8	12.6	17.5	20.9	18.6	13.8	8.5	0.0	0.0	0.0
72	0.0	0.0	7.4	12.5	16.7	19.1	17.6	13.6	8.9	0.0	0.0	0.0
70	0.0	0.0	7.9	12.4	16.1	18.0	16.8	13.4	9.2	2.5	0.0	0.0
68	0.0	1.6	8.4	12.3	15.6	17.2	16.2	13.2	9.4	4.2	0.0	0.0
66	0.0	3.6	8.7	12.2	15.2	16.6	15.8	13.0	9.6	5.2	0.0	0.0
64	0.0	4.7	8.9	12.2	14.9	16.1	15.4	12.9	9.8	5.9	0.0	0.0
62	0.0	5.4	9.1	12.1	14.6	15.7	15.0	12.8	9.9	6.4	1.5	0.0
60	2.4	6.1	9.4	12.1	14.3	15.4	14.7	12.7	10.1	6.9	3.3	0.0
58	3.7	6.6	9.5	12.0	14.1	15.0	14.4	12.6	10.2	7.3	4.3	2.2
56	4.6	7.0	9.7	12.0	13.9	14.7	14.2	12.5	10.3	7.7	5.1	3.5
54	5.3	7.3	9.8	11.9	13.7	14.5	14.0	12.4	10.4	7.9	5.7	4.4
52	5.8	7.7	9.9	11.9	13.5	14.2	13.8	12.3	10.4	8.2	6.2	5.1
50	6.3	7.9	10.0	11.9	13.4	14.0	13.6	12.2	10.5	8.4	6.6	5.7
48	6.7	8.2	10.1	11.8	13.2	13.8	13.4	12.2	10.6	8.6	7.0	6.2
46	7.1	8.4	10.2	11.8	13.1	13.6	13.3	12.1	10.6	8.8	7.3	6.6
44	7.4	8.6	10.2	11.8	12.9	13.5	13.1	12.1	10.6	9.0	7.6	6.9
42	7.7	8.8	10.3	11.7	12.8	13.3	13.0	12.0	10.7	9.1	7.9	7.3
40	7.9	9.0	10.4	11.7	12.7	13.1	12.9	11.9	10.7	9.3	8.1	7.6
38	8.2	9.1	10.4	11.6	12.6	13.0	12.8	11.9	10.8	9.4	8.4	7.8
36	8.4	9.3	10.5	11.6	12.5	12.9	12.6	11.8	10.8	9.6	8.6	8.1

Notes: Day-lengths (in hours) calculated for the 15th of each month from Latitudes 80 degrees N to 36 degrees N. *Period of light-hours defined for $PAR > 200 \mu\text{mol m}^{-2} \text{s}^{-1}$.

If we let mm1 and mm2 be the start and end of the growing season for a particular vegetation type, N_d (mm) be the number of days per month, N_L (mm) be the number of light-hours per day (Table 5.1), and T_{mm} be the monthly mean temperature, for month "mm", yearly emissions can be evaluated with:

$$\text{Emis(isoprene)} = \sum_{mm=mm1}^{mm2} A.D. \gamma_{iso}(T_{mm}). N_d(mm). N_L(mm)$$

Emissions of terpenes from species displaying MTL behaviour are also described by this equation.

For the yearly emissions of species displaying the MTS-type behaviour there is no light-dependency, and we perform the calculation for 24 hours per day:

$$\text{Emis(monoterpenes)} = \sum_{mm=mm1}^{mm2} A.D. \gamma_{mt}(T_{mm}). N_d(mm). 24$$

Similarly,

$$\text{Emis(OVOC)} = \sum_{mm=mm1}^{mm2} A.D.\gamma_{voc}(T_{mm}).N_d(mm).24$$

5.2 Hourly calculation

If desired, and appropriate meteorological data are available, the environmental correction factors (γ) may be evaluated on an hourly basis using local surface temperature and sunlight conditions. The algorithms, temperature and light corrections, C_T and C_L , are as given in Equations 1-7 above.

Refinements:

Age distribution of forest

Isidorov et al. (1993) have pointed out that a land-use data-base built up with knowledge of not only the area but also the age distribution within each region can give a better description of the biomass densities. This approach requires more data but helps to eliminate a potentially large area of uncertainty.

Seasonal variation

Foliar density varies markedly over the year, and this can be straightforwardly incorporated into the above calculations if data are available through the use of seasonal-dependent foliar biomass density.

Altitude temperature correction

Atmospheric temperature generally decreases with height at a rate of ca. 6 degrees C per km. Thus, data obtained from a meteorological station at a given height may be corrected to temperatures in another location (e.g. on a mountainside) before applying any of the detailed methodologies.

6 RELEVANT ACTIVITY STATISTICS

Vegetation coverage in terms of the vegetation types discussed in section 8 is required, together with foliar biomass estimates (D), and estimates of growing seasons. Commercial forestry at least is usually well documented. Other wooded land is a common category where definitions are more problematic.

For a good inventory it is actually most important to specify the correct foliar biomass density to accompany any given area of vegetation. This is because "area" is an ill-defined quantity in many instances, e.g. 1 km² of wooded area may include very dense forest with an average foliar biomass density of, say, 1400 g m⁻², or it may contain scattered trees with only 100g m⁻².

The new SNAP codes have been designed to encourage the use of data for each tree species separately for at least the most common trees. Very nice examples of this type of compilation

are provided by Andreani-Aksoyoglu and Keller (1995) for Switzerland, and Ortiz and Dory (1990) for Spain, the latter tabulating area coverage and mean biomass factors for all 50 level III territorial units.

Categories such as mixed forest should be avoided as this gives no information on species content. If species-specific data are not available, then genus-level data should be used. Only as a last resort should more general categories be supplied.

As pointed out by Veldt (1989) common vegetation names are often confusing, and care should be taken to provide Latin names of species as well as common names of all species. Translations of some common tree species names are included in Table 14.1, taken from EC (1996).

Foliar Biomass densities

For the simpler methodology, seasonal average foliar biomass densities may be used. Default values are suggested below, and in section 8. These suggestions appear to fit quite well a wide range of measurements, but the variability of Mediterranean vegetation may cause some problems. For example, Ortiz & Dory (1990) mention a land-use class, Monte hueco, which consists of a mixture of species, with biomass densities as low as 100 g m⁻². For coniferous forests, Veldt suggests densities of 700-1400 g m⁻² for different species < 60 deg N latitude, whereas Ortiz and Dory use 400 g m⁻². Even further north, variations are great. Andreani-Aksoyoglu and Keller, 1995, quote a biomass factor for oak species of 530 g m⁻². Some variations are systematic; Isidorov et al. (1993) points out that foliar biomass as a proportion of total tree biomass increases in harsher conditions, and with age.

*** Therefore, it is **STRONGLY RECOMMENDED** that foliar biomass densities appropriate to the local vegetation are used. These may well be a factor of 2 or three different to the default values. ***

Table 6.1: Default foliar biomass densities (adapted from Veldt, 1989)

Land Use Type		Foliar Biomass Density, D (g m ⁻²)
Broadleaf:		
Deciduous Oaks		320
Birch (<i>Betula</i>)		320
Poplar, aspen (<i>Populus</i>)		320
Default deciduous broadleaved		300
Evergreen broadleaved		500
Conifers		
Norway spruce (<i>Picea abies</i>)	> 60° N lat.	800
	55-60° N lat.	1400
	< 55° N lat.	1600
Sitka spruce (<i>Picea sitchensis</i>)		1400
Other spruce		1400
Scots pine (<i>Pinus sylvestris</i>)	> 60° N lat.	500
	< 60° N lat.	700
Other <i>Pinus ssp.</i>		700
(Fir) <i>Abies ssp.</i>		1400
Douglas Fir (<i>Pseudotsuga menziesii</i>)		1000
Larch (<i>Larix</i>)		300
Other coniferous		1000

Comment on Satellite data

Satellites provide a spatially comprehensive method of mapping vegetation with very high-resolution. Use of such data is encouraged, but a strong warning should be issued that ground-validation is essential if biogenic emissions are to be estimated. The apparent beauty and detail of a satellite image should not be mistaken for accuracy! Satellite data are easily misinterpreted (wrong species, problems with non-dominant vegetation, etc.) and even in the United States where biogenic emission inventories are very advanced, discrepancies of up to a factor of 5 are still found between satellite-derived isoprene emissions and ground-based determinations (Lamb et al., 1997).

7 POINT SOURCE CRITERIA

No point sources

8 EMISSION FACTORS, QUALITY CODES AND REFERENCES

Emission potentials (ϵ) are required separately for isoprene, monoterpenes and OVOC. Further, for monoterpenes two classes of behaviour are distinguished. For most trees emissions are temperature-only dependant, controlled by the γ -mts environmental factor (equation 5a). For evergreen oaks the MTL algorithm is used (eqn. 5b).

Emission potentials for a wide variety of species have recently been compiled by Guenther et al. (1994, 1997), Geron et al.(1994) for American woodlands, and by Steinbrecher (1997) and Seufert et al. (1997) for European species. Very little reliable experimental data on the emissions of OVOCs is available, and consequently Guenther et al. (1994) recommended the use of a uniform emission rate of $1.5 \mu\text{g g}^{-1} \text{h}^{-1}$ for all tree species, recognising that this was a first order approximation to a ten-fold range ($0.5\text{-}5 \mu\text{g g}^{-1} \text{h}^{-1}$). The data of König et al.(1995) fall within this range, and so until further European data are available then $1.5 \mu\text{g g}^{-1} \text{h}^{-1}$ also seems a reasonable choice for preliminary, first-order, estimates of OVOC emissions in Europe.

The emission potentials are given in Table 8.1.

Table 8.1: Standard emission potentials ($\mu\text{g g}^{-1} \text{h}^{-1}$ at 30 deg. C and PAR=1000 $\mu\text{mol m}^{-2} \text{s}^{-1}$) for European trees.

Isoprene and monoterpene emission potentials are taken from Guenther et al., 1994, 1997, or Geron et al., 1994, except where European measurements can provide a basis, as indicated by additional refs. For terpenes ϵ -mtl denote emissions controlled by light and temperature (using γ -mtl), whereas ϵ -mts denote emissions controlled by temperature only. All isoprene rates are branch-level, often derived from leaf-level U.S. estimates by division by 1.75.

Common name (example)	Latin name	Type	Foliar	Iso.	Terpenes		O- VOC	Additional Refs.	
			biomass density, D g m ⁻²	ϵ -iso	ϵ -mtl	ϵ -mts	ϵ -ovoc	Iso.	Terp.
Fir	<i>Abies</i>	e	1400	0	0	3	1.5		
Maple/Sycamore*:	<i>Acer</i>	d	320	0	0	3	1.5	S93	S93
Common Alder	<i>Alnus</i>	d	320	0	0	1.5	1.5	S93	S93
Birch	<i>Betula</i>	d	320	0	0	0.2	1.5	K	P,K
Hornbeam	<i>Carpinus</i>	d	320	0	0	0.65	1.5	K	K
Cedar	<i>Cedrus</i>	e	700	0	0	1.5	1.5		
Orange	<i>Citrus sp.</i>	d	320	0	0	1.5	1.5		
Italian cypress	<i>Cupressus</i>	d	700	0	0	0.65	1.5		
Blue gum	<i>Eucalyptus sp.</i>	e	400	20	0	3	1.5	Str97b	H
European beech	<i>Fagus</i>	d	320	0	0	0.65	1.5	P,S93,K, Sh	K,Sh
Ash	<i>Fraxinus</i>	d	320	0	0	0	1.5	S93	S93
Walnut	<i>Juglans</i>	d	320	0	0	3	1.5		
Common juniper	<i>Juniperus</i>	e	700	0	0	0.65	1.5	O	O
European larch	<i>Larix</i>	d,c	300	0	0	1.5	1.5	S93	S93
Olive	<i>Olea</i>	e	200	0	0	0	1.5		
Date palm	<i>Phoenix</i>			20	0	0	1.5		
Spruce	<i>Picea sp.</i>	e	Varies	1	1.5	1.5	1.5	As P.abies	
Norway spruce	<i>Picea abies</i>	e	Varies	1	1.5	1.5	1.5	S94,Ke;	J,Ke, S94,LP
	<i>Picea omorika</i>	e	Varies	10	0	0.65	1.5		
Blue spruce	<i>Picea pungens</i>	e	Varies	1	0	0.65	1.5		
Sitka spruce	<i>Picea sitchensis</i>	e	Varies	6	0	3	1.5	Str96,97b,Sm	
Pines:	<i>Pinus sp.</i>	e	700	0	0	3	1.5		-
Aleppo pine	<i>Pinus halepensis</i>	e	700	0	0	0.65	1.5		H
Umbrella pine	<i>Pinus pinea</i>	e	700	0	0	6	1.5	Ks,Std,Str97a,Sf	
Maritime pine	<i>Pinus pinaster</i>	e	700	0	0	0.2	1.5		Si
Scots pine	<i>Pinus sylvestris</i>	e	Varies	0	0	1.5	1.5		J
Pistachio	<i>Pistacia sp.</i>			0	0	3	1.5	H,Ha	H,Ha
Americ. sycamore*	<i>Platanus</i>	d	320	34	0	0	1.5		
Poplar	<i>Populus</i>	d	320	60	0	0	1.5	H	
Cherry#4	<i>Prunus</i>	d	300	0	0	0	1.5		
Douglas Fir	<i>Pseudotsuga</i>	e	1000	0	0	1.5	1.5		D

Oaks:	-			-	-	-		-;	-
Default deciduous Oak#1	-	d	320	60	0	0.2	1.5	Sf	Sf
Default evergreen Oak#2	-	e	500	0	20	0	1.5	Sf	Sf,
Turkey oak	<i>Quercus cerris</i>	d	320	0	0	1	1.5	S97	S97
Kermes/Holly oak	<i>Quercus coccifera</i>	e	500	0	20	0	1.5	SH	SH
Hungarian oak	<i>Quercus frainetto</i>	d	320	100	0	0.2	1.5	S97,Sf,	Sf
Holm oak	<i>Quercus ilex</i>	e	500	0	20	0	1.5	Be,Ks96,Str97,Sf	
Sessile oak	<i>Quercus petraea</i>	d	320	60	0	0.2	1.5	K,S97,Str97b,Sf	
Downy oak	<i>Quercus pubescens</i>	d	320	60	0	0.2	1.5	S97	S97
European oak#3	<i>Quercus robur</i>	d	320	60	0	0.2	1.5	S93,I	S93
Cork oak	<i>Quercus suber</i>	e	500	0	0	0.2	1.5	Sf	Sf
Locust	<i>Robinia pseudoacacia</i>	d	320	10	0	-	1.5		
Willow	<i>Salix</i>	d	150	34	0	0.2	1.5	O	S93
Saw-palmetto	<i>Serenoa</i>	d	320	10	0	0	1.5		
Lime tree/Basswood	<i>Tilia</i>	d	320	0	0	0	1.5		
Elm	<i>Ulmus</i>	d	320	0	0	0.2	1.5		

Notes:

Type gives evergreen (e),deciduous (d), or (d,c) for *Larix* deciduous coniferous

#1 e.g. *Q. rubra*, *Q. faginea*, *Q. lusitanica*.

#2 e.g. *Q. rotundifolia*, *Q. calliprinos*, *Q. ithhaburiensis*, *Q. coccifera*

#3 also known as English oak, Pendunculate oak

#4 includes almond,apricot, blackthorn (sloe), peach.

* Sycamore = *Acer pseudoplatanus*, not to be confused with the American sycamore, *Platanus occidentalis*

Refs:

Be Bertin et al, 1997; D Duyzer, 1993; H Hewitt, C.N. and Owen, S., pers.comm.; I Isidorov et al., 1985; J Janson, 1993; K Koenig et al., 1995; Ks96,97 Kesselmeier et al., 1996, 1997; LP Lindskog and Potter, 1995; Ha Hanson et al., 1997; O Owen et al., 1997; P Puxbaum, 1997; Sh Schuh et al., 1997; Si Simon et al., 1994; Sf Seufert et al., 1997; Sm Simpson et al., 1995; Std Staudt et al., 1997; S94 Steinbrecher, R., 1994; S93,97 Steinbrecher et al., 1993, 1997; Str96,97a,97b Street et al., 1996,1997a,1997b.

9 SPECIES PROFILES

Emission (γ) potentials have been given separately for isoprene, terpenes, and OVOC, and this division represents the most important level of speciation. However, there are many species represented within the class of terpenes and OVOC covering a wide range of chemical behaviour. This section attempts some guidance as to likely breakdowns among the monoterpene and OVOC classes.

Monoterpenes

Although many types of monoterpenes exist, most plants emit only 2-3 major species, with the reactive α -pinene often dominating emissions from species such as Norway spruce and Scots pine (Janson, 1993). The ratio of one compound to another is very variable, both with season and temperature, so it is very difficult to specify the speciation in a quantitative way

(Janson, 1993). In order to illustrate the major compounds, Table 9.1 compares the ratios of several monoterpenes to α -pinene obtained from several studies. Table 9.2 groups a number of species in order of their relative frequency of emission.

Table 9.1: Relative composition of hydrocarbon-mix emitted by vegetation as reported by different authors, adapted from Duyzer (1993). Numbers in % are given relative to α -pinene (α -pinene is 100%)

	Veldt:91	Janson:93	Janson:93	Steinb.'93	Simon:93	Duyzer:93
	Average of several pines	Scots pine	Norway spruce	Norway spruce	Maritime pine	Douglas fir
β -pinene	40	33	5	17	105	40-100
3-carene	30	111	6-800		50	30-80
Limonene	26	61	5-15	13	44	20-60

Table 9.2: Examples of monoterpenes emitted by vegetation into the atmosphere (Zimmerman, 1979; Isidorov, 1985, as given by Guenther et al., 1994).

Major	Frequent	Occasional
Δ^3 - Carene	α Thujene	α Fenchene
d-Limonene	Tricyclene	β -Fenchene
Myrcene	Terpinolene	δ -Fenchene
α -pinene	α -Terpinene	ϵ -Fenchene
β -pinene	β -Terpinene	Bornylene
Sabinene	γ -Terpinene	Alloocimene
Camphene	p-Cymene	Methyl chavicol
1,8-Cineole	α -Phellandrene	p-Cymen-8-ol
β -Phellandrene	trans-Ocimene	Linalool
	cis-Ocimene	2-Methyl-6-methylene-1,7-octadiene-3-one
	2-Carene	Pinocarvone
		Verbenone
		Fenchone
		Thujone
		Camphor

OVOC

The identification and quantification of OVOC emissions from plants has proven one of the most difficult problems in evaluating total biogenic emissions. OVOC consists of a wide variety of compounds, many of which have been difficult to measure. Examples are alcohols, ketones, esters, ethers, aldehydes, alkenes and alkanes. Useful reviews can be found in Puxbaum (1997), Bode et al. (1997), Guenther et al. (1994) and Kotzias et al. (1997). The most extensive quantitative European data-set appears to be that of König et al. (1995), otherwise some screening studies are also available (Hewitt and Street, 1992, Steinbrecher, 1994, Isidorov, 1992, Goldstein et al., 1996, Arey et al., 1991a,b).

10 UNCERTAINTY ESTIMATES

None of the biogenic emission inventories used in Europe can be compared in terms of complexity or accuracy with those generated in the U.S.. All European methodologies have been severely limited by the availability of data on a European scale. Several key items are either missing or known to only a limited extent, necessitating some rather arbitrary choices.

Assessment of the uncertainties inherent in calculations of biogenic VOC emissions in Europe is rather difficult. As a starting point, estimates of the uncertainty of even recent U.S. inventories have suggested up to a factor of 3 for isoprene (Guenther et al., 1994). Further, even though much progress has been made in emission potentials and algorithms (Guenther et al., 1993, 1997, Seufert et al., 1997), awareness has grown of the large uncertainties associated with specifying land-cover for particular species. Even in the U.S., where land-use databases exist over the whole country in consistent format, uncertainties associated with specifying forest coverage are still significant (Guenther et al., 1994). In Europe such uncertainties are very much greater because such coherent land-use data sets have not yet become available.

We discuss some of the important factors contributing to the total uncertainty of the European emission estimates below.

Emission potentials

Even with large campaigns such as BEMA (Seufert et al., 1997) emission factors for European species are very few, and taken from a very limited set of conditions and samples. Genus-level potentials derived in the U.S. are often not appropriate for Europe because the species mix within a genus is often very different. It is clear that many more measurements are required before emissions in Europe can be described with any confidence, but meanwhile the first positive steps that can be taken are to collect good land-use data as a basis for any inventory.

Land-use data

The focus of most forest statistics appears to be the area of productive, coniferous forest, rather than the categories of most interest for biogenic inventories. Even for the coniferous forest category definitions vary greatly; 1 km² of coniferous forest appears to mean that 50% of the stem-volume is coniferous in Finland and Norway, 70% in Sweden, 80% in Ireland, and 100% in the U.K. (UN ECE, 1985). Whichever definition is used, the aim should be to get the best description of foliar biomass for the area and tree species concerned.

Biomass data

Although the biomass data given in the simpler methodology can be used as default values if no other information is available, factor of two uncertainties can easily be introduced. Clearly the best solution is for each country to specify biomass densities appropriate to local conditions.

OVOC emissions

Guenther et al. (1993) noted that the recommended emission rate of $1.5 \mu\text{g g}^{-1} \text{h}^{-1}$ is associated with a 10-fold range ($0.5 - 5 \mu\text{g g}^{-1} \text{h}^{-1}$) in possible emissions, and that even this may underestimate some emissions.

Final remarks

It has been recognised that the minimum level of uncertainty in global biogenic emission estimates is a factor of 3 (Guenther et al., 1995), but this is likely to represent a lower limit for the accuracy of European emission estimates. Further, this figure relates to estimates of annual emissions. Uncertainties for episodic calculations must obviously be substantially greater.

11 WEAKEST ASPECTS/PRIORITY AREAS FOR IMPROVEMENT IN CURRENT METHODOLOGY

The emission factors and knowledge of land-use within each region are certainly the weakest aspects. The emission factors can only be improved with more measurements. The land-use problem is primarily one of data collection, as presumably forestry and agricultural Institutes hold quite detailed data for most countries. Collection of this land-use data is of the greatest priority.

12 SPATIAL DISAGGREGATION CRITERIA FOR AREA SOURCES

Follows from land-use and climate data.

13 TEMPORAL DISAGGREGATION CRITERIA

The details of hourly calculations are given in section 5, detailed methodology.

It is worth noting that annual emissions of biogenic emissions give only a limited insight into the importance of these compounds. For assessing their impacts on photochemical ozone formation it is the biogenic emissions during the warmest and sunniest days, which are of interest. In practice therefore, photochemical oxidant models all calculate their own biogenic emission rates internally using short period temperature, radiation data in conjunction with land-use data.

14 ADDITIONAL COMMENTS

Recent developments and re-evaluations of previous methodologies have resulted in significant changes in the emission factors, which should be used in inventorying biogenic VOC emissions. This chapter has presented information on the new emission factors for a range of species derived from the latest American and European evaluations. In addition, much progress has been made in developing algorithms to describe the emission-temperature-sunlight relationships for isoprene, monoterpenes and other VOC. Still, these algorithms will certainly be changed in the future as knowledge of the underlying processes improves. Suggestions for modifications to include long-term (seasonal changes) to the emission potentials, or other improvements have been presented by Guenther (1997), Schnitzler et al., 1997, and Schuh et al. (1997), although more work is needed to evaluate these algorithms before they can be recommended for the Guidebook.

Canopy models

It is possible to apply complex 'canopy' approaches in which forest canopy models are used to estimate levels of temperature and radiation at different heights within a canopy (e.g. Pierce and Waldruff, 1991, Lamb et al., 1993), and such an approach was tested in Simpson et al. (1995). Canopy approaches should be used together with 'leaf-level' emission factors, as opposed to the 'branch-level' factors given in section 8. However, the difference in emissions estimates between a canopy model and simple use of branch-level estimates is relatively small (up to 20%). Given the much larger uncertainties in the emission potentials, uncertainties introduced by the forest-canopy model itself (e.g. in temperature profiles within the canopy), and the lack of evaluation of such models in European conditions, we do not recommend applying such a model for European emissions at this stage.

The emission factors given in section 8 are therefore exclusively for use where emission-canopy models are not used.

It should be noted that this section still presents a simpler methodology for calculating emissions than can be found in Guenther et al., 1995. We make no attempt to account for factors such as net primary production, leaf-area index, or vegetation index. No canopy radiative model is used. Such factors might improve the accuracy of the estimates somewhat, but until the basic emission factors for European vegetation are more firmly established too much sophistication in the inventory procedure seems unnecessary. Groups having the data and resources to implement such methods are referred to Guenther et al., 1994, Guenther et al., 1995 or Geron et al., 1994.

Table 14.1: Generic names of tree species in different European languages

Botanical Name	French	German	Greek	Italian
<i>Fagus sylvatica</i>	Hêtre	Rotbuche	Οξνα δασική	Faggio
<i>Quercus petraea</i>	Chêne rouvre	Traubeneiche	Δρνς αποδισκο	Rovere
<i>Quercus robur</i>	Chêne pédonculé	Stieleiche	Δρνς ποδισκοφορος	Farnia
<i>Quercus ilex</i>	Chêne vert	Steineiche	Αρια	Leccio
<i>Quercus suber</i>	Chêne liège	Korkeiche	Φελλοδρνς	Sughera
<i>Pinus sylvestris</i>	Pin sylvestre	Gemeine Kiefer	Δασικ πενκη	Pino silvestre
<i>Pinus nigra</i>	Pin noir	Schwarzkiefer	Μανη πενκη	Pino nero
<i>Pinus pinaster</i>	Pin maritime	Seestrandkiefer	Οαλασσια πενκη	Pino marittimo
<i>Pinus halepensis</i>	Pin d'Alep	Aleppokiefer	Χαλεπιος πενκη	Pino d'Aleppo
<i>Picea abies</i>	Épicéa commun	Rotfichte	Ερνθρελατη νψηλη	Abete rosso
<i>Picea sitchensis</i>	Épicéa de Sitka	Sitkafichte	Ερνθρελατη	Picea di Sitka
<i>Abies alba</i>	Sapin pectiné	Weißtanne	Λενκη ελατη	Abete bianco
<i>Larix decidua</i>	Mélèze d'Europe	Europäische Lärche	Λαριξ ενρωπαικη	Larice

Botanical Name	Portuguese	Russian	Spanish	Swedish
<i>Fagus sylvatica</i>	Faia	áoê éãñííé	Haya	Bok
<i>Quercus petraea</i>	Carvalho branco Americano	áoá ñêãëúíúé	Roble albar	Bergek
<i>Quercus robur</i>	Carvalho roble	áoá ñãðáàòúé	Roble común	Ek
<i>Quercus ilex</i>	Azinheira	áoá éàíáííúé	Encina	Stenek
<i>Quercus suber</i>	Sobreiro	áoá ñðíáêíáúé	Alcornoque	Korkek
<i>Pinus sylvestris</i>	Pinheiro silvestre	ññíá ñáúéíñáííáÿ	Pino silvestre	Tall
<i>Pinus nigra</i>	Pinheiro Austriaco	ññíá ñ ðíáÿ	Pino laricio	Svarttall
<i>Pinus pinaster</i>	Pinheiro bravo	ññíá ñ ðèííðñêáÿ	Pino negral	Terpentintall
<i>Pinus halepensis</i>	Pinheiro de alepo	ññíá ñ èãññêáÿ	Pino carrasco	Aleppotall
<i>Picea abies</i>	Picea	áëú ááðííáéñêáÿ	Abeto rojo	Gran
<i>Picea sitchensis</i>	Picea de Sitka	áëú ñèòèíñêáÿ	Picea de Sitka	Sitkagran
<i>Abies alba</i>	Abeto blanco	íèòòá ááèáÿ	Abeto común	Sivergran
<i>Larix decidua</i>	Larício Europeu	èèñòááííèòá ááðííáéñêáÿ	Alerce	Europeisklärk

Botanical Name	Danish	Dutch	English	Finnish
<i>Fagus sylvatica</i>	Bøg	Beuk	Common beech	Pyökki
<i>Quercus petraea</i>	Vintereg	Wintereik	Sessile oak	Talvitammi
<i>Quercus robur</i>	Stilkeg	Zomereik	European oak	Metsätammi
<i>Quercus ilex</i>	Steneg	Steeneik	Holm oak	Rautatammi
<i>Quercus suber</i>	Korkeg	Kurkeik	Cork oak	Korkkitammi
<i>Pinus sylvestris</i>	Skovfyr	Grove den	Scots pine	Metsämänty
<i>Pinus nigra</i>	Østrisk fyr	Oostenrijkse/ Corsicaanse zwarte den	Corsican/Austrian black pine	Euroopanmusta- mänty
<i>Pinus pinaster</i>	Strandfyr	Zeeden	Maritime pine	Rannikkomänty
<i>Pinus halepensis</i>	Aleppofyr	Aleppoden	Aleppo pine	Aleponmänty
<i>Picea abies</i>	Rødgran	Fijnspar	Norway spruce	Metsäkuusi
<i>Picea sitchensis</i>	Sitkagran	Sitkaspar	Sitka spruce	Sitkankuusi
<i>Abies alba</i>	Ædelgran	Zilverden	Silver fir	Saksanpihta
<i>Larix decidua</i>	Lærk	Europese lariks	European larch	Euroopanlehti- kuusi

15 SUPPLEMENTARY DOCUMENTS

The American Biogenic Emission Inventory System (BEIS) has resulted in extensive lists of emission potentials. The latest published version is Geron et al. (1994). The updated BEIS-3 version is currently under preparation by Guenther et al. (1998). (Some of these rates have been already adopted in Table 8.1).

A qualitative list of isoprene and monoterpene emitting species is held at:

Hewitt, C. N., Street R.A. and Scholefield P.A. (1998):
Isoprene and monoterpene-Emitting Species Survey 1998.
<http://www.es.lancs.ac.uk/es/people/pg/pas/download.html>

16 VERIFICATION PROCEDURES

If satellite data have been used in the land-use mapping process it is essential that these be independently verified by on-the-ground surveys. Large errors are possible in the identification of vegetation types and biomass from remote sensing methods.

In general all of the emission potentials are built upon very few data. More measurements are required of at least the major sources, and several different measurement techniques need to be applied in order to eliminate the artefacts (usually enhanced emissions) easily generated by disturbances to the vegetation.

17 REFERENCES

- Andreani-Aksoyoglu S. and Keller J., 1995, Estimates of monoterpenes and isoprene emissions from the forests of Switzerland, *J. Atmos. Chem.*, 20, 71-87.
- Arey J., Winer A., Atkinson R., Aschman S., Long W., Morrison C. and Olszyk D. (1991) Terpenes emitted from agricultural species found in California's central valley, *J. Geophys. Res.*, 96D, 9329-9336.
- Arey J., Winer A. M., Atkinson R., Aschmann S. M., Long W. D. and Morrison C. L. (1991): The emission of (z)-3-hexen-1-ol, (z)-3-hexenylacetate and other oxygenated hydrocarbons from agricultural plant species. *Atmos. Environ.* 25A , 1063-1075.
- Bode K., Helas G. and Kesselmeier J. (1997): Biogenic contribution to atmospheric organic acids. In: *Biogenic volatile organic compounds in the atmosphere*, Helas G., Slanina J. and Steinbrecher R. (eds.), SPB Academic Publishing bv Amsterdam, pp. 157-170.
- Bertin N., Staudt M., Hansen U., Seufert G., Ciccioli P., Foster P., Fugit J.L. and Torres, L. (1997) Diurnal and seasonal course of monoterpene emissions from *Quercus ilex* (L.) under natural conditions - applications of light and temperature algorithms, *Atmos. Environ.*, 31, S1, 135-144.

- Boissard C., Cao X. L., Street R. A., Shuttleworth S. M., Juan C.-Y., Duckham S. C., S. C. Hewitt S. C., Beverland I. J., ONeil D. H., Moncrieff J. B., Milne R. and Fowler D. (1996): Quantification of Non-methane Hydrocarbon Emissions from Two Terrestrial Ecosystems in the UK. in: Proceedings of EUROTRAC Symposium 96, Borrell P. M., Borrell P., Cvitas T., Kelly K. and Seiler W. (eds.), Computational Mechanics Publications, Southampton, pp. 163-167.
- Corchno, S.B., Arey J. and Atkinson R. (1992): Hydrocarbon emission from twelve urban shade trees of the Los Angeles Basin. *Atmos. Environ.* 3 (26B), 339-348.
- Duyzer J., 1993, Measurements of the emissions of monoterpenes from Douglas fir forest, Technical Report IMW-R 93/312, TNO, Delft, NL.
- EC, 1996, European programme for the intensive monitoring of forest ecosystems, General information on the permanent observation plots in Europe (level II); European Commission, DG VI, Brussels.
- Evans R. C., Tingey D. T. and Gumpertz M. L. (1985): Interspecies variation in terpenoid emissions from Engelmann and Sitka spruce seedlings. *Forest Sci.* 31, 132-142.
- Evans R.C., Tingey D.T., Gumpertz M.L., and Burns W.F. (1982) Estimates of isoprene and monoterpene emission rates in plants, *Bot. Gaz.*, 143, 304-310.
- Fehsenfeld F., Calvert J., Fall R., Goldan P., Guenther A.B., Hewitt C.N., Lamb B., Liu S., Trainer M., Westberg H. and Zimmerman P. (1992) Emissions of volatile organic compounds from vegetation and the implications for atmospheric chemistry, *Global Biogeochem. Cycles*, No. 6, 389-430.
- Geron C., Guenther A. and Pierce T. (1994) An improved model for estimating emissions of volatile organic compounds from forests in the Eastern United States, *J. Geophys. Res.*, 99, 12773-12792.
- Geron C.D., Pierce T.E. and Guenther A.B. (1995) Reassessment of biogenic volatile organic compound emissions in the Atlanta area, *Atmos. Environ.*, 29, No. 13, 1569-1578.
- Goldstein A. H., Fan S. M., Goulden M. L., Munger J. W. and Wofsy S. C. (1996): Emissions of ethene, propene, and 1-butene by a midlatitude forest. *J. Geophys. Res.* 101(D4), 9149-9157.
- Guenther A., Greenberg J., Helmig D., Klinger L., Vierling L., Zimmerman P, and Geron C. (1996) Leaf, branch, stand and landscape scale measurements of volatile organic compound fluxes from U.S. woodlands. *Tree Physiology*, 16, 17-24.
- Guenther A.B., Monson R.K. and Fall R. (1991) Isoprene and monoterpene rate variability: observations with Eucalyptus and emission rate algorithm development, *J. Geophys. Res.*, 96, No. D6, 10799-10808.
- Guenther A.B., Zimmerman P.R., Harley P.C., Monson R.K. and Fall R. (1993) Isoprene and monoterpene rate variability: model evaluations and sensitivity analyses, *J. Geophys. Res.*, 98, No. D7, 12609-12617.
- Guenther A., Zimmerman P. and Wildermuth M. (1994) Natural volatile organic compound emission rate estimates for U.S. woodland landscapes, *Atmos. Environ.*, 28, 1197-1210.

Guenther A., Hewitt C.N., Erickson D., Fall R., Geron C., Graedel T., Harley P., Klinger L., Lerdau M., McKay W.A., Pierce T., Scholes R., Steinbrecher R., Tallamraju R., Taylor J. and Zimmerman P., 1995, A global model of natural volatile organic compound emissions, *J. Geophys. Res.*, 100, No. D5, 8873-8892.

Guenther A., Geron C., Pierce T., Lamb B., Harley P. and Fall R. (1998) (in preparation) Natural emissions of volatile organic compounds, carbon monoxide, and oxides of nitrogen from North America.

Hanson U. et al. (1997), Biogenic emissions and CO₂ gas exchange investigated on four Mediterranean shrubs, *Atmos. Environ.*, 31, S1, 157-166.

Hewitt C.N. and Street R.A. (1992) A qualitative assessment of the emissions of non-methane hydrocarbon compounds from the biosphere to the atmosphere in the U.K.: present knowledge and uncertainties, *Atmos. Environ.*, 26A, No. 17, 3069-3077.

Hewitt C. N., Street R.A. and Scholefield P.A. (1997): Isoprene and monoterpene-Emitting Species Survey 1997. <http://www.es.lancs.ac.uk/es/people/pg/pas/download.html>

Isidorov V.A., Zenkevich I.G. and Ioffe B.V. (1985) Volatile organic compound in the atmosphere of forests, *Atmos. Environ.*, Vol. 19, No. 1, pp1-8.

Isidorov V.A. (1992) Non-methane hydrocarbons in the atmosphere of boreal forests: composition, emission rates, estimation of regional emission and photocatalytic transformation, *Ecological Bulletins*, Vol. 42, pp71-76.

Isidorov V.A., Povarov V.G., Klokova E.M., Prilepsky E.B., Churilova, Yu.Yu., (1993) Estimation of photochemically active VOC emission by forests of the European part of the Former USSR, In: Proc. Sixth Europ. Symp. Phys.Chem.Behav.Atmos.Pollut., Varese, Italy, 18-22 October, 1993, V.1, pp31-40.

Janson R. W. (1993) Monoterpenes emissions from Scots pine and Norwegian spruce, *J. Geophys. Res.*, 98, No. D2, 2839-2850.

Kempf K., Allwine E., Westberg H., Claiborn C. and B. Lamb (1996): Hydrocarbon emissions from spruce species using environmental chamber and branch enclosure methodes. *Atmos. Environ.* 30(9), 1381-1389.

Kesselmeier J., Schäfer L., Ciccioli P., Brancaleoni E., Cecinato A., Frattoni M., Foster P., Jacob V., Denis J., Fugit J.L., Dutaur L. and Torres L. (1996) Emission of monoterpenes and isoprene from a Mediterranean oak species *Quercus ilex* L. measured within the BEMA (Biogenic emissions in the Mediterranean area) project, *Atmos. Environ.*, 30, Nos 10/11, 1841-1850.

Kesselmeier J. et al. (1997) Emissions of short chained organic acids, aldehydes and monoterpenes from *Quercus ilex* L. and *Pinus pinea* L. in relation to physiological activities, carbon budgets, and emission algorithms, *Atmos. Environ.*, 31, S1, 119-134.

König G., Brunda M., Puxbaum H., Hewitt C.N., Duckham S.C. and Rudolph J. (1995) Relative contribution of oxygenated hydrocarbons to the total biogenic VOC emissions of selected mid-European agricultural and natural plant species, *Atmos. Environ.*, 29, No. 8, 861-874.

- Lamb B., Gay D., Westberg H. and Pierce T. (1993) A biogenic hydrocarbon emission inventory for the U.S.A. using a simple forest canopy model, *Atmos. Environ.*, 27, No. 11, 1673-1690.
- Lamb B., Hopkins B., Westberg H. and Zimmerman P. (1997) Evaluation of biogenic emission estimates using ambient VOC concentrations in Western Washington, presented at Workshop on biogenic hydrocarbons in the atmospheric boundary layer, August 24-27, 1997, University of Virginia.
- Lindskog A. and Potter A. (1995) Terpene emissions and ozone stress, *Chemosphere*, 30, No. 6, 1171-1181.
- Lübker B. and Schöpp W. (1989) A model to calculate natural VOC emissions from forests in Europe, International Institute for Applied Systems Analysis . Working paper WP-89-082.
- Ortiz A. and Dory M.A.G. (1990) The estimation of non methane VOC emissions from nature in Spain for CORINAIR inventory, In Pacyna J. M. and Joerss K.E., editors, EMEP Workshop on emission inventory techniques, Regensburg, Germany, 3-6 July, 1990. Norwegian Institute for Air Research, Kjeller, Norway, EMEP/CCC-Report 7/90.
- Owen S., Boissard C., Street R.A., Duckham C. Csiky O., Hewitt C.N. (1997) The BEMA project: screening of 18 Mediterranean plant species for volatile organic compound emissions, *Atmos Environ.*, 31, No. S1, 101-118.
- Pierce T.E. and Waldruff P.S. (1991) PC-BEIS: a personal computer version of the biogenic emissions inventory system, *J. Air Waste Manage. Assoc.*, 41, No. 7, 937-941.
- Pierce T.E. (1991) User's guide to the personal computer version of the biogenic emissions inventory system (PC-BEIS), Atmospheric research and exposure assessment laboratory, U.S. E.P.A., Research Triangle Park, NC. Report EPA/600/8-90/084.
- Pio C., Nunes T., Valente A. and Brito S. (1994) Forest emissions of hydrocarbons, annual report 1993, Part 4 BIATEX, published by EUROTRAC
- International scientific secretariat, Fraunhofer Institute (IFU), Garmisch-Partenkirchen, Germany, July 1994.
- Puxbaum H. (1997): Biogenic emissions of alcohols, esters, ether and higher aldehydes. in: Biogenic volatile organic compounds in the atmosphere, G. Helas, J. Slanina, and R. Steinbrecher (eds.), SPB Academic Publishing bv Amsterdam, pp. 79-99.
- Rasmussen R. A. (1978): Isoprene plant species list. Special report of air pollution Research Section Washington State University
- Rudolph J., Plass-Dülmer C., Benning L., Brandenburger U., Brauers T., Dorn H.-P., Hausmann M., Hofzumahaus A., Holland F., Parusel E., Ramacher B., Wahner A. Wedel A., Duckham C., Hewitt N., König G. and Puxbaum H. (1998) The POPCORN campaign 1994, an intensive field study of biogenic and man made volatile organic compounds in the atmosphere: an overview. *J. Atmos. Chem.* in press.
- Schnitzler J.-P., Lehning A., Steinbrecher R. (1997): Seasonal pattern of isoprene synthase activity in *Quercus robur* leaves and its impact on modeling isoprene emission rates. *Botanica Acta* 110, 240-243.

- Schuh et al. (1997) Emissions of volatile organic compounds from sunflower and beech: dependence on temperature and light intensity, *J. Atmos. Chem.*, 27, 291-318.
- Seufert G., Bartzis J., Bomboi T., Ciccioli P., Cieslik S., Dlugi R., Foster P., Hewitt N., Kesselmeier J., Kotzias D., Lenz R., Manes F., Perez-Pastor R., Steinbrecher R., Torres L., Valentini R. and Versino B. (1997): The BEMA-project: and overview of the Castelporziano experiments. *Atmos. Environ*, 31, No. S1, 5-18.
- Simon V., Clement B., Riba M.L. and Torres L. (1994) The Landes Experiment: monoterpenes emitted from maritime pine. *J. Geophys. Res.* 99(D8), 16501-16510.
- Simpson D., Guenther A., Hewitt C.N. and Steinbrecher R. (1995) Biogenic emissions in Europe 1. Estimates and uncertainties, *J. Geophys. Res.*, 100, No. D11, 22875-22890.
- Simpson D. (1995) Biogenic emissions in Europe 2: Implications for ozone control strategies, *J. Geophys. Res.*, 100, No. D11, 22891-22906.
- Staudt et al. (1997) Seasonal and diurnal patterns of monoterpene emissions from *Pinus pinea* (L.) under field conditions, *Atmos. Environ.*, 31, S1, 145-156.
- Steinbrecher R. (1994) Emission of VOCs from selected European ecosystems: the state-of-the-art, In et al., Borrel P., editor, *Transport and Transformation of Pollutants in the Troposphere*, Proceedings EUROTRAC symposium 1994, pages 448-455. SPB Acad. Publish. bv., the Hague, Netherlands.
- Steinbrecher R. (1996) Reaktive organische Luftkomponenten (C6-C12) anthropogenen und biogenen Ursprungs in Laub- und Nadelwldern. Abschlussbericht BMBF-Forschungsvorhaben 07 EU 816/7, pp. 88.
- Steinbrecher R. (1997) Emission factor table of BVOC of plant species in Europe, IFU-Garmisch (vers. December 1997).
- Steinbrecher R., Hahn J., Stahl K., Eichstdter G., Lederle K., Rabong R., Schreiner A.-M. and Slemr J. (1997): Investigations on emissions of low molecular weight compounds (C2-C10) from vegetation. In: *Biosphere-Atmosphere exchange of pollutants and trace substances*, S. Slanina (ed.) Springer Verlag Berlin, Vol. 4, 342-351.
- Steinbrecher R., Schürmann W., Schreiner A-M. and Ziegler H. (1993) Terpenoid emissions from Common oak (*Quercus robur* L.) and Norway spruce (*Picea abies* [L.] karst, In Slanina J. et al., editor, *Proceedings of the Joint CEC/BIATEX Workshop, Aveiro (P)*, 4-7 May 1993 on the General Assessment of Biogenic Emissions and Deposition of Nitrogen Compounds, Sulfur compounds and Oxidants in Europe, pages 251-261. CEC Research Programme Report 47.
- Street R.A. (1995) Emissions of non-methane hydrocarbons from three forest ecosystems, PhD thesis, Lancaster Univ., Lancaster, England.
- Street R. A., Duckham S. C., Boussard C. and Hewitt C. N. (1997): Emissions of VOCs from Stressed and Unstressed Vegetation. In: *Biosphere-Atmosphere Exchange of Pollutants and Trace Substances*, Slanina S. (ed.), Springer-Verlag Berlin, Heidelberg 1997, pp. 366-371.
- Street R.A., Hewitt C.N. and Mennicken S. (1997) Isoprene and monoterpene emissions from a Eucalyptus plantation in Portugal, *J. Geophys. Res.*, D13, 102, 15875-15887.

UN-ECE, 1992, *The environment in Europe and North America: annotated statistics 1992*.

Veldt C. (1988) Inventorying natural VOC emissions for the CORINAIR project, Apeldoorn, The Netherlands, MT-TNO Report 88-275; Also published in Coriniar Technical Annexes Volume 2, Default emission factor handbook, European Commission EUR 12586/2, pp101-128.

Veldt C. (1989) Leaf biomass data for the estimation of biogenic VOC emissions, Apeldoorn, The Netherlands, MT-TNO Report 89-306.

Veldt C. (1991) The use of biogenic VOC measurements in emission inventories, Apeldoorn, The Netherlands, MT-TNO Report 91-323.

Winer A. M., Fitz D. R., Miller P. R., Atkinson R., Brown D. E., Carter W. P. L., Dodd M. C., Johnson C. W., Myers M. A., Neisess K. R., Poe M. P. and Stephens E. R. (1983): Investigation of the role of natural hydrocarbons in photochemical smog formation in California. Final Report, Contract No. AO-056-32, California Air Resources Board, Statewide Air Pollution Research Center, University of California, Riverside, California 92521.

Winer A.M., Arey J., Atkinson R., Aschmann S.M., Long W.D., Morrison C.L. and Olszyk, D. (1992) Emission rates of organics from vegetation in California's central valley. *Atmos. Environ.* 26A, 2647-2659

Zimmerman P. (1979) Testing of hydrocarbon emissions of vegetation, leaf litter and aquatic surfaces and development of a methodology for compiling biogenic emission inventories, EPA 450/4-79-004.

18 BIBLIOGRAPHY

Guenther A.B., Zimmerman P.R., Harley P.C., Monson R.K. and Fall R. (1993) Isoprene and monoterpene rate variability: model evaluations and sensitivity analyses, *J. Geophys. Res.*, 98, No. D7, 12609-12617.

Guenther A., Zimmerman P. and Wildermuth M. (1994) Natural volatile organic compound emission rate estimates for U.S. woodland landscapes, *Atmos. Environ.*, 28, 1197-1210.

Guenther A., Hewitt C.N., Erickson D., Fall R., Geron C., Graedel T., Harley P., Klinger L., Lerdau M., McKay W.A., Pierce T., Scholes R., Steinbrecher R., Tallamraju R., Taylor J. and Zimmerman P. (1995) A global model of natural volatile organic compound emissions, *J. Geophys. Res.*, 100, No. D5, 8873-8892.

Hewitt C. N., Street R.A. and Scholefield P.A. (1997), Isoprene and Monoterpene-Emitting Species Survey 1997: <http://www.es.lancs.ac.uk/es/people/pg/pas/download.html>.

Simpson D., Guenther A., Hewitt C.N. and Steinbrecher R. (1995) Biogenic emissions in Europe 1. Estimates and uncertainties, *J. Geophys. Res.*, 100, No. D11, 22875-22890.

Simpson D., Winiwarter W., Börjesson G., Cinderby S., Ferreira A., Guenther A., Hewitt C.N., Janson R., Khalil M.A.K., Owen S., Pierce T.E., Puxbaum H., Shearer M., Steinbrecher R., Tarrason L. and Öquist M.G., Inventorying emissions from Nature in Europe, submitted.

Steinbrecher R. (1997) Emission factor table of BVOC of plant species in Europe, IFU Garmisch (vers. December 1997).

Veldt C. (1989) Leaf biomass data for the estimation of biogenic VOC emissions, Apeldoorn, The Netherlands, MT-TNO Report 89-306.

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20 POINT OF ENQUIRY

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SOURCE ACTIVITY TITLE:	OTHER SOURCES AND SINKS
	<i>Non-managed deciduous forests soils (excluding CO₂)</i>
	<i>Non-managed coniferous forests soils (excluding CO₂)</i>
	<i>Natural grassland and other vegetation soils (excluding CO₂)</i>
	<i>Managed deciduous forests soils (excluding CO₂)</i>
	<i>Managed coniferous forests soils (excluding CO₂)</i>

NOSE CODE:

NFR CODE: N/A

1 ACTIVITIES INCLUDED

This chapter covers emissions from non-agricultural areas that are produced biogenically in soils. Although the magnitude of emissions from soils may be perturbed and controlled by human activities, the actual processes are considered natural. This version of the chapter deals with only NO_x emissions, mainly in the form of nitric oxide (NO), which are produced by micro-organisms in soil. Natural ecosystems tend to have modest fluxes, but soils that are nitrogen-enriched, especially agricultural regions, may have NO_x fluxes approaching those of anthropogenic sources (Williams et al., 1992). Fluxes of CH₄ are not dealt with here as fluxes are expected to be to the ground, not to the atmosphere.

This chapter contains the information required to calculate emissions from soils in five SNAP categories (non-managed and managed, deciduous and coniferous, forests, and natural grasslands). Emissions from agricultural soils are covered in 100100 (cultures with fertilisers) and 100200 (cultures without fertilisers), although the methodologies are similar.

2 CONTRIBUTIONS TO TOTAL EMISSIONS

Emissions of NO_x from soils are estimated to be as much as 16% of the global budget of NO_x in the troposphere (Logan, 1983). The contribution of soil NO emissions from agricultural lands has previously been estimated to be 15% of the total European NO_x emissions inventory (Simpson et al., 1995), but emissions from non-agricultural areas are certainly much smaller than this.

These activities are not believed to be a significant source of PM_{2.5} (as of December 2006).

3 GENERAL

3.1 Description

Nitric oxide (as well as N_2 and N_2O) are produced as intermediate steps in microbial nitrification and denitrification processes. As emissions depend on the amounts of nitrogen going through these processes, agricultural soils, subject to direct fertilisation and manure, are responsible for the great majority of emissions, and in some regions may have NO_x fluxes approaching those of anthropogenic sources (Williams et al., 1992).

Soils emit NO_x mainly through biological pathways, and emission rates can be categorised by land use. The quantity of NO_x emissions from agricultural land is dependent on the rate of fertiliser application and the subsequent microbial nitrogen processing in the soil, together with a multitude of other environmental factors. A large number of studies have been discussed in relation to possible controlling factors in Skiba et al. (1997).

Although the magnitude of soil NO_x emissions may be small in overall comparison to anthropogenic NO_x emissions, there is considerable uncertainty in the estimates. Further, soil NO emissions occur in low- NO_x regions where ozone formation is most sensitive to NO_x availability, and the highest fluxes of NO occur in the warmer months of the year--times when photochemical smog is of concern.

3.2 Definitions

Soil NO emissions: nitric oxide produced by micro-organisms in soils, which ultimately "leaks" into the atmosphere.

3.3 Techniques

Current estimation techniques are based on empirical algorithms that account for land use cover and possibly N-inputs and/or soil temperature. These algorithms are based on a limited number of field chamber measurements.

Soil emissions of NO_x are dependent on the crop type and fertilisation rate and on a multitude of other environmental factors. The simple technique provides an annual estimate based upon N-inputs only, whereas the detailed technique is intended more for modelling purposes and uses the algorithm of Novak and Pierce (1993) that calculates emissions of NO based on land use and temperature.

3.4 Emissions

The current draft of this chapter considers only NO emissions. Other trace gases, such as methane, N_2O , and CO , are known to be emitted from soils but are not yet included in this methodology. For methane, soils, especially within forests, are probably a major sink so it is not clear how emissions should be defined.

3.5 Controls

Nitric oxide emissions from soils are emitted by a natural process, microbial activity in soils. This activity may be influenced by the amount of nitrogen-based fertiliser added to soils, but

this discussion is mainly relevant to agricultural emissions, as discussed in chapter 100100 and Skiba et al. (1997).

4 SIMPLER METHODOLOGY

The simpler methodology is derived from the work of Skiba et al. (1997), who suggested that 0.3% of applied N is returned to the atmosphere as NO. For non-agricultural areas applied N would consist of animal manure and atmospheric deposition. Additionally, a background N-emission rate of $0.1 \text{ ng NO-N m}^{-2} \text{ s}^{-1}$ is assumed. (For agricultural areas the fertiliser application is considered, see 100200 and 100100).

Despite its simplicity, the Skiba et al. approach has the advantage of explicitly relating emissions to applied N amounts.

5 DETAILED METHODOLOGY

Given the lack of evaluation of any parameterisations in Europe, a detailed methodology is probably not worthwhile for estimates of annual emissions. However, in case hourly changes in emissions are required (e.g. for modelling) the following methodology is proposed. Unfortunately, the simple and detailed methodologies are not consistent - they produce quite different annual estimates. However, until more is known about the merits of either method a consistent description cannot be provided.

This methodology is taken from Novak and Pierce (1993) and is known commonly as the second-version of the Biogenic Emissions Inventory System (BEIS-2). BEIS-2 can estimate NO emissions for forests, agricultural crops, urban trees, and grasslands. BEIS-2 calculates a range of emission flux rates based on land use types and soil temperature. The basis of the BEIS-2 calculation for soil NO emissions originates with the following equation (Williams et al., 1992):

$$F_{\text{NO}} = A \times \exp(0.071 \times T_s)$$

where

F_{NO}	=	NO flux, ($\text{ng N m}^{-2} \text{ s}^{-1}$)
T_s	=	Soil temperature, degrees Celsius
A	=	Experimentally derived constant for the land use types of grasslands and pasture, forests and wetlands.

The parameter A is given in Table 8.1. Emissions from soils at sub-zero temperatures can be assumed to be zero for inventory purposes.

6 RELEVANT ACTIVITY STATISTICS

For all approaches land use coverage is required, at least to distinguish agricultural and forest and other soils. For the simple methodology an estimate of N-inputs is required. This can be obtained from national deposition estimates or EMEP modelling. For the detailed approach air temperature statistics are needed.

7 POINT SOURCE CRITERIA

No point sources.

8 EMISSION FACTORS, QUALITY CODES AND REFERENCES

A large number of emission factors based on field measurement data are given in the literature. Williams et al. (1992), Yienger and Levy (1995) and Skiba et al. (1997) provide excellent reviews on these data. Emission factors are given as a function of land use and other environmental conditions, such as temperature, soil moisture, and soil nitrate levels. The variation in these measurements is considerable, resulting in a wide range of uncertainty in current emission factors. Williams et al. estimates an annual uncertainty of about a factor of three. The quality code for soil NO emission factors thus should be considered a D.

The factors required for the application of the detailed methodology (recommended for modelling hourly emissions, rather than annual) are given in Table 8.1.

Table 8.1: Empirical coefficients for BEIS-2 system, from Novak and Pierce, 1993

Land use category	A	Function to compute T_s (°C) from ambient temperature (T_a).
Grasslands + pasture	0.9	$T_s = 0.67 T_a + 8.8$
Forest	0.07	$T_s = 0.84 T_a + 3.6$
Wetlands	0.004	$T_s = 0.92 T_a + 4.4$

Valid for $0 < T_s < 35$ (°C).

9 SPECIES PROFILES

Nitric oxide (NO) is considered to be the predominant NO_x compound emitted from soils.

10 UNCERTAINTY ESTIMATES

The uncertainty of soil NO emission estimates is reported by Williams et al (1992) to be about a factor of three, when averaged over the United States on an annual basis. In view of the poor coverage of data across Europe, especially in Mediterranean areas, a factor of five uncertainty seems reasonable. Additional field studies comparing atmospheric measurements of NO_x fluxes with soil emissions derived from chamber measurements are needed to reduce this uncertainty.

11 WEAKEST ASPECTS/PRIORITY AREAS FOR IMPROVEMENT IN CURRENT METHODOLOGY

In developed areas of the world, such as Europe, the greatest uncertainty in total soil NO emissions is the amount of NO emitted from heavily-fertilised farmland. Little information is available on emissions from natural grasslands (see e.g. Skiba et al., 1997). Studies are needed to determine the fraction of nitrogen inputs that are subsequently released into the atmosphere as NO. The role of plant canopies in mitigating the flux of NO into the free atmosphere also needs to be explored.

12 SPATIAL DISSAGGREGATION CRITERIA FOR AREA SOURCES

The spatial disaggregation of soil NO emissions depends on the spatial distribution of land use, with agricultural cropland being the most important.

13 TEMPORAL DISSAGGREGATION CRITERIA

Many meteorological processes affect the temporal distribution of soil NO emissions. These processes and parameters include soil temperature and rainfall (which affects soil moisture).

14 ADDITIONAL COMMENTS

Emissions of NO from soils is both a natural and anthropogenic-influenced source. Although emissions occur as a result of biogenic processes, the magnitude of soil-generated NO is influenced by human cultivation, fertiliser application, animal management and deposition onto the soil.

A more detailed method than those suggested here has been presented in Yienger and Levy (1995) and was initially used for global soil NO_x emissions. In this approach, the variation in soil NO emissions is associated with biomass burning, history of soil moisture (pulsing), temperature, soil moisture, vegetation cover type (biome), canopy reductions, and fertilisation rate. Temperature is calculated from air temperatures using the same empirical relationships used in Novak and Pierce for wet soils, and by adding 5° C to dry soils, based on observations reported in Johansson et al., 1988. In dry soils, rather than an exponential increase, emissions increase with temperature, in a weak linear relationship.

Apart from the differences in methodology, the Yienger-Levy and Skiba approaches differ greatly in their assumption of the fraction of applied N (fertiliser, etc.) which is released as NO. Yienger and Levy assume 2.5%, whereas Skiba et al. assume 0.3% (NO as N). As the Skiba et al. figure is based upon a larger literature than Yienger and Levy and includes many European measurements it is probably a better estimate for European inventories, but the range illustrates well the uncertainties associated with this emission source.

15 SUPPLEMENTARY DOCUMENTS

16 VERIFICATION PROCEDURES

Because the emission factors are largely based on soil chambers, independent verification of the fluxes into the free atmosphere is needed. This verification is expensive, but may be accomplished using micrometeorological techniques that examine either the gradient in NO_x concentration differences with height or direct eddy correlation of NO_x fluxes. The rapid conversion of freshly-emitted NO into NO₂ in the presence of O₃ complicates the measurement and interpretation of micrometeorological flux data.

17 REFERENCES

Chameides-W, Kasibhatla-P, Yienger-J, Levy-H. (1994). Growth of continental-scale metro-agro-plexes, regional ozone production, and world food production, *Science*, vol. 264, 74-77.

Johansson-C, Rodhe-H, Sanhueza-E. (1988). Emission of NO from savannah soils during rainy season. *J. Geophysical Research*, 93, 14193-14198.

Logan-J. (1983). Nitrogen Oxides in the Troposphere: Global and Regional Budgets. *Journal of Geophysical Research*, vol. 88, 10,785-10,807.

Novak-J, Pierce-T. (1993). Natural emissions of oxidant precursors. *Water, Air, and Soil Pollution*, 67, 57-77. 340,342-352,353

Skiba, U., D. Fowler and K.A. Smith, 1997, Nitric oxide emissions from agricultural soils in temperate and tropical climates: sources, controls and mitigation options, *Nutrient Cycling in Agroecosystems*, 48, 139-153.

Williams-E, Guenther-A, Fehsenfeld-F. (1992). An Inventory of Nitric Oxide Emissions from Soils in the United States, *Journal of Geophysical Research*, vol. 97, 7511-7519.

Yienger-J, Levy-H. (1995). Empirical model of global soil-biogenic NO_x emissions. *J. Geophysical Research*, 100, 11447-11464.

18 BIBLIOGRAPHY

19 RELEASE VERSION, DATE AND SOURCE

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SNAP CODE: 1103
110301
110302

SOURCE ACTIVITY TITLE: OTHER SOURCES AND SINKS
Forest and Other Vegetation Fires
Man-induced
Other

NOSE CODE: 301.03.01
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1 ACTIVITIES INCLUDED

Burning (naturally or man-induced) of non-managed and managed forests and other vegetation, excluding agricultural burning of stubble, etc.

2 CONTRIBUTIONS TO TOTAL EMISSIONS

On a global scale biomass burning in all its forms is estimated to make very significant contributions to greenhouse gases (Andreae et al., 1988, Andreae, 1991). However, most of this burning is human-initiated, and takes place in the tropics. Of an estimated 3550 Tg CO₂ (as C) formed yearly from biomass burning, only 117 Tg C is ascribed to fires in the temperate and boreal regions, and only a small fraction of these take place in Europe (Levine, 1994).

Considering the European continent as a whole, the vast majority of these fires occur in the Eurasian part of Russia, where recent estimate suggest annual areas burnt of between 2-7 million ha (Conard and Davidenko, 1997, and references contained therein).

However, we here deal with the European part of the Russian Federation, along with the other European countries, where the area burnt is estimated at 0.5-1 million ha (range is from several years, Stannars and Bourdeau, 1995).

According to the CORINAIR-1990 inventory, forest fires account for 0.2% of European NO_x emissions, 0.5% of NMVOC emission, 0.2% of CH₄ emissions, 1.9% of CO emissions, 1.2% of N₂O emissions and 0.1% of NH₃ emissions. On the whole forest fires appear to contribute only a small percentage of emissions. However, uncertainties are very large and in some areas emissions might make appreciable contributions to ground level concentration, especially as fires occur over short periods of the time.

On a global scale biomass burning is a very significant source of CO₂ and a number of other gases to the atmosphere. However, most burning takes place in the tropical and subtropical regions, so emissions from European fires have received very little attention.

3 GENERAL

3.1 Description

Forest fires have always been a feature of forest ecosystems. However, although 'natural' forest fires may be initiated by lightning, recent estimates indicate that on a global scale almost all biomass burning is human-initiated and is increasing with time (Andreae, 1991, Levine, 1994). Much of the global emission results from so-called slash-and-burn agriculture in the tropics, but such practices are much less common in Europe. Prescribed burning, a management practice common in North America, and upon which most emission-factor measurements are based, is also not common in Europe. However, fires in Europe are still heavily associated with human influence, although often by accident rather than design.

The frequency and extent of fires in Europe is very variable from year to year, reflecting year-to-year climatological variability.

3.2 Definitions

Biomass - for forest fires the biomass of interest is the mass of all living and dead vegetation per unit area, in units of kg/m^2 . Both above-ground and below-ground biomass are significant and must be distinguished. This use of the term biomass should be distinguished from the 'foliar biomass' of importance for VOC emission estimates from foliar activity (e.g. SNAP 1101,1102).

3.3 Techniques

3.4 Emissions

The major products of biomass burning are CO_2 and water vapour. However a large number of particulates and trace gases are produced, including the products of incomplete combustion (CO , NMHCs) and nitrogen and sulphur species. These arise partly from nitrogen and sulphur contained in the vegetation and organic matter in the surface soils. Additionally, emissions can arise from the re-volatilisation of substances which have been deposited (Hegg et al., 1987, 1990).

Some emissions are not considered further here as they have little relevance for tropospheric chemistry, but they are worth mentioning for their stratospheric impacts. These include H_2 , COS and to a lesser extent CH_3Cl (Crutzen et al., 1979, Andreae, 1991).

Many other trace emissions have been measured, but which seem to contribute little to total emissions, e.g. methanesulphonate (MSA), aldehydes, organic acids (Andreae et al., 1988).

A secondary effect of fires is that emissions from the land-area after burning can be significantly enhanced relative to unburned areas. Such effects are not considered here.

3.5 Controls

Many forest fires are set deliberately or accidentally as a result of human activities. For example, data from Russia suggests that 68% of fires occur within 5 km of a road (Korovin, 1996). The main control options then consist of improved fire-prevention and fire-extinction.

Little information appears to be available on methodologies to reduce emissions during controlled forest burns. However, in the Agricultural sector it is known that time of burning and meteorological conditions have important effects on both emissions and ground level concentrations.

4 SIMPLER METHODOLOGY

From annual statistics of forest burnt one may simply multiply the area burnt by the emission factors given in Table 8.1. These emission factors are in fact identical to those given in the detailed methodology if default biome characteristics are used.

5 DETAILED METHODOLOGY

Emissions are obtained in a two-step process:

- (i) Estimate the emissions of carbon from the burned land.
- (ii) Estimate the emissions of other trace gases using emission ratios with respect to carbon.

The basic calculation of the mass of carbon emitted, $M(C)$, follows the methodology of Seiler and Crutzen (1980):

$$M(C) = 0.45 \times A \times B \times \alpha \times \beta$$

Where

0.45 is the average fraction of carbon in fuel wood

"A" is the area burnt (m^2)

"B" is the average total biomass of fuel material per unit area (kg/m^2),

" α " is the fraction of the above average above-ground biomass relative to the total average biomass B,

" β " is the burning efficiency (fraction burnt) of the above-ground biomass.

Values of B, " α " and " β " are given for relevant biomes in Table 5.1. These data are taken from Seiler and Crutzen (1980), although we have added a new forest category, "Mediterranean forest" to account for the low biomass density of this region. The " α " and " β " fractions assumed for this biome are derived from the Spanish CORINAIR 1990-93 inventories, see also Rodriguez Murrilo (1994).

Table 5.1: Biome characteristics for forest-fire emission estimates

Biome	Biomass (kg/m ²)	Above ground biomass fraction	Burning efficiency
	B	"α"	"β"
Boreal forest	25	0.75	0.2
Temperate forest	35	0.75	0.2
Mediterranean forest (1)	15	0.75	0.25
Scrubland (2)	7.5	0.64	0.5
Grassland (Steppe)(2)	2	0.36	0.5

Notes: all data from Seiler+Crutzen (1980), except:

(1) new forest category, assuming lower biomass density

(2) which is a subjective estimate, assuming burning efficiency of European grass/shrublands is less than the data on tropical biomes for which Seiler+Crutzen suggest 0.8

The emission of any particular species can then be obtained by multiplying the mass of carbon formed by the emission ratios (in g/kg C) from section 8.

As an example, if we use the factors presented above for Boreal forests, we can evaluate the mass of carbon generated in one hectare of burned boreal forest:

$$M(C) = 0.45 \times A \times B \times a \times b$$

$$= 0.45 \times 10000 \text{ m}^2 \times 25 \text{ kg/m}^2 \times 0.75 \times 0.2 = 16875 \text{ kg}$$

The emission ratio for NO_x is given in section 8 (Table 8.1) as 8 g/kg C emitted, therefore the emission factor to be applied is 135 kg NO_x/ha.

This factor is somewhat higher than the factor recommended for CORINAIR-90, namely 75 kg/ha for oceanic climate type forest, but as the background to the previous recommendation is not known we cannot discuss this further.

6 RELEVANT ACTIVITY STATISTICS

The area of forest burnt (A) must be known. The ecosystem-dependent biomass and burning "B", "α" and "β" should ideally be estimated from local data, otherwise the values given in Table 5.1 provide a default.

7 POINT SOURCE CRITERIA

No point sources.

8 EMISSION FACTORS, QUALITY CODES AND REFERENCES

Emission factors of trace gases relative to CO₂ formed in burning are given in Table 8.1, based upon the recommendations of Andreae, 1991.

Table 8.1: Emission ratios for biomass fires, relative to carbon emitted as CO₂.

	moles X per 100 moles CO ₂ emitted			g X/kg C emitted as CO ₂
	Field Measurements	Laboratory Studies	"Best Guess"	"Best Guess"
CO	6.5-140	59-105	100	230
CH ₄	6.2-16	11-16	11	15
NMHCs	6.6-11	3.4-6.8	7	21
NO _x	2-8	0.7-1.6	2.1	8
NH ₃	0.9-1.9	0.08-2.5	1.3	1.8
N ₂ O	0.18-2.2	0.01-0.05	0.1	0.4
SO _x	0.1-0.34	-	0.3	1.6

Notes: average mass of NMHC assumed to be 37, derived from speciation obtained by Radke et al., 1991. NO_x as NO₂, SO_x as SO₂.

Source: Andreae (1991)

For the simplified methodology we can use the data given in Tables 5.1 and 8.1 to calculate default emission factors per hectare of land. These default emission factors are given in Table 8.2.

Table 8.2: Default emission factors (kg/ha) for forest and vegetation fires.

	CO	CH ₄	NMVOC	NO _x	NH ₃	N ₂ O	SO _x
Boreal forest	3881	253	354	135	30	8	30
Temperate forest	5434	354	496	189	43	6	43
Mediterranean forest	1456	95	133	51	11	3	11
Shrubland	828	54	76	29	7	1.6	7
Grass/Steppe	373	24	30	13	3	0.7	3

Notes: NO_x as NO₂, SO_x as SO₂.

Quality codes for all forest fire emissions should probably be "D".

Table 8.3: AP 42 Particulate Matter Emission Factors* for forest and vegetation burning (US EPA 1996)

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Fire/Fuel Configuration	Phase ^a	Emission Factor (g/kg)			Fuel Mix (%)	Rating
		PM _{2.5}	PM ₁₀	TSP		
Broadcast logging slash						
Hardwood	F	6.0	7 ^b	13.0	33.0	A
	S	13.0	14 ^b	20.0	67.0	A
	Fire	11.0	12 ^b	18.0		A
Conifer						
Short needle	F	7.0	8 ^c	12.0	33.0	A
	S	14.0	15 ^c	19.0	67.0	A
	Fire	12.0	13 ^c	17.0		A
Long needle	F	6.0	6 ^d	9.0	33.0	B
	S	16.0	17 ^d	25.0	67.0	B
	Fire	13.0	13 ^d	20.0		B
Logging slash debris						
Dozer piled conifer						
No mineral soil ^d	F	4.0	4.0	5.0	90.0	B
	S	6.0	7.0	14.0	10.0	B
	Fire	4.0	4.0	6.0		B
10 to 30% Mineral soil ^e	S	ND	ND	25.0	ND	D
25% Organic soil ^e	S	ND	ND	35.0	ND	D
Range fire						
Juniper slash	F	7.0	8.0	11.0	8.2	B
	S	12.0	13.0	18.0	15.6	B
	Fire ^f	9.0	10.0	14.0	12.5	B
Sagebrush	F	15.0	16.0	23.0		B
	S	13.0	15.0	23.0		B
	Fire ^f	13.0	15.0	23.0		B
Chaparral shrub						
communities	F	7.0	8.0	16.0		A
	S	12.0	13.0	23.0		A
	Fire	10.0	11.0	20.0		A
Line fire						
Conifer						
Long needle (pine)	Heading ^g	ND	40.0	50.0		D
	Backing	ND	20.0	20.0		D
Palmetto/gallberry ^g	Heading	ND	15.0	17.0		D
	Backing	ND	15.0	15.0		D
	Fire	ND	8-22	ND		D
Chaparralk	Heading	8.0	9.0	15.0		C
Grasslands ^g	Fire	ND	10.0	10.0		D

^a Unless otherwise noted, determined by field testing of fires 1 acre size. F = flaming. S = smoldering.

Fire = weighted average of F and S. ND = no data.

^b For PM-10, EMISSION FACTOR RATING: C.

^c For PM-10, EMISSION FACTOR RATING: D.

^d For PM-10, EMISSION FACTOR RATING: D.

^e Determined using laboratory combustion hood.

^f Fuel mix uncertain, because of short, intense flaming phase. Use fire average for emission inventory purposes.

^g Determined using laboratory combustion hood.

* = In the absence of more appropriate data use the AP 42 emission factors

9 SPECIES PROFILES

For NMHC emissions from a number of forest fires, Radke et al. (1991) obtained an average species profile of 35% C₃H₆, 30% C₂H₆, 16% C₂H₂, 14% C₃H₈, 5% n-C₄H₁₀ (by mass).

10 UNCERTAINTY ESTIMATES

Andreae (1991) suggests that the emissions of CO₂ are uncertain by about 50% and a factor of 2 for the other trace gases. The fact that emission ratios so far determined seem to be consistent from Brazil to Canada (see Andreae, 1991, and references therein) lends some confidence to extrapolating results into Europe. However, one possible cause for concern lies in results reported by Hegg et al. (1987) which suggested that areas which had experienced substantial N-deposition emission ratios for NO_x could be an order of magnitude greater than those obtained in rural areas. Indeed, emissions of purely man-made species such as F12 are also observed from forest fires, again the result of resuspension of previously deposited pollutants (Hegg et al., 1990). Such re-suspension is very likely in many areas of Europe,

Overall, a factor of 3 uncertainty would seem a reasonable first guess for emissions of gases such as NO_x from Europe.

11 WEAKEST ASPECTS/PRIORITY AREAS FOR IMPROVEMENT IN CURRENT METHODOLOGY

Very few measurements are available of emissions from natural forest fires, and all emission rates and biome-factors reported here are based upon studies in North America or the Amazon. Evaluation of these data against European conditions should be a priority.

Despite all the complex interactions involved in forest fire emissions, the emission ratios as given in Table 2 do seem quite consistent between various workers.

The burning efficiency is here set to 0.2 for forest fires, following Seiler and Crutzen (1980). However, efficiencies of 0.76 have been reported from wild fires in Australia (Hurst et al., 1996), or 0.1 for fires in Siberia (Dixon and Krankino, 1993).

Additionally, the uncertainty in the area burned can be one of the limiting factors in establishing emissions. Estimates for Russia for example have varied by a factor of ten, partly due to the fact that official statistics do not include fires in areas not receiving fire-protection (Conard and Davidenko, 1996).

12 SPATIAL DISSAGGREGATION CRITERIA FOR AREA SOURCES

From statistics, satellite observation, etc.

13 TEMPORAL DISSAGGREGATION CRITERIA

14 ADDITIONAL COMMENTS

Some estimates of emissions from biomass burning distinguish between different phases of burning. In the 'smouldering' phase emissions tend to be higher than in the burning phase (Cofer et al., 1991), as it is the most easily combustible material which burns in the early phases. During the smouldering phase the less oxidised products (CO, HCs, etc.) are produced in higher proportions (Cofer et al., 1989, 1991). However, all phases of burning display a mixture of complete and incomplete combustion. Given the lack of data on typical European fires, and the lack of significant emissions from this source sector, such distinctions are not recommended for inventory development at this stage.

15 SUPPLEMENTARY DOCUMENTS

16 VERIFICATION PROCEDURES

17 REFERENCES

Andreae, M.O., Browell, E.V., Garstang, M., Gregory, G.L., Harriss, R.C., Hill, G.F., Jacob, D.J., Pereira, M.C., Sachse, G.W., Setzer, A.W., Silva Dias, P.L., Talbot, R.W., Torres, A.L., and Wofsy, S.C., 1988, Biomass-burning emissions and associated haze layers over Amazonia, *J. Geophys. Res.*, 93, No. D2, 1509-1527.

Andreae, M. O., 1991. Biomass burning. Its history, use, and distribution and its impact on environmental quality and global climate. In J.S. Levine (ed.), *Global Biomass Burning: Atmospheric, Climatic, and Biospheric Implications*. MIT Press, Cambridge, Massachusetts, pp. 3-21.

Cofer, W.R. III, Levine, J.S., Sebacher, D.J., Winstead, E.L., Riggan, P.J., Stocks, B.J., Brass, J.A., Ambrosia, V.G., and Boston, P.J., 1989, Trace gas emissions from chaparral and boreal forest fires, *J. Geophys. Res.*, 94, 2255-2259.

Cofer, W.R. III, Levine, J.S., Winstead, E.L., and Stocks, B.J., 1991, Trace gas and particulate emissions from biomass burning in temperate ecosystems, in *Global Biomass Burning: Atmospheric, Climatic, and Biospheric Implications*, J.S. Levine (Ed.), MIT Press, Cambridge, Mass., 203-208.

Conard, S.G. and Davidenko, E.P., 1996, Fire in Siberian Boreal Forests--Implications for Global Climate and Air Quality 1, presented at the International symposium on Air pollution and change effects on forest ecosystems, Feb. 5-9, 1996, Riverside, California.

Crutzen, P.J., Heidt, L.E., Krasnec, J.P., Pollock, W.H. and Seiler, W., 1979, Biomass burning as a source of atmospheric gases CO, H₂, N₂O, NO, CH₃C and COS, *Nature*, Vol. 282, 253-256.

Dixon, R.K. and Krankina, O.N., 1993, Forest fires in Russia: carbon dioxide emissions to the atmosphere, *Can. J. For. Res.*, Vol. 23, 700-705.

Hegg, D.A., Radke, L. F., P. V. Hobbs, and Brock, C.A., 1987, Nitrogen and sulphur emissions from the burning of forest products near large urban areas. *J. Geophys. Res.*, 92, No. D12, 14701-14709.

Hegg, D.A., Radke, L. F., P. V. Hobbs, R.A. Rasmussen, and P. J. Riggan, 1990, Emissions of some trace gases from biomass fires, *J. Geophys. Res.*, 95, No. D5, 5669-5675.

Hurst, D.F., Griffith, D.W.T., and Cook, G.D., 1996, Trace-gas emissions from biomass burning in Australia, in In J.S. Levine (ed.), *Biomass Burning and Global Change*, Vol.2, Biomass burning in South America, Southeast Asia, and Temperate and Boreal Ecosystems, and the Oil Fires of Kuwait. MIT Press, Cambridge, Massachusetts, pp. 787-792.

Korovin, G. N. 1996. Analysis of the distribution of forest fires in Russia. In: Goldammer, J.G. and Furyaev, V.V., eds. *Fire in ecosystems of boreal Eurasia*; Netherlands: Kluwer Academic Publishers; 112-128.

Levine, J. S., 1990. Global biomass burning: Atmospheric, climatic and biospheric implications. *EOS, Transactions, American Geophysical Union* 71, 1075-1077.

Levine, J. S., 1994, Biomass burning and the production of greenhouse gases, *Climate Biosphere Interaction: Biogenic Emissions and Environmental Effects of Climate Change*, Ed. Righard G. Zepp, ISBN 0-471-58943-3 Copyright 1994 John Wiley and Sons, Inc.

Radke, L.F., Hegg, D.A., Hobbs, P.V., Nance, J.D., Lyons, J.H., Laursen, K.K., Weiss, R.F., Riggan, P.J., and Ward, D.E., 1991, Particulate and trace gas emissions from large biomass fires in North America, In J.S. Levine (ed.), *Global Biomass Burning: Atmospheric, Climatic, and Biospheric Implications*, MIT Press, Cambridge, Mass., pp. 209-224.

Rodriguez Murillo, J.C. 1994, The carbon budget of the Spanish forests, *Biogeochemistry*, 25, 197-217.

Seiler, W. and P. J. Crutzen, 1980. Estimates of gross and net fluxes of carbon between the biosphere and the atmosphere from biomass burning. *Climatic Change* 2, 207-247.

Stannars, D. and Bourdeau, P. (eds.), 1995, *Europe's Environment. The Dobbris Assessment*, European Environmental Agency, Copenhagen, Denmark.

US EPA (1996) *Compilation of Air Pollutant Emission Factors Vol.1. Stationary, Point and Area Sources*. Report AP-42 (5th ed.).

18 BIBLIOGRAPHY

19 RELEASE VERSION, DATE AND SOURCE

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	110402
	110403
	110404

SOURCE ACTIVITY TITLE:	OTHER SOURCES AND SINKS
	<i>Natural Grassland and Other Vegetation</i>
	<i>Grassland</i>
	<i>Tundra</i>
	<i>Other Low Vegetation</i>
	<i>Other Vegetation (Mediterranean scrub,)</i>

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	301.04.04

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1 ACTIVITIES INCLUDED

This chapter deals with NMVOC emissions from all types of grasslands and other types of vegetation (natural, semi-natural and in some cases cultivated) which do not fit easily into the forest classification. This includes especially the Mediterranean maquis/garrigue and other low scrub-type vegetation, heathland, Tundra, etc. Table 1.1 outlines some examples within the SNAP codes.

Most of the grasslands in mid- and northern Europe are agriculturally used for either harvesting hay (meadows) or for grazing (pastures). Natural grasslands can be found in alpine regions above the timberline (alpine Tundra), at lower elevation northwards of the timberline (boreal Tundra), in dry climatic regions with poor soil (Steppe), on saltfloors and on moorland.

Low vegetation (< 5 m height), apart from grassland, is widespread across Europe. For example, in many parts of Europe moorland and heathland cover large areas, with *Erica sp.*, *Ulex sp.*, *Calluna sp.*, *Pteridium sp.*, and similar species being common. In the Mediterranean region maquis, garrigue and jaral are characteristic landscapes (see definitions, 3.2).

Many types of vegetation could be covered within this system, and many overlap with other SNAP categories. E.g. reed vegetation, or maritime coastal vegetation (halopsammophytic), could be included here or under SNAP1105 (wetlands). Unfortunately, we have no information on emission rates yet, so this particular problem does not arise. More importantly, methods are given here for calculating VOC emissions from Agricultural crops such as wheat, as the methodology is identical to that for other vegetation. These emissions should be entered under SNAP-level 10-Agriculture, and not SNAP-11 - Other sources and sinks.

Emissions of N₂O are assumed to follow IPCC methodologies and so are not covered here. Emissions of NO_x from the soil are dealt with in a separate chapter covering all types of forests and grasslands. Emissions arising from fires are covered in SNAP 1103 (Forest and other vegetation fires). Emissions of CH₄ should strictly be treated also in the soils section, but in any case are assumed to be zero (the flux is probably to the ground, not to the atmosphere).

Table 1.1: Classification scheme for grassland and other non-forest vegetation in Europe

110401	Natural grassland Pastures, Meadows, Steppe
110402	Tundra alpine Tundra, boreal (treeless) Tundra
110403	Other low vegetation Heathland, Moorland Miscellaneous dwarf shrub vegetation(Garrigue etc.)
110404	Other vegetation Maquis
100205	Grasslands (Agricultural) Agricultural grassland of low and medium productivity (< 8 t ha ⁻¹ yr ⁻¹ yield), Agricultural grassland with high productivity (> 8 t ha ⁻¹ yr ⁻¹ yield)

2 CONTRIBUTIONS TO TOTAL EMISSIONS

According to the CORINAIR-1990 inventory, natural grassland accounts for 0.6% of European NMVOC emission, 0.4% of CH₄ emissions, 2.9% of N₂O emissions and 0.3% of NH₃ emissions. However emission rates for NMVOC need to be substantially revised. Simpson et al., 1998, using the recommended defaults in this chapter estimate that pastures and meadows may contribute nearly 1 Mt to European NMVOC emissions (ca. 4%) and crops also ca. 1 Mt. Uncertainties are still very large and in some other areas emissions might be appreciable, e.g. the NH₃ emission from pastures (due to animal droppings) and meadows (in particular when fertilised with manure).

The area coverage of grasslands in Europe is second highest after forests, however the biomass density of grasslands is often lower than the foliar biomass density of forests.

The emissions from other low vegetation were not covered in CORINAIR-90/94, however their emissions may have been included by some countries under the "forest" SNAP codes.

These activities are not believed to be a significant source of PM_{2.5} (as of December 2006).

3 GENERAL

3.1 Description

Emissions of NMVOC from plants are usually divided into isoprene, monoterpene, and OVOC (other VOC) emissions for inventory purposes. In general isoprene and monoterpene emissions are the most photochemically reactive and hence of most interest for ozone studies. However, for grasslands the major emission probably consists of OVOC, and these may be significant in mass terms.

Only a small number of screening studies have been undertaken until now to survey biogenic VOC emissions from non-forest vegetation.

Hewitt & Street (1992) tested the 21 most abundant grass and herbaceous species in the U.K with a qualitative method. Only purple moor grass (*Molinia caerulea*), bracken (*Pteridium aquilinum*) and common gorse (*Ulex europaeus*) were found to emit isoprene, and only ivy (*Hedera helix*) and cocksfoot grass (*Dactylis glomerata*) found to emit monoterpenes. 28 species of agricultural crops were also screened, of which only blackcurrant (*Ribes nigrum* v. Ben Sark and Ben Lomond) produced any significant emissions. The only major species, in terms of abundance, of which no varieties were tested was winter barley (*Hordeum vulgare*). As pointed out by Hewitt & Street, genetically different varieties of the same species may display different emission characteristics to those found above, but these data strongly suggest that isoprene as well as terpene emissions from crops and grasses are not important in the U.K. This result supports the previous findings that grasses and grass related crops are generally no or low emitters for isoprene and terpenes. (However, other VOC emissions are probably very significant).

A compilation of biogenic VOC emissions from crops and "hay" (meadows for hay production) from US sites is found in Lamb et al. (1993). Except for Tomato and Maize (Corn) all agricultural crops tested as well as "hay" were classified as "low emitters" (emission rate of all VOC determined $< 1 \mu\text{g g}^{-1} \text{h}^{-1} \text{DW}$) in the Lamb et al. (1993) study (Table 8.4). However, in their compilation the chemical nature of VOC other than isoprene and terpenes (OVOC) has not been specified, though for several crop species the OVOC emission rate exceeded the emission rates of the "classical" biogenic VOCs isoprene and terpenes. Arey et al. (1991) and Winer et al. (1992) investigated emission rates from a number of agricultural crops and from a perennial grassland plot in the US in more detail (Table 8.1). They specified the "OVOCs" and found (*Z*)-3-hexenol ("leaf alcohol") and (*Z*)-3-hexenylacetate ("leaf ester") in many crops as the two most dominant compounds of the group of "OVOCs".

Very little is known of emissions from shrubs and bushes, except that obtained as a result of a limited number of intensive field campaigns held at a few locations in the north-western Mediterranean region, as part of the Biogenic Emissions in the Mediterranean Region (BEMA) project (e.g. Owen et al, 1997) and in the U.K. (e.g. Cao et al., 1997). Additionally a limited amount of screening work has been carried out on these ecosystems (e.g. Hewitt and Street, 1992).

However, the vegetation species found in these ecosystems are often very aromatic and hence may be expected to emit a very wide and complex range of volatile organic compounds. This is especially so for Mediterranean vegetation. By far the majority of efforts to date have been focused on the emissions of isoprene and monoterpenes, so it is difficult to quantify the emissions of these other VOC, including the oxygenated compounds. Additionally, nothing is known of the emission of nitrogen and sulphur compounds from these plants.

There are very few data about VOC emissions from single herbaceous species which may occur in certain areas in relatively large quantities. An example is *allium ursinum* (wild garlic) which grows in mid and northern Europe in beech and other mixed hardwood forests in spring with biomass densities up to 300 g m⁻². Although wild garlic emits no isoprene and only a little of terpenes, the emission rate of OVOCs was found to be 2.6 µg g⁻¹ h⁻¹ DW (Puxbaum & König 1997) and thus it can be classified as "high OVOC - emitter". Similarly Tanner & Zylinska (1994) found relatively high emission rates of oxygenated terpenoids (> 4 µg g⁻¹ h⁻¹ DW) in undercover vegetation (tarweed) in the San Joaquin Valley. Although these examples are for species under forests - and not in grasslands, they might indicate that there might be grassland biomes which contain herbaceous plants with higher emission rates than those to be recommended in section 8.

König et al. (1995) tested VOC emissions from agricultural plants such as wheat, rye, rape, grape and three types of grassland in East Austria. They used the Arey et al. (1991) approach to include also specified OVOC emissions. In terms of prevalence of one of the groups of emitted VOCs (isoprene, terpenes, OVOC) wheat, rye, oilseed rape, grape and two of the grass plots examined were "OVOC" - emitters. However, for one of the examined grass plots, terpene and OVOC - emissions were of equal importance (grassland A3, Table 8.2). After mowing of one of the grass plots the emissions of terpenes and OVOCs increased roughly by a factor of three. The same group performed a measurement campaign in Northern Germany in 1995. Although the results are not published until now, we include the data for an examined grass plot (grassland G in Table 8.2, Puxbaum et al., in preparation). The data fit well into the results for grasslands from the above mentioned studies from the US and Austria.

3.2 Definitions

OVOC - Other non-methane VOC, excluding isoprene and terpene. Usually used to encompass a wide range of emitted VOC - see section 9.

BMD - Biomass density (g m⁻² DW) averaged over vegetation period

DW - dry weight of plants(used for emission rates), as opposed to fresh weight.

NPP - Net primary production (g Carbon m⁻² yr⁻¹) Buildup of biomass carbon during a year

PAR- Photosynthetically active radiation, typically about 45-50% of total global radiation.

Grasslands are areas which are dominated by grassy plants, but usually also containing other herbs. There are mainly two families of grassy plants: *poaceae* ("sweet grasses") and *cyperaceae* ("acidic grasses"), the first of the two occurring most frequently in European grasslands.

Tundra - vast level treeless (almost) Arctic region where subsoil is frozen.

Names and definitions of Mediterranean landscape classes vary from country-to-country and from author-to-author (Di Castri et al., 1981). However the following are in common usage:

Maquis

- also known as matorral denso, espinal (ES), chaparral (UK, USA), macchia alta (IT).
- comprising evergreen shrubs and small trees, typically olive (*Olea oleaster*), carob (*Ceratonia siliqua*), dwarf *Quercus ilex* and *Erica multiflora*

Garrigue

- also know as matorral claro (ES), scrub (UK), macchia bassa (IT)
- comprised of mid-height shrubs, 0.6-2 m high on calcereous soils, typically *Pistachia lentiscus*, *Arbutus unedo*, *Myrtus communis* and *Ulex sp.*

Garrigue is sometimes used for vegetation less than 0.6 m high also, in which case **lande** (FR), **tomillar** (ES), **gairriga** (IT), **phyrgana** (GR) are alternative names.

Jaral

- similar sized shrubs on siliceous soils, e.g. *Erica sp.*, *Cistus sp.*

3.3 Techniques

Not applicable

3.4 Emissions

This chapter deals with NMVOC - emissions of grassland and other low vegetation including crops. As for forests, NMVOC species are classified into three groups: Isoprene, Terpenes (Mono- and Sesquiterpenes), OVOC (other VOC). The composition of OVOC is discussed in section 9.

3.5 Controls

Principally there seems to be no control to natural emissions by definition, however land-use changes obviously can significantly affect total emissions (e.g. very early changes which date back to the bronze age, when forests were cleared in Europe to gain agricultural land and meadows, or more recent changes due to nitrogen deposition where heather gets converted to grassland).

4 SIMPLER METHODOLOGY

Grasslands and other low vegetation ecosystems consist generally of plant communities (except for crops which are usually monocultures). Often a few species dominate the

community. We introduce a classification scheme for grassland and other low vegetation in Europe (Table 8.2), but emission rate data for grasslands are not available for the different types. Similarly for Mediterranean shrubland landscapes it does not seem possible at present to subdivide into species for inventory purposes, and northern moorlands and heathlands must also be treated together.

We therefore recommend the use of ecosystem-wide rates for dealing with these types of vegetation. However, we give emission rates for individual species where known. These may be used to devise more appropriate ecosystem rates for particular regions, or to conduct a species-specific approach if desired.

For all types of vegetation, an appropriate system describing the emissions flux on an hourly basis is that of Guenther et al. (1996):

$$\text{Flux } (\mu\text{g m}^{-2} \text{ yr}^{-1}) = \int \varepsilon \cdot D \cdot \gamma \, dt \quad (1)$$

where ε is the average emission potential ($\mu\text{g g}^{-1} \text{ h}^{-1}$) for any particular species, "D" is the foliar biomass density (g dry weight foliage m^{-2}), and γ is a unitless environmental correction factor representing the effects of short-term (e.g. hourly) temperature and solar radiation changes on emissions. For isoprene emissions, and light-activated terpene emissions (so far only quantified for two forest species, *Picea abies* and *Quercu ilex*), γ is a function of light and temperature, and is denoted γ -iso. Terpene and OVOC emissions from most vegetation types are simply dependant on temperature, in which case γ is temperature-only dependant, and denoted γ -mt.

The simplified methodology consists of modifying equation (1) to be a seasonal rather than an hourly calculation.

$$F = \varepsilon \cdot D \cdot \Gamma \quad (2)$$

Where Γ represents the integrated value of γ over the growing season of the vegetation concerned.

The total emission from an area is then obtained in a detailed methodology by calculating F every hour for each vegetation category and group of VOC compounds and multiplying by the appropriate areas.

Using meteorological data from the EMEP MSC-W models the integrated values, Γ -iso and Γ -mts, have been calculated for both 6 monthly (May-October) and 12 monthly growing seasons, as averages over each country. These have been calculated from hourly γ values, and thus have units of hours. The Γ values are tabulated in Table 4.1. With this simplified methodology we could estimate for example the OVOC emissions from 1 km^2 of grassland as simply:

$$\begin{aligned} \text{Emission} &= \text{Area} \times \varepsilon \cdot D \cdot \Gamma\text{-mts} \\ &= 1000\,000 \text{ m}^2 \times 1.5 \mu\text{g g}^{-1} \text{ h}^{-1} \times 500 \text{ g m}^{-2} \times \Gamma\text{-mts h} \end{aligned}$$

For Austria, for example, Table 4.1 gives Γ -mts (= Γ -ovoc) for 6-months as 588, therefore we have:

$$\text{Emission} = 1000\ 000\ \text{m}^2 \times 1.5\ \mu\text{g}\ \text{g}^{-1}\ \text{h}^{-1} \times 500\ \text{g}\ \text{m}^{-2} \times 588\ \text{h} = 441\ \text{kg}\ \text{km}^{-2}$$

Table 4.1: Country average values of integrated environmental correction factors, Γ -iso and Γ -mts for 6 and 12 month growing seasons (unit = hours).

	Γ -mts (= Γ -ovoc)		Γ -iso	
	6-month	12-month	6-month	12-month
Albania	745	976	563	719
Austria	588	734	452	540
Belarus	753	895	581	684
Belgium	739	969	580	712
Bosnia Herzegovina	709	893	561	686
Bulgaria	824	1029	620	755
Croatia	883	1121	667	815
Czech_Republic	712	885	533	633
Denmark	518	704	373	485
Estonia	565	669	422	491
Finland	458	523	339	379
France	840	1107	669	829
Germany	698	890	525	632
Greece	1076	1440	816	1057
Hungary	966	1188	730	874
Ireland	467	713	337	478
Italy	904	1208	711	902
Latvia	636	757	486	572
Lithuania	675	813	516	613
Luxembourg	786	1003	620	745
Macedonia,F.Y.R.	631	783	492	597
Moldova, Rep. of	858	1040	649	771
Netherlands	676	901	513	643
Norway	327	397	240	284
Poland	736	912	558	669
Portugal	1015	1388	853	1093
Romania	783	964	587	706
Russia, Fed.	808	917	637	717
Slovakia	797	977	607	724
Slovenia	745	940	562	682
Spain	982	1301	806	1004
Sweden	423	508	315	368
Switzerland	465	580	368	432
Turkey	976	1263	783	983
United_Kingdom	493	720	358	492
Ukraine	856	1023	656	771
Yugoslavia	752	937	557	674

5 DETAILED STATE OF THE ART METHODOLOGY

For a more detailed calculation the environmental correction factors (γ -iso, γ -mts) may be calculated explicitly on a monthly or an hourly basis if relevant meteorological data are available. The procedure is identical to that presented for forest emissions in 1101, 1102, and is not repeated here.

6 RELEVANT ACTIVITY STATISTICS

The relevant statistics are vegetation cover, foliar biomass density, and possibly monthly and/or hourly temperature and radiation parameters if the detailed methodology is to be pursued.

Vegetation coverage in terms of the vegetation types discussed in section 8 are required, together with foliar biomass estimates (D), and estimates of growing seasons. Grasslands are found in land use statistics generally under grassland, pastures and possibly meadows. Care should also be taken not to double count species/vegetation types.

We have not found a comprehensive discussion of biomass densities. However, there is information about the annual net primary production (Ruimy et al., 1994, Lieth and Whittaker, 1975). Data for natural grasslands are compiled in Table 6.1.

Table 6.1: Compilation of Annual Net Primary Production for Grasslands ($\text{g C m}^{-2} \text{ yr}^{-1}$) according to Ruimy et al. (1994) and Lieth and Whittaker (1975) and estimate for default biomass density values. BMD = Biomass density.

	"REF" Ruimy et al.	"MEAN" Lieth&W	default BMD $\text{g m}^{-2} \text{ DW}$
Tundra	100	50	100
Savanna	530	400	500
Temperate grasslands	470	300	450

Notes: The default biomass densities (BMD) are derived in the following way: The net primary production is the build up of new biomass in the vegetation period in $\text{g C m}^{-2} \text{ yr}^{-1}$. The conversion factor from C (Carbon) to biomass DW is 2.2 (Ruimy et al., 1994). It is assumed that 50 g m^{-2} biomass is remaining from the past year. The new (at the end of the vegetation period) and old biomass value is averaged over the vegetation period.

For Alpine grasslands the following defaults are recommended:

	D ($\text{g m}^{-2} \text{ DW}$)
Alpine pastures above timberline:	50
Mid productivity alpine grassland (1-3 cuts)	200

For heathlands and moorlands very little data is available. We recommend a default based upon the biomass density of Gorse, which is widespread in the UK, assuming 50% coverage:

	D ($\text{g m}^{-2} \text{ DW}$)
Heathland /moorland	175

For Mediterranean scrublands, the following are recommended:

Maquis	400
Garrigue/low-scrubland	200
Monte-hueco*	100

* mixed pastures and trees, mainly Holm and Cork oaks.

For Karelian (Russian) spruce forests (marshy-grassy-types) the following data are available. The ground-biomass densities of Russian forests are probably larger though than those of many managed forests in other parts of Europe:

	D (g m ⁻² DW)
Forest grass-biomass	90
Forest ferns	14
lichen+mosses	100-300
shrub, including berries	10-30

For a more detailed evaluation of D, local information is required. Some insight about the biomass of meadows can be obtained from harvest yields of hay. In Austria the yield of hay is of the order of 6-10 t ha⁻¹ and more, which is equivalent to 600-1000 g m⁻² harvested dry biomass. The frequency of harvests per year is 1-6, depending of altitude and fertilisation. If we assume that after harvest a biomass density of 50 g m⁻² remains, and there is a linear growth rate between harvests, the annual average biomass density is estimated to 200-300 g m⁻². This number is for medium productive grassland in the alpine region and is lower than Lamb et al's (1987) estimate for meadows for hay production in the US of 540 g m⁻². However, in highly productive grasslands in flat terrain in Europe the biomass density might be as high as the Lamb et al. 1987 estimate.

For agricultural grassland an estimation of the vegetation period averaged biomass density can be obtained by the following formula:

$$\text{BMD} = [\text{Y} * 100/2 * \text{n}] + 50$$

Where BMD is biomass density (g m⁻² DW), Y is yield of biomass DW per vegetation period (t ha⁻¹), n is the number of cuts per year, and 50 is the biomass remaining after cutting

7 POINT SOURCE CRITERIA

No point sources

8 EMISSION FACTORS, QUALITY CODES AND REFERENCES

NMVOC

For grassland areas the emission potentials (ϵ , equation 1) should be given as standardised values at 30 °C and for full sunlight=1000 $\mu\text{mole photons m}^{-2} \text{s}^{-1}$ PAR. A summary of default ecosystem-type emission factors which seem appropriate especially for European species is given in Table 8.1 below

Table 8.1 Ecosystem-default emission potentials (ϵ) for isoprene (iso), terpenes (mts) and OVOC, and biomass densities. ϵ ($\mu\text{g g}^{-1} \text{DW h}^{-1}$) is given for 30°C and 1000 $\mu\text{mol PAR}$.

Ecosystem	D ($\text{g m}^{-2} \text{DW}$)	ϵ -iso	ϵ -mts	ϵ -ovoc	Main reference
Grass	400*	0	0.1	1.5	K
Maquis	400	8	0.65	1.5	O,G95
Garrigue	200	8	0.65	1.5	O, G95
Monte-hueco	100	1	10**	1.5	***
Moorland/heathland	350	8	0.65	1.5	C

Notes: * but see section 6 for Alpine grasslands; ** Calculate with γ -iso. All other terpenes with γ -mts; *** Assumes ca. 50% *Q.illex*, 50% *Q.suber*; K=König et al., 1985, C=Cao et al., 1997, G95=Guenther et al., 1995, O=Owen et al. 1997

It is important to note that the 1.5 $\mu\text{g g}^{-1} \text{h}^{-1}$ rate for OVOC given here is a default (from Guenther et al., 1995) with a wide uncertainty range. Almost all measurements have used methods for the determination of VOCs, which are not capable of finding and quantifying polar compounds with less than four carbon atoms (e.g. methanol, formaldehyde, etc.). For this reason the OVOC results in these tables also list separately OVOC $\geq \text{C}_4$ and OVOC $< \text{C}_4$ if known. Recently it was shown that plants may emit methanol (MacDonald & Fall 1993), low molecular weight aldehydes (Kotzias et al. 1997) and low molecular weight organic acids (Bode et al. 1997) in relevant quantities. For grasslands and crops no data are available however to quantify this further.

The following tables give species-specific emission potentials and some more detail about the OVOC split.

Table 8.2: Species-specific emission potentials in $\mu\text{g g}^{-1} \text{h}^{-1}$ dry weight of plants for grassland (SNAP 110401) normalised to 30°C, PAR levels not specified

	Biomass density g m^{-2}	ϵ -iso	ϵ -mts	ϵ -ovoc	OVOC split ($>C_4, <C_4$)	Refs
USA:						
"hay"	540	0.07	0.175	1.5	(0.11*,n.a.)	L
Grassland [#]		n.d.	0.015	1.5	(0.06,n.a.)	A/W
Europe:						
Grassland A1		0.001/C [§]	0.02	1.5	(0.015,n.a.)	K
Grassland A2	300	0.002/S [§]	0.015	1.5	(0.06,n.a.)	K
Grassland A3	420	0.003/S [§]	0.07	1.5	(0.08,n.a.)	K
Grassland A3m (after mowing)	420	0.002/S [§]	0.20	1.5	(0.27,n.a.)	K
Grassland G	230	n.d./xx [§]	0.03	1.5	(0.15,n.a.)	P/
Recommended default for grassland	400**	0	0.1	1.5		

Notes: n.d. not detected; n.a. not analyzed; S[§] measured under sunny conditions; C[§] measured under cloudy conditions; *OVOC not specified; ** see also section 6. ; # perennial natural grassland; A1) grassland under oak forest; A2) grassland with no flowers, 35 cm height; A3) grassland with some flowers, 25 cm height; A3m) grassland 3 mowed; G grassland in northern Germany.

Refs: L: Lamb et al. 1987, 1993, A/W: Arey et al. 1991, Winer et al. 1992, K: König et al. 1995, S: Street et al. 1997, P: Puxbaum and König 1997, P/: Puxbaum et al. in preparation, Tanner and Zylinska 1994.

Table 8.3: Species-specific emission potentials in $\mu\text{g g}^{-1} \text{DW h}^{-1}$ for shrubs and low vegetation, standardised to 30 °C and 1000 $\mu\text{mol m}^{-2} \text{s}^{-1}$ PAR.

NOTE: Many tree species can be shrub like - their emission potentials may be found for these in the chapter covering forests (SNAP1101,etc.)

Common name (example)		ϵ -iso.	ϵ -mts	ϵ -ovoc\$	References	
					Iso	Terp.
Wild garlic	Allium_ursinum	0	0	3	P	P
-	Anthyllis	0.1	0.2	1.5	O	O
Strawberry tree-	Arbutus	0.1	0.2	1.5	B3,O	O
-	Arundo	60	0.2	1.5	O,H90,B3	B3
	Artemisia	0	0.2	1.5	P/	P/
Dwarf_boxwood	Buxus	10	0.2	1.5	O,B3	B3
Carob	Ceratonia	0.1	0.65	1.5	O	O
	Chrysanthemum	0.1	0.65	1.5	O	O
Rockrose	Cistus	0.1	0.2	1.5	O	O
Broom	Cytisus	20	0.2	1.5	S:Sf;B3	B3,-O
Tree_heath	Erica	5	0.2	1.5	S:Sf;O	B3,O
	Helichrysum	0.1	3	1.5	O	O
Tarweed	Holocarpa	0	3	3	TZ	TZ
Juniper	Juniperus	0.1	0.65	1.5	S:Sf;B3	B3,O
Lavender	Lavendula	0.1	0.65	1.5	S	O
Common_myrtle	Myrtus	34	0.2	1.5	S:Sf;B3	O
	Phillyrea	0.1	0.65	1.5	O	O
	Rhamus	20	0	1.5	O,B3	B3
Rosemary	Rosmarinus	0	1.5	1.5	S:Sf	Ha
Sage	Salvia	0.1	1.5	1.5	B3	B3
Broom	Spartium	5	0.2	1.5	S:Sf;O	B3,O
Bil/blueberry	Vaccinium	0.1	0	1.5	B3	B3
Gorse	Ulex	8	0.65	1.5	S:B,i6;C;B3	B
Grape	Vitis	0.1	0.1	1.5	B3	A,-K

Notes: \$We have used a default ϵ -OVOC of 1.5 $\mu\text{g g}^{-1} \text{h}^{-1}$ for all compounds except Wild garlic, which includes 1.6 $\mu\text{g g}^{-1} \text{h}^{-1}$ oxygenated compounds, 1.0 $\mu\text{g g}^{-1} \text{h}^{-1}$ h carbon from sulphurous organics, and tarweed for which TZ give higher rates. References as for Table 8.1, plus B3=Guenther et al. 1998, O=Owen et al.,1997, Sf=Seufert et al., 1997.

Table 8.3: Emission rates in $\mu\text{g h}^{-1} \text{g}^{-1}$ dry weight of plants for crops, normalised to 30°C, PAR levels not specified

	Biomass density g m^{-2}	ϵ -iso	ϵ -mts	ϵ -ovoc	Measured OVOC split ($>C_4, <C_4$)	Authors
Wheat (<i>Triticum</i>):						
USA	740	0.002	0.008	1.5	(0.03*,n.a.)	L
USA	n.a.	0	0	1.5	(0.05,0.5*)	A/W
Europe (after bloom)	800	0/S [§]	0	1.5	(0.016,n.a.)	K
Rye (<i>Secale</i>):						
USA	2430	0.003	0.008	1.5	(0.005*,n.a.)	L
Europe	400	0/S [§]	0.10	1.5	(0.25,n.a.)	K
Barley (<i>)</i>:						
USA	1290	0.006	0.015	1.5	(0.009,n.a.)	L
Oats (<i>Avena</i>):						
USA	750	0.01	0.026	1.5	(0.0015*,n.a.)	L
Recommended default for grass related crops	800	0.002	0.1	1.5		
Other crops:						
"High emitters":						
Maize/Corn US	1610	0	0.22	1.5	(0.88*,n.a.)	L
Maize/Corn Europe	n.a.	0	0	1.5	6.4 1.0	S R
Tomato (S.)	n.a.	0	13.2	1.5	(0.4,n.a.)	A/W
Tomato (C.)	n.a.	0	21.8	1.5	(1.2,n.a.)	A/W
Miscellaneous						
Alfalfa	3250	0.005	0.2	1.5	(0.6,n.a.)	L, A/W
Safflower	n.a.	0	0.3	1.5	(0.7,n.a.)	A/W
Sorghum	3180	0.002	0.03	1.5	(1.0,n.a.)	L, A/W
Rice	1050	0.10	0.24	1.5	(0.15*,n.a.)	L
Tobacco	490	0	0.12	1.5	(0.48*,n.a.)	L
Soybeans	740	0.03	0	1.5	(n.a.,n.a.)	L
Sunflower	n.a.	0.05	0.7	1.5	(0.3,n.a.)	SCH
Oilseed Rape	400	0/S [§]	0.12	1.5	(0.23,n.a.)	K
Grape:						
USA (T.S.)	n.a.	0	0	1.5	(1.4,n.a.)	A/W
USA (F.C.)	n.a.	0	0.07	1.5	(1.3,n.a.)	A/W
Europe (Ch.)	410/l.o.	0.002/S [§]	0.002	1.5	(0.05,n.a.)	K
Misc. crops default	1335	0.09	0.13	1.5	(0.6,0.9)***	

Notes: 0 not detected. n.a. not analyzed. S[§] measured under sunny conditions.

*OVOC not specified; *** the OVOC $<C_4$ emission rate is a guess based on very little data l.o. leaves only.

Refs: L: Lamb et al. 1987, 1993, A/W: Arey et al. 1991, Winer et al. 1992, K: König et al. 1995, S: Street et al. 1997, R: Rudolph et al. (in press).

9 SPECIES PROFILES

As with forest NMVOC emissions, biogenic emissions from grasslands consist of a wide variety of species, including isoprene, monoterpenes, (alpha-pinene, beta-pinene, limonene, etc.), and 'other' VOC. The 'other' VOC (OVOC) species consist of a large number of oxygenated compounds (alcohols, aldehydes, etc.), and have proven difficult to quantify in atmospheric samples. Progress in quantification of OVOC from European vegetation has been made recently (König et al. 1995, Puxbaum 1997), although many more measurement data will be required before reliable attempts to inventory specific OVOC can be made.

Section 8 has already presented separate emission rates for isoprene, terpenes, and <C₄, >C₄ OVOC. However, within each of these groups a wide range of species are emitted, as indicated by Table 9.1.

Table 9.1. Main emitted single VOC species (Rank 1-3) emitted from grassland plots and various crops (compiled from König et al. 1995 and Puxbaum et al. in preparation). Note that <C₄ organic compounds have not been determined in these studies.

Plot	Rank 1	Rank 2	Rank 3
Grassland A1	a-Pinene	Leaf ester	Hexanal
Grassland A2	Leaf ester	Leaf alcohol	Limonene
Grassland A3	Leaf ester	1,8-Cineol	a-Pinene
Grassland G	Pentanal	Leaf ester	Limonene
Wheat	Leaf ester	Hexanal	2-Pentanone
Rye	1-Hexanol	Leaf alcohol	2-Methyl-1-propanol
Rape	Leaf ester	Limonene	Sabinene
Grape (Chardonnay)	Butanone	Leaf ester	Hexanal

Leaf ester: (Z)-3-hexen-1-ol-acetate, Leaf alcohol: (Z)-3-hexen-1-ol

10 UNCERTAINTY ESTIMATES

With so few data it is very difficult to quantify the uncertainties. Quality codes for all grassland vegetation should probably be "E".

11 WEAKEST ASPECTS/PRIORITY AREAS FOR IMPROVEMENT IN CURRENT METHODOLOGY

Very few measurements are available of VOC emissions from natural grasslands. Emissions of "other" VOC < C₄ in particular (see e.g. aldehyde and ketone emissions from corn as determined by Street et al. 1997) are possibly significant but virtually unquantified for grasslands. In the same way emissions of "other" VOC < C₄ for crops are unknown.

More data are needed about NMVOC emissions for major grassland and shrub-type biomes in Europe, in particular also from Northern, Eastern, and Southern Europe. E.g. there is hardly any information about heather, Tundra, grasslands in the mountainous regions in Northern Europe with ferns and other scrub, alpine pastures, Steppe, etc.

12 SPATIAL DISAGGREGATION CRITERIA FOR AREA SOURCES

Given by vegetation coverage and climate.

13 TEMPORAL DISAGGREGATION CRITERIA

Given by equation (1) if required. It has to be kept in mind, that grassland vegetation in Europe is generally perennial, although in some cases snow-covered. However no data about emissions in the cold season are known.

A detailed treatment could also take into account the changes in biomass density over the growing season. Methods are given in Guenther et al., 1995, for many vegetation types which allows for the gradual changes in biomass dependant on NPP. A more extreme temporal variation is caused by cutting on agricultural or semi-natural areas: an example an alpine meadow at lower elevation with 3 cuts in a season is shown in Figure 13.1.

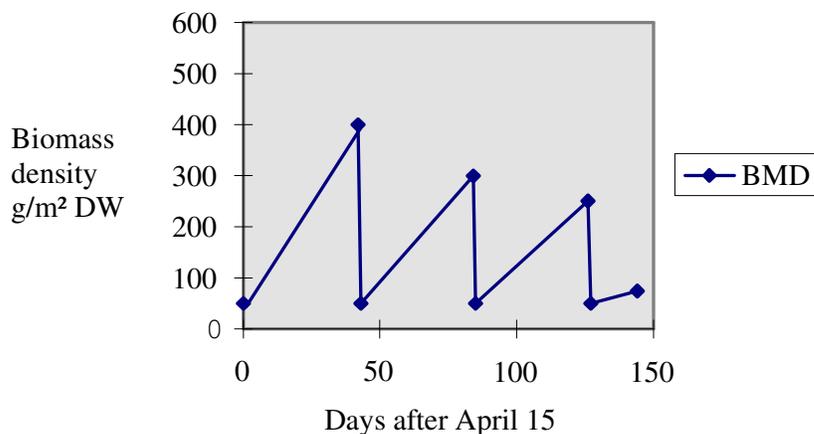


Figure 13.1: Biomass development on a 3-cut alpine meadow

14 ADDITIONAL COMMENTS

For modelling purposes the terpenes and OVOC emissions have to be speciated in some way. As the major emission is OVOC, one could use methanol for the $<C_4$ OVOC and hexenylacetate for $>C_4$ OVOC.

15 SUPPLEMENTARY DOCUMENTS

A land surface characterisation for global mapping purposes is described by DeFries et al. (1995). An interesting feature in this approach is the discrimination between C_3 and C_4 plants.

The American Biogenic Emission Inventory System (BEIS) has an extensive list of emission potentials. The latest version is documented by Guenther et al. (1998).

16 VERIFICATION PROCEDURES

If satellite data have been used in the land-use mapping process it is essential that these are independently verified by on-the-ground surveys. Large errors are possible in the identification of vegetation types and biomass from remote sensing methods.

In general all of the emission potentials are built upon very few data. More measurements are required of at least the major sources, and several different measurement techniques need to be applied in order to eliminate the artefacts (enhanced emissions) easily generated by disturbances to the vegetation.

17 REFERENCES

Arey, J., Winer, A.M., Atkinson, R., Aschmann, S.M., Long, W.D., and Morrison, C.L., 1991, The emission of (Z)-3-hexen-1-ol, (Z)-3-hexenylacetate and other oxygenated hydrocarbons from agricultural plant species. *Atmos. Environ.* 25A, 1063-1075.

Bode, K. Helas, G., Kesselmeier, J., 1997, Biogenic contribution to atmospheric organic acids. In: Helas, G., Slanina, J., and Steinbrecher R. (eds.) *Biogenic volatile organic carbon compounds in the atmosphere*, pp.79-99, SPB Academic Publishing, Amsterdam.

Cao, X.-L., Boissard, C., Juan, A.J., Hewitt, C.N., and Gallagher, M., 1997, Biogenic emissions of volatile organic compounds from gorse (*Ulex europaeus*): Diurnal emission fluxes at Kelling Heath, England, *J. Geophys. Res.*, 102, No. D15, 18903-18917.

De Fries R.S. and 14 co-authors , 1995, Mapping the land surface for global atmosphere-biosphere models: Towards continuous distributions of vegetation's functional properties, *J. Geophys. Res.* 100, 20,867-20,882.

Geron, C.D. , Pierce, T.E. , and Guenther, A.B. , 1995, Reassessment of biogenic volatile organic compound emissions in the Atlanta area, *Atmos. Environ.*, 29, No. 13, 1569-1578.

Guenther, A.B. , Zimmerman, P.R. , Harley, P.C. , Monson, R.K. , and Fall, R. , 1993, Isoprene and monoterpene rate variability: model evaluations and sensitivity analyses, *J. Geophys. Res.*, 98, No. D7, 12609-12617.

Guenther, A. , Zimmerman, P. , and Wildermuth, M. , 1994, Natural volatile organic compound emission rate estimates for U.S. woodland landscapes, *Atmos. Environ.*, 28, 1197-1210.

Guenther, A. , Hewitt, C.N. , Erickson, D. , Fall, R. , Geron, C. , Graedel, T. , Harley, P. , Klinger, L. , Lerdau, M. , McKay, W.A. , Pierce, T. , Scholes, R. , Steinbrecher, R. , Tallamraju, R. , Taylor, J. , and Zimmerman, P. , 1995, A global model of natural volatile organic compound emissions, *J. Geophys. Res.*, 100, No. D5, 8873-8892.

Guenther, A. (1997): Seasonal and spatial variations in the natural volatile organic compound emissions. *Ecological Applications* 7(1) 34-45.

Guenther, A., J. Greenberg, D. Helmig, L. Klinger, L. Vierling, P. Zimmerman, and C. Geron (1996) Leaf, branch, stand and landscape scale measurements of volatile organic compound fluxes from U.S. woodlands. *Tree Physiology*, 16, 17-24.

Guenther, A., C. Geron, T. Pierce, B. Lamb, P. Harley, and R. Fall, 1998 (in preparation) Natural emissions of volatile organic compounds, carbon monoxide, and oxides of nitrogen from North America.

Guenther, A.B. , Monson, R.K. , and Fall, R. , 1991, Isoprene and monoterpene rate variability: observations with Eucalyptus and emission rate algorithm development, *J. Geophys. Res.*, 96, No. D6, 10799- 10808.

Hewitt, C.N., R. K. Monson, and R. Fall (1990): Isoprene emission from the grass *Arundo donax* L. are not linked to photorespiration. *Plant Science*, 66, 130-144.

Hewitt, C.N. and Street, R.A. , 1992, A qualitative assessment of the emissions of non-methane hydrocarbon compounds from the biosphere to the atmosphere in the U.K.: present knowledge and uncertainties, *Atmos. Environ.*, 26A, No. 17, 3069-3077.

König, G. , Brunda, M. , Puxbaum, H. , Hewitt, C.N. , Duckham, S.C. , and Rudolph, J., 1995, Relative contribution of oxygenated hydrocarbons to the total biogenic VOC emissions of selected mid-European agricultural and natural plant species, *Atmos. Environ.*, 29, No. 8, 861-874.

Lamb, B., Guenther A., Gay, D., and Westberg, H., 1987, A national inventory of biogenic hydrocarbon emissions, *Atmos. Environ.* 21, 1695-1705.

Lamb, B., Gay, D., Westberg, H., and Pierce, T., 1993, A biogenic hydrocarbon emission inventory for the U.S.A. using a simple forest canopy model, *Atmos. Environ.* 27A, 1673-1690.

Lieth, H., and Whittaker, R.H., 1975, Primary production of the major vegetation units of the world. In: *Primary productivity of the biosphere*, Eds. Lieth, H., and Whittaker, R.H., Springer-Verlag, New York, pp. 204-215.

MacDonald R.C., and Fall. R., 1993, Detection of substantial emissions of methanol from plants to the atmosphere. *Atmos. Environ.* 27A, 1709-1713

Ortiz, A. and Dory, M.A.G. , 1990, The estimation of non methane VOC emissions from nature in Spain for CORINAIR inventory, In Pacyna, J. and Joerss, K.E. , editors, EMEP Workshop on emission inventory techniques, Regensburg, Germany, 3-6 July, 1990. Norwegian Institute for Air Research, Kjeller, Norway, EMEP/CCC-Report 7/90.

Owen, S., Boissard, C., Street, R.A., Duckham, C., Csiky, O., Hewitt, C.N., 1997, The BEMA project: screening of 18 Mediterranean plant species for volatile organic compound emissions, *Atmos Environ.*, 31, No.S1, 101-118.

Pierce, T.E. and Waldruff, P.S. , 1991, PC-BEIS: a personal computer version of the biogenic emissions inventory system, *J. Air Waste Manage. Assoc.*, 41, No. 7, 937-941.

Pierce, T.E. , 1991, User's guide to the personal computer version of the biogenic emissions inventory system (PC-BEIS), Atmospheric research and exposure assessment laboratory, U.S.E.P.A., Research Triangle Park, NC. Report EPA/600/8-90/084.

Puxbaum, H. , 1997, Biogenic emissions of alcohols, ester, ether and higher aldehydes. In: Helas, G., Slanina, J., and Steinbrecher R. (eds.) Biogenic volatile organic carbon compounds in the atmosphere, pp.79-99, SPB Academic Publishing, Amsterdam.

Puxbaum, H. , and König G., 1997, Observation of Dipropenyldisulfide and other organic sulfur compounds in the atmosphere of a beech forest with *Allium ursinum* ground cover. Atmos. Environ. 31, 291-294.

Rudolph, J., Plass-Dülmer, C., Benning, L., Brandenburger, U., Brauers, T., Dorn, H.-P., Hausmann, M., Hofzumahaus, A., Holland, F., Parusel, E., Ramacher, B., Wahner, A., Wedel, A., Duckham, C., Hewitt, N., König, G., and Puxbaum, H., 1998, The POPCORN campaign 1994, an intensive field study of biogenic and man made volatile organic compounds in the atmosphere: an overview. *J. Atmos. Chem.* in press

Ruimy, A., Saugier, B., and Dedieu, G., 1994, Methodology for the estimation of terrestrial net primary production from remotely sensed data. *J. Geophys. Res.* 99, 5263-5283.

Seufert, G., J. Bartzis, T. Bomboi, P. Ciccioli, S. Cieslik, R. Dlugi, P. Foster, N. Hewitt, J. Kesselmeier, D. Kotzias R. Lenz, F. Manes, R. Perez-Pastor, R. Steinbrecher, L. Torres, R. Valentini, and B. Versino (1997): The BEMA-project: and overview of the Castelporziano experiments. *Atmos. Environ.* 31, S1, 5-18.

Simpson, D. , Guenther, A. , Hewitt, C.N. , and Steinbrecher, R. , 1995, Biogenic emissions in Europe 1. Estimates and uncertainties, *J. Geophys. Res.*, 100, No. D11, 22875-22890.

Simpson, D., Winiwarter, W., Börjesson, G., Cinderby, S., Ferreira, A., Guenther, A., Hewitt, C.N., Janson, R., Khalil, M.A.K., Owen, S., Pierce, T.E., Puxbaum, H., Shearer, M., Steinbrecher, R., Tarrason, L., and Öquist, M.G., Inventorying emissions from Nature in Europe, submitted.

Steinbrecher, R. , 1994, Emission of VOCs from selected European ecosystems: the state-of-the-art, In Borrell , P., editor, Transport and Transformation of Pollutants in the Troposphere, Proceedings EUROTRAC symposium 1994, pages 448-455. SPB Acad. Publish. bv., the Hague, Netherlands.

Street, R.A. , 1995, Emissions of non-methane hydrocarbons from three forest ecosystems, PhD thesis, Lancaster Univ., Lancaster, England.

Street, R.A., Duckham S.,C., Boissard, and Hewitt, C.N., 1997, Emissions of VOCs from stressed and unstressed vegetation, In: Slanina, S. (ed.) Biosphere-Atmosphere Exchange of Pollutants and Trace Substances, Vol.4, Series Transport and Chemical Transformation of Pollutants in the Troposphere, Springer, Berlin.

Tanner R.L., and Zylinska B., 1994, Determination of the biogenic emission rates of species contributing to VOC in the San Joaquin Valley of California. *Atmos. Environ.* 28, 1113-1120

Tingey, D.T. , Manning, M. , Ratsch, H.C. , Burns, W.F. , Grothaus, L.C. , and Field, R.W. , 1978a, Monoterpene emission rates from slash pine, Final Report EPA CERL-045. Environmental Protection Agency, Research Triangle Park, North Carolina.

Tingey, D.T. , Ratsch, H.C. , Manning, M. , Grothaus, L.C. , Burns, W.F. , and Peterson, 1978b, Isoprene emissions rates from live oak, Final Report EPA CERL-040. Environmental Protection Agency, Research Triangle Park, North Carolina.

- UN-ECE, 1992, The environment in Europe and North America: annotated statistics 1992.
- Veldt, C. , 1988, Inventorying natural VOC emissions for the CORINAIR project, Apeldoorn, The Netherlands, MT-TNO Report 88-275; Also published in Corinair Technical Annexes Volume 2, Default emission factor handbook, European Commission EUR 12586/2, pp101-128.
- Veldt, C. , 1989, Leaf biomass data for the estimation of biogenic VOC emissions, Apeldoorn, The Netherlands, MT-TNO Report 89-306.
- Veldt, C. , 1991, The use of biogenic VOC measurements in emission inventories, Apeldoorn, The Netherlands, MT-TNO Report 91-323.
- Winer, A.M., Arey, J., Atkinson, R., Aschmann, S.M., Long, W.D., Morrison, C.L., and Olszyk, D., 1992, Emission rates of organics from vegetation in California's central valley. *Atmos. Environ.* 26A, 2647-2659
- Zimmerman, P. , 1979, Testing of hydrocarbon emissions of vegetation, leaf litter and aquatic surfaces and development of a methodology for compiling biogenic emission inventories, EPA 450/4-79-004.

18 BIBLIOGRAPHY

- Guenther, A.B. , Zimmerman, P.R. , Harley, P.C. , Monson, R.K. , and Fall, R. , 1993, Isoprene and monoterpene rate variability: model evaluations and sensitivity analyses, *J. Geophys. Res.*, 98, No. D7, 12609-12617.
- Guenther, A. , Zimmerman, P. , and Wildermuth, M. , 1994, Natural volatile organic compound emission rate estimates for U.S. woodland landscapes, *Atmos. Environ.*, 28, 1197-1210.
- Guenther, A. , Hewitt, C.N. , Erickson, D. , Fall, R. , Geron, C. , Graedel, T. , Harley, P. , Klinger, L. , Lerdau, M. , McKay, W.A. , Pierce, T. , Scholes, R. , Steinbrecher, R. , Tallamraju, R. , Taylor, J. , and Zimmerman, P. , 1995, A global model of natural volatile organic compound emissions, *J. Geophys. Res.*, 100, No. D5, 8873-8892.
- Hewitt, C. N., R.A. Street and P.A. Scholefield (1998), Isoprene and Monoterpene-Emitting Species Survey 1998: <http://www.es.lancs.ac.uk/es/people/pg/pas/download.html>
- Veldt, C. , 1989, Leaf biomass data for the estimation of biogenic VOC emissions, Apeldoorn, The Netherlands, MT-TNO Report 89-306.

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SNAP CODE:	110501
	110502
	110503
	110504
	110505
	110506
	110601

SOURCE ACTIVITY TITLE:	OTHER SOURCES AND SINKS
	<i>Wetlands (Marshes-Swamps)</i>
	<i>Undrained Marshes</i>
	<i>Drained Marshes</i>
	<i>Bogs</i>
	<i>Fens</i>
	<i>Swamps</i>
	<i>Floodplains</i>
	<i>Lakes</i>

NOSE CODE:	301.05.01
	301.05.02
	301.05.03
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1 ACTIVITIES INCLUDED

This chapter covers emissions of methane (CH₄) and to a lesser extent sulphur produced in naturally saturated soils, in areas either permanently or seasonally flooded with fresh water. Note that this chapter covers shallow lakes (110601), typically defined by depths of less than 2 m, as well as the wetland (1105) SNAP-codes. Lakes of greater than 2m depth should not generally be treated as wetlands. The chapter does not cover agricultural wetlands such as rice fields, though the biogeochemical processes are the same. (See Schütz et al., 1989, for experimental measurements from Italian rice fields.)

The main emission, CH₄, is produced by anaerobic bacteria (methanogens) in the soil, diffused through soil water and transported to the atmosphere by plants, ebullition, or diffusion. Type of vegetation soil characteristics, and local climate are three important factors affecting methane emissions; data about these factors are used to make global and regional estimates.

Natural sulfur gases such as OCS (carbonyl sulfide), DMS (dimethyl sulfide), H₂S and CS₂ are emitted from brackish wetlands and wetlands with high soil sulfur, usually as the result of

microbiological activity, though partly by chemical reduction of sulfate (some H₂S) or possibly by algae or other plants (DMS). These gases will be briefly discussed as they are not considered a significant source of pollutants. The bacteria which produce the sulfur gases usually out-compete methanogens, so methane production is inhibited by saline conditions. Brackish marshes have usually been omitted from inventories of methane emissions.

Wetland areas are affected by human management when drained for agriculture or construction; maintained for wildlife habitat or water treatment; or built/converted for water storage and transport such as canals or farm ponds. These changes in area may be estimated if adequate data are available from local sources.

2 CONTRIBUTIONS TO TOTAL EMISSIONS

Wetlands are estimated to produce about 20% of the annual global methane emissions. Recent global estimates have been 100 - 110 Tg (10¹² g) per year, with a range of about 50 - 150 Tg CH₄ emitted per year. These estimates are reviewed in Matthews (1993).

Biogenic sulfur gases emitted from wetlands and soils are estimated to be less than 2% of the total sulfur budget; 5-12 Tg S per year out of a total of 310 Tg. Less than 10% of the world's soils are in brackish marsh, so sulfur emissions from saline marshes are on the order of 1-2 Tg; insignificant compared to anthropogenic sources (Warneck, 1988; Andreae, 1984). Early studies which indicated a much larger source of biogenic sulfur gases from wetlands were either not reproduced, or may have been an artifact of the sampling process (see Chin and Davis, 1993, for further discussion).

These activities are not believed to be a significant source of PM_{2.5} (as of December 2006).

3 GENERAL

3.1 Description

CH₄ is produced by anaerobic bacteria (methanogens) in the soil, diffused through soil water and transported to the atmosphere by plants, ebullition, or diffusion. Ground water table position, type of vegetation, soil characteristics, available substrates and local climate are all important factors affecting methane emissions. Further, methanogenesis is the final step in the anaerobic degradation chain, requiring organic by-products from other bacteria as food, and emitting methane as a waste (Gujer and Zehnder, 1983). For this reason, methane emission usually requires days to weeks to become significant at the beginning of the season. Methane in turn is a food source for aerobic bacteria called methanotrophs, so it can be oxidised in the aerobic root zone of plants or aerobic layers in soil or water. Approximately 10 - 40% of the methane produced in saturated soil is eventually emitted to the atmosphere. (See Conrad, 1996, and references therein.)

Biogenic sulfur gases are formed during anaerobic decomposition, from chemical reactions with the sulfate ion, and possibly also by some species of marsh vegetation (Patrick and DeLaune, 1977; Warneck, 1988; Chin and Davis, 1993).

3.2 Definitions

Many terms are used to describe naturally occurring flooded areas: wetland, mire, bog, fen, wet tundra, swamp, wet meadow and marsh are among the most common. In common usage the terms are imprecise and sometimes interchangeable. For the purposes of this chapter:

Wetland is used as an overall term for any area of permanently or seasonally flooded soils, where soils are saturated long enough for the soil to become reduced, a methanogen population established, and methane emitted from the soil. The types of wetlands are differentiated by their vegetation, which affects the amount of organic substrate available and transport of CH₄; and by season of flooding or thawing.

The following definitions are derived from Zoltai and Pollet (1983), Aselmann and Crutzen (1989), though a similar scheme was used by Matthews and Fung (1987).

A bog is a peat forming wetland, usually with mossy vegetation, sometimes with boreal forest, waterlogged from precipitation only.

A fen is a peat forming wetland with water flowing through the system, usually with grasses and sedges in addition to moss; less acidic than a bog and more productive.

Bogs and fens make up most of the boreal wetlands in tundra ecosystems, though they may be found at other latitudes.

Swamps are forested wetlands, with much less accumulated organic matter than bogs, usually found in temperate or tropical latitudes.

A marsh is a wetland with grass, sedges or reed vegetation.

A flood plain is the area seasonally covered by water along rivers or lakes. They are significant sources of methane principally in South America and Africa.

A shallow lake is a body of water warm enough for methane to be produced in sediment and shallow enough (<2m) that methane can diffuse or bubble to the surface. Canals and farm ponds might also be considered in this category as well as natural bodies of water.

The SNAP classifications "undrained and brackish marshes", "drained marshes" are preserved for consistency with previous work, but essentially all marshes which still fit unto the definition of wetland are treated identically in the following.

3.3 Techniques

Methane fluxes from wetlands have commonly been estimated by measuring its accumulation in closed chambers. In the past few years, area estimates from various types of eddy correlation measurements have become more common. Areas of wetlands have been estimated from maps, Gore (1983) for example, and from digitized databases of soils and vegetation. Season of methane emission is usually estimated from local climate data.

3.4 Emissions

Wetlands emit methane, carbon dioxide and biogenic sulfur gases, together with minor quantities of N₂O and NO. However, methane is the only gas emitted that is globally significant. Biogenic CO₂ is simply recycled (although wetlands do play a role in the global carbon cycle as the amounts of C stored in peatlands are significant - ca 412 Gt of C world wide; Woodwell et al., 1995). Biogenic sulfur gases are insignificant compared to anthropogenic sulfur emissions.

3.5 Controls

Natural wetlands have commonly been drained in temperate and tropic zones for agriculture, construction and peat harvest. These activities have "controlled" emissions by destroying the wetlands. Arctic and high latitude boreal wetlands are not drained because the ground is frozen much of the year; no controls appear reasonable.

4 SIMPLER METHODOLOGY

Methane emission from wetlands (W_{CH_4} , in mass units) is estimated by:

$$W_{CH_4} = \sum_i^7 (A_i \cdot F_i \cdot S_i \cdot cf) \quad (01)$$

Where $i = 1, 2, \dots, 7$ for the 7 wetland types; A_i is the area in each wetland type; F_i is the seasonal average flux (in mass/area/time units, usually $mg\ CH_4\ m^{-2}\ day^{-1}$); S_i is the length of the season of methane emission. The season is the time the soil is thawed for boreal and northern temperate wetlands, and the length of time the soil is inundated for flood plains and seasonal marshes and swamps. "cf" is the appropriate units conversion factor.

5 DETAILED STATE OF THE ART METHODOLOGY

The detailed methodology is essentially the same as the simple methodology. The estimates may be improved by introducing wetland types characterized specifically by country; or using local flux measurements rather than the averaged ones given in section 8. Any information specific to a country rather than a global database should improve the precision of the estimate.

6 RELEVANT ACTIVITY STATISTICS

Wetland area data are found in a series of tables in Aselmann and Crutzen, 1989. They show percent wetland area in 2.5° latitude x 5° longitude cells. Matthews and Fung (1987) used a different classification scheme and divided their estimate into 1°x 1° cells. Their data base is documented by Matthews (1989) and is available by FTP from the US NCAR (National Center for Atmospheric Research) data site: ncardata.ucar.edu.

Maps of some wetland areas in Europe may be found in Gore (1983) volume 4A: General Studies and volume 4B: Regional Studies. Great Britain, Ireland, Finland and Sweden are covered in particular detail. Most of these maps are based on research done in the country of origin.

Local government agencies and researchers may be able to provide rainfall and temperature data to determine seasonality; and more precise land use data for wetland areas.

7 POINT SOURCE CRITERIA

All wetland sources are considered area sources.

8 EMISSION FACTORS, QUALITY CODES AND REFERENCES

Methane:

Bartlett and Harriss (1993) did a thorough review of flux measurements from wetlands and shallow lakes for the purpose of making global estimates. The following table is adapted from their work. They combined measurements from fens and bogs.

Climate Zone	Flux by Wetlands Type ($\text{mgm}^{-2}\text{d}^{-1}$)					
	Bogs	Fens	Marsh	Swamp	Flood-plain	Shallow Lakes
Arctic	96	96				
Boreal	87	87	87	87		35
Temperate	135	135	70	75	48	60
Tropical	199	199	233	165	182	148

The climate zones are arctic: 60 - 90° latitude; boreal: 45 - 60° latitude; temperate: 20 - 45° latitude; tropical: 0 - 20° latitude. These climate zones apply best to the American continents, as most of the northern hemisphere studies are from Canada and the U.S., and most southern hemisphere studies were done in Brazil.

Biogenic Sulfur gases:

Stuedler and Peterson estimated a total annual emission of $5.8 \text{ g S m}^{-2} \text{ yr}^{-1}$ in a study which measured all principal biogenic sulfur gases emitted from a brackish marsh over the period of a year.

9 SPECIES PROFILES

10 UNCERTAINTY ESTIMATES

The data quality for making an estimate of methane emissions from wetlands is moderate (D rating).

Wetland flux estimates are probably the greatest source of uncertainty in making global estimates of methane. Although there are measurements in all wetland types from the principal wetland areas, fluxes may vary over several orders of magnitude at a single site. Inter-annual variation of seasonal averages can vary as much as an order of magnitude. Most boreal and temperate zone flux measurements have been made in North America and Scandinavia, and most tropical zone measurements have been made in Central and South America. Since there are few or no other measurements of methane flux from other parts of the world, the uncertainty of using the available measurements cannot be calculated, but may be large. Measurements of methane flux in Europe have, however, fit in the range of other boreal and high temperate zone measurements.

The estimated areas of wetlands may differ greatly depending on the underlying vegetation databases. The differences in area estimates between Matthews and Fung (1987) and Aselmann and Crutzen (1989) are discussed at length in the latter paper and in Bartlett and Harris (1993). Their total areas are very close but their distribution differs greatly, particularly in the tropics. Their estimates of total area for the northern hemisphere temperate and boreal zones are very close, but their vegetation classes are not strictly comparable.

The flux estimates for biogenic sulfur gases is poorer (an E rating). There are few measurements of all sulfur gases and the measured emissions are extremely variable.

The comments on the uncertainties of flux measurements of methane also apply to the biogenic sulfur gases. Additional variability is due to flux which varies with the tide (H_2S), or with daylight (DMS). Since not all researchers have measured all gases, it is difficult to get a total sulfur estimate. Since there is still possible contamination of the samples during measurement for the earlier data, there can be four orders of magnitude difference between measurements made in the same area by different researchers.

11 WEAKEST ASPECTS/PRIORITY AREAS FOR IMPROVEMENT IN CURRENT METHODOLOGY

As noted in 10, the emissions flux estimates are probably the greatest source of uncertainty. Additionally, linking flux estimates to wetland classification is an important problem. As it is not known exactly which parameters affect flux, then it is difficult to devise good parameterisation schemes. A further problem arises from differences in techniques used in measuring fluxes - such factors may explain some of the variability found of measurements.

Development of better techniques for remote sensing and evaluation is probably an essential component of inventory improvement.

12 SPATIAL DISAGGREGATION CRITERIA

Methane emissions are estimated by the different types of wetlands defined in section 3.2.

13 TEMPORAL DISAGGREGATION CRITERIA

Methane emissions vary seasonally, usually following soil temperature, plant growing season or saturation season, though exceptions may be found (Svensson and Rosswall, 1984; Whalen and Reeburgh, 1992; Westermann, 1993). For example, in the high northern latitudes wetlands are usually classified as bogs, forested bogs, and fens with maximum emissions from June to September. Methane emission increases when soil temperature increases above 0 degrees but has been measured at very low levels from frozen soil. Seasonal wetlands such as flood plains will only emit methane during the wet season, and methane emissions vary within wetlands along moisture gradients (Svensson, 1976; Moore et al., 1990; Granberg et al., 1997). Dry, aerated soils are usually sinks of methane; drought or other change in water table may cause a source area to become a sink (Harriss et al., 1982, Whalen et al., 1991, Oechel, 1993).

All fluxes given in section 8 are averaged diurnally and seasonally.

14 ADDITIONAL COMMENTS:

Very recently, Cao et al. (1996) and Christensen et al. (1996) have modeled the carbon system and methane emissions from wetlands. This type of model is considerably more complicated but allows modeling changes of methane emissions due to changes in climate. At present these models are validated against global estimates using measured fluxes (Matthews and Fung, 1987; Aselmann and Crutzen, 1989; Bartlett and Harris, 1993). The models are not yet generally available.

15 SUPPLEMENTARY DOCUMENTS

16 VERIFICATION PROCEDURES

17 REFERENCES

Andreae, M.O. The emission of sulfur to the remote atmosphere: Background paper. In: *The Biogeochemical Cycling of Sulfur and Nitrogen in the Remote Atmosphere*. J.N. Galloway, R.J. Charlson, M.O. Andreae, and H. Rodhe, Eds. Nato ASI Series C Vol. 159, D. Reidel, Holland, 1984.

Aselmann, I. and P.J. Crutzen. Global distribution of natural freshwater wetlands and rice paddies, their net primary productivity, seasonality, and possible methane emissions. *Journal of Atmospheric Chemistry* 8:307-358, 1989.

Bartlett, K.B. and R.C. Harriss. Review and assessment of methane emissions from wetlands. *Chemosphere* 26:261-320, 1993.

Cao, M., S. Marshall, and K. Gregson. Global carbon exchange and methane emissions from natural wetlands: Application of a process based model. *J. of Geophys. Res.* 101:14,399-14,414, 1996.

Christensen, T.R., I.C. Prentice, J. Kaplan, A. Haxeltine and S. Stich. Methane flux from northern wetlands and tundra, an ecosystem modeling approach. *Tellus* 48B:652-661, 1996.

Chin, M. and D.D. Davis. Global sources and sinks of OCS and CS₂ and their distributions. *Global Biogeochemical Cycles* 7:321-337, 1993.

Conrad, R. Soil microorganisms as controllers of atmospheric trace gases (H₂, CO, CH₄, OCS, N₂O, and NO). *Microbiological Reviews* 60:609-640, 1996.

Gore, A.J.P. Introduction. In: *Ecosystems of the World 4A, Mires: Swamp, Bog, Fen, and Moor, General Studies*. A.J.P. Gore, Ed. Elsevier Scientific Publ. Co., 1983.

Granberg, G., C. Mikkilä, I. Sundh, B.H. Svensson, and M. Nilsson. Sources of spatial variation in methane emission from mires in northern Sweden: A mechanistic approach in statistical modeling. *Global Biogeochemical Cycles* 11:135-150, 1997.

Matthews, E. and I. Fung. Methane emission from natural wetlands: Global distribution, area, and environmental characteristics of sources. *Global Biogeochemical Cycles* 1:61,86, 1987.

Matthews, E. Global Databases on distribution, characteristics and methane emission of natural wetlands: documentation of archived data tape. NASA Technical Memorandum 4153, 1989.

Matthews, E. Wetlands. In: *Atmospheric Methane: Sources, Sinks, and Role in Global Change*. M.A.K. Khalil, Ed. NATO ASI Series: Global Environmental Change, Vol. 13, Springer-Verlag, Berlin, 1993.

Moore, T., N. Roulet, and R. Knowles. Spatial and temporal variations of methane flux from subarctic/northern boreal fens. *Global Biogeochemical Cycles* 4:29-46, 1990.

Oechel, W. C., et al. (1993). "Recent change of Arctic tundra ecosystems from a net carbon dioxide sink to a source." *Nature* **361**: 520-523.

Patrick, W.H. Jr. and R.D. DeLaune. Chemical and biological redox systems affecting nutrient availability in the coastal wetlands. *Geoscience and Man* 18:131-137, 1977.

Schütz, H., A. Holzapfel-Pschorn, R. Conrad, H. Rennenberg, and W. Seiler. A 3-year continuous record on the influence of daytime, season, and fertilizer treatment on methane emission rates from an Italian rice paddy. *J. Geophys. Res.* 94(D13):16,405-16,416, 1989.

Stuedler, P.A. and B.J. Peterson. Annual cycle of gaseous sulfur emissions from a New England *Spartina alterniflora* marsh. *Atmospheric Environment* 9:1411-1416, 1985.

Svensson, B.H. Methane production in tundra peat. *Microbial Production and Utilization of Gases (H₂, CH₄, CO)*. H.G. Schlegel, G. Gottschalk, N. Pfennig and E. Goltze, editors (Gottingen). p. 135-139, 1976.

Warneck, P. *Chemistry of the Natural Atmosphere*. International Geophysics Series, Vol. 41, Academic Press Inc., USA. p.498-505; 540-542, 1988.

Whalen, S.C., W.S. Reeburgh, and K.S. Kizer. Methane consumption and emission by taiga. *Global Biogeochemical Cycles* 5:261-273, 1991.

Zoltai, C.S. and Pollett, F.C. 1983. Wetlands in Canada. In: *Ecosystems of the World. Vol 4B, Mires: Swamp, Bog, Fen, and Moor*. A.J.P. Gore (ed). Elsevier Sci. Publ., New York, NY, pp. 245-268.

18 BIBLIOGRAPHY

Bartlett, K.B., R.C. Harriss, and D.I. Sebacher. Methane flux from coastal salt marshes. *J. Geophys. Research* 90:5710-5720, 1985

Botch, M.S. and V.V. Masing. Mire ecosystems in the U.S.S.R. In: *Ecosystems of the World 4B, Mires: Swamp, Bog, Fen, and Moor, Regional Studies*. A.J.P. Gore, Ed. Elsevier Scientific Publ. Co., Amsterdam, The Netherlands. 1983.

Gujer, W. and Zehnder, A.J.B. 1983. Conversion processes in anaerobic digestion. *Water Sci. Technol.* 15: 127-167.

Clymo, R.S. and E.F.J Reddaway. Productivity of *Sphagnum* (Bog-moss) and peat accumulation. *Hidrobiologia* 12:181-192, 1971.

Freeman, C., M.A. Lock and B. Reynolds. Fluxes of CO₂, CH₄, and N₂O from a Welsh peatland following simulation of water table draw-down: potential feedback to climate change. *Biogeochemistry* 14:51-60, 1993.

Gallagher, M.W., T.W. Choularton, K.N. Bower, I.M. Stromberg, K.M. Beswick, D. Fowler, K.J. Hargreaves. Measurement of methane fluxes on the landscape scale from a wetland area in north Scotland. *Atmospheric Environment* 28:2421-2430, 1994.

Holzapfel-Pschorn, A., R. Conrad, W. Seiler. Production, oxidation and emission of methane in rice paddies. *FEMS Micro. Ecol.* 31:343-351, 1985.

Jørgensen, B.B. and B. Okholm-Hansen. Emissions of sulfur gases from a Danish estuary. *Atmospheric Environment* 19:1737-1749, 1985.

Patrick, W.H. Jr. and R.D. DeLaune. Chemical and biological redox systems affecting nutrient availability in the coastal wetlands. *Geoscience and Man* 18:131-137, 1977.

Ruuhijärvi, R. The Finnish mire types and their regional distribution. In: *Ecosystems of the World 4B, Mires: Swamp, Bog, Fen, and Moor, Regional Studies*. A.J.P. Gore, Ed. Elsevier Scientific Publ. Co., Amsterdam, The Netherlands. 1983.

Sebacher, D.I., R.C. Harriss, and K.B. Bartlett. Methane emissions to the atmosphere through aquatic plants. *J. Environ. Qual.* 14:40-46, 1985.

Sundh, I., M. Nilsson, G. Granberg, and B.H. Svensson. Depth distribution of microbial production and oxidation of methane in northern boreal peatlands. *Microbial Ecology* 27:253-265, 1994.

Svensson, B.H. Methane production in tundra peat. *Microbial Production and Utilization of Gases (H₂, CH₄, CO)*. H.G. Schlegel, G. Gottschalk, N. Pfennig and E. Goltze, editors (Gottingen). p. 135-139, 1976.

Svensson, B.H. and T. Rosswall. In situ methane production from acid peat in plant communities with different moisture regimes in a subarctic mire. *Oikos* 43:341-350, 1984.

Westermann, P. and B.K. Ahring. Dynamics of methane production, sulfate reduction, and denitrification in a permanently waterlogged alder swamp. *Appl. Environ. Microbiol.* 53:2554-2559.

Westermann, P. Temperature regulation of methanogenesis in wetlands. *Chemosphere* 26:321-328, 1993.

Whalen, S.C. and W.S. Reeburgh. Interannual variations in tundra methane emission: A 4-year time series at fixed sites. *Global Biogeochemical Cycles* 6:139-159, 1992.

Woodwell, G.M., MacKenzie, F.T., Houghton, R.A., Apps, M.J., Gorham, E., Davidson, E.A. 1995. Will the warming speed the warming? In: *Biotic Feedbacks in the Global Climatic System*

Woodwell, G.M. and MacKenzie, F.T. (eds.) Oxford University Press, Oxford, UK, pp 393-411.

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SNAP CODE: 110600

SOURCE ACTIVITY TITLE: OTHER SOURCES AND SINKS
Waters

NOSE CODE:

NFR CODE: N/A

A specific methodology for these activities has not been prepared because the contribution to total national emissions is thought to be currently insignificant, i.e. less than 1% of national emissions of any pollutant.

If you have information contrary to this please contact the expert panel leaders.

This activity is not believed to be a significant source of PM_{2.5} (as of December 2006).

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SNAP CODE: 110701
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110703

SOURCE ACTIVITY TITLE: OTHER SOURCES AND SINKS: ANIMALS
Termites
Mammals
Other Animals

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1 ACTIVITIES INCLUDED

This section covers the emissions from wild-living animals. Both the emissions from the intestines and from excreta are included. Not covered are emissions from animal husbandry (chapter 10.4: Enteric fermentation) or from pets, which are partly similar, but may be considered influenced by human behavior in many respects. Still included here however are emissions from humans (breath, sweat, etc.; excreta are dealt with in chapter 9.1.7, latrines, or 9.10.2, waste water treatment), as they do not appear anywhere else and should be perceived differently to other anthropogenic emissions.

2 CONTRIBUTIONS TO TOTAL EMISSIONS

The information available is very sparse. With respect to the global situation, animal methane emissions have been attributed to termites, which are hardly relevant for Europe. The relatively high emissions of ammonia given for humans in some publications include emissions from pets, and thus need to be considered with caution for the purpose of this chapter. Nevertheless the figures presented may give some guidance which levels of emissions are to be expected.

For the UK [1], ammonia emissions from humans (without pets) have been estimated at 0.7 % of total ammonia emissions, and wild animals (deer and birds) at 0.2 %. Global emissions of ammonia were estimated at 4.8 % for humans, and at 0.2 % for wild animals [2]. The estimate for humans here however includes emissions from pets (which in [1] are estimated to total three times the amount of human emissions) and from latrines. Estimates for methane are not available for Europe, but using global estimates [3] or the emission factors provided below the contribution of emissions appears to be smaller than 1% of the total.

These activities are not believed to be a significant source of PM_{2.5} (as of December 2006).

3 GENERAL

3.1 Description

Metabolic processes especially in the intestines of animals, but also processes in their excretions are responsible for gas formation. One important pathway leading primarily to methane formation is the anaerobic degradation of plant cellulose by symbiotic microflora (methanogenic bacteria, but also acetogenic bacteria) in the intestines. Major kinds of animals that are known to emit methane are mammals (primarily ruminants and rodents) and termites. A completely different pathway of emissions is the decay of urea or uric acid to ammonia in animal manure (mammals or birds). This pathway may also lead to N₂O formation. Emissions however are much more pronounced for domestic animals, where manure is actually collected and kept liquid for longer periods of time, or other sites where animals live in a very dense population (point emissions from bird breeding colonies on small islands, e.g. in the North Sea). Other relevant emissions are volatile organic compounds like isoprene, however no specific information could be obtained as this source is probably negligible.

It is very important to discuss the difference and the reasons of the difference between domestic and wild animals. Domestic animals are generally kept more densely, such that manure management is needed and the manure has to be stored for a longer period of time. Chemical processes in the manure (decay of urea to ammonia) are completely different and much less relevant for natural animals. Also, the diet is quite different between natural and domestic animals, influencing the feed nitrogen content, which is important for ammonia formation. The diet also affects the methane yield, the proportion of food energy content emitted in the form of methane. Nevertheless emissions need to be considered comparable to some extent, especially due to the absence of any better data (see section 8).

For a gas which deposits quite efficiently as ammonia, also a canopy effect may be taken into account. Gases released effectively from the animals may well be absorbed immediately in the forest canopy or in the grass before ever actually escaping to the lowest layers of the atmosphere. These emissions will never have any apparent effect on the atmosphere.

3.2 Definitions

Wild-living animals: Animals which are not severely affected in their feeding behavior or their mobility by anthropogenic influences, and are not controlled by humans.

3.3 Controls

Not applicable.

3.4 Emissions

Emissions are mainly methane and ammonia. Some NMVOC emissions are also possible, but probably small. Considering similar processes as for domestic animals, also nitrous acid emissions should be expected. For instance, formic acid emissions have been attributed to formicine ants [4]. These emissions have never been actually quantified and may not be relevant anywhere outside the tropical rain forests.

3.5 Controls

There is no controls to natural emissions by definition.

4 SIMPLER METHODOLOGY

Apply emission factors given in section 8.

5 DETAILED METHODOLOGY

For detailed emission assessment, emission factors as given in section 8 should be adapted towards national particularities. Such an approach has been used in [5]. Animal weights may vary within a species as much as a factor of 2, leading to considerably different emission scaling factors, depending on which variety of a species is dominant in a certain country. Also, the feeding habits should be taken into account, both in terms of energy content in order to assess methane emissions [3], and in nitrogen content for scaling ammonia emissions [2].

6 RELEVANT ACTIVITY STATISTICS

Information from wildlife specialists, hunting statistics etc. on number and kind of animals present. For big game species, hunting accounts for about 20-30% of the winter population (which resembles the annual population minimum).

7 POINT SOURCE CRITERIA

There are no point sources.

8 EMISSION FACTORS, QUALITY CODES AND REFERENCES

As measuring emission factors for wild living animals is almost by definition very difficult, the data quality is poor (D-E). Most information is taken from similarities and analogies between domestic and wild animals. The choice of emission factors for ammonia has been discussed in detail [1]. Ammonia emission rates have been given for red deer (0.9 kg/individual and year, [1]) and for reindeer (1 kg N per individual and year, [2]). The emission factors seem to be similar enough to be combined for Table 1. Not considered here however was possible redeposition of ammonia in forests at plant surfaces before emissions actually can reach the atmosphere (canopy effect), as discussed in [2].

For methane, data presented in this guidebook for enteric fermentation were used [6]. Large uncertainty is associated with deriving deer emissions from cattle emission factors. Scaling of these emissions for moose and for red deer was performed using estimated excretion of nitrogen [2] as an indicator of their metabolic activity. These emission factors are about 50 % larger than those suggested previously [3]. However as methane emissions from animal droppings are not included in either of the data given (an additional 25 % according to [6]), the emission factors proposed here still should not be considered upper limits. Methane

emissions from humans, mainly in human breath, have been assessed from measured values [3]. The resulting emission factor of 0.07 kg/person and year is notably lower to that of pigs, which may have a comparable metabolism. Considering the food uptake of humans, which is about one third of that of pigs, an emission factor of 0.5 kg/person would be expected. Much of this discrepancy may be due to a different diet but no full explanation is possible. We thus propose to apply an emission factor of 0.1 kg/person and year.

As weights for different game species vary considerably, we recommend to further scale the emissions by the life weight in a linear fashion. A more complex scaling proportional to the $\frac{3}{4}$ power of weight has been suggested [3], which may describe the food demand more closely, but other parameters also contribute to methane emissions such that it does not seem justified to perform an increase in complexity. The average weights of species have been simplified from much more detailed literature data [7]. Thus the average weight of red deer and reindeer is taken at 100 kg, fallow deer and white-tailed deer 90 kg, roe deer 15 kg, chamois 35 kg, ibex 70 kg and mouflon 25 kg. Moose emissions were assumed to be twice those of reindeer, according to estimates of nitrogen excretion [2]. The resulting methane emission factors are consistent with estimations by the Swiss Federal Office of Environment [8]. Ammonia emission factors agree in part with data from the Czech Republic [5]. There are discrepancies of almost a factor of 3 for red deer however, as the dominant variety is the unusually heavy Carpathian deer (170 kg).

Table 8.1: Emission factors for wild animals' emissions (in kg per animal/person and year)

	Assumed life weight [kg]	CH ₄	NH ₃	Literature
deer (red deer, reindeer)	100	25	1.1	derived from [6],[1]
moose	350	50	2.2	derived according to [2]
Roe deer	15	4	0.2	scaled from red deer*
boar		1.5	1	[6], derived from [1]
birds	0.8	--	0.12	[1]
Large birds	2.4	--	0.36	[1]
humans		0.1	0.05	derived from [3], [1]

* Use animal weights to similarly scale emissions for other species

No information at all was available for rodents. Here also linear scaling by weight should be performed. While this probably underestimates the metabolic activity of small animals somewhat, the methane yield, given in [3] as the fraction of food energy content that is emitted as methane, has been assumed to be clearly smaller for any species other than ruminants. A Czech study [5], taking into account the nitrogen content of feed, assumes ammonia emissions from hares to be about 8 times of what should be expected from weight scaling. On the other hand, for smaller animals living close to or under the ground, the canopy effect should be expected very large. All of this is to be considered part of overall uncertainty. Not included were termite emissions, which are currently assumed negligible for the European continent, even if termites have become established in Southern Europe, or emissions from other invertebrates.

9 SPECIES PROFILES

No profiles are needed for methane or ammonia emissions. Information on NMVOC is missing.

10 UNCERTAINTY ESTIMATES

Uncertainty is to be considered very high (data quality D, for methane emissions from deer E).

11 WEAKEST ASPECTS/PRIORITY AREAS FOR IMPROVEMENT IN CURRENT METHODOLOGY

Emission rates are primarily inferred from domestic animals.

12 SPATIAL DISSAGGREGATION CRITERIA FOR AREA SOURCES

Forest area or grassland area, depending on animal species considered.

13 TEMPORAL DISSAGGREGATION CRITERIA

Source is too small such that no detailed temporal disaggregation is needed.

14 ADDITIONAL COMMENTS

Wild living animals are generally to be considered as causing natural emissions, even if their number is to a large extent dependent of human interest (in both directions: animals in competition to domestic animals, but also animal feeding in winter because of hunting interests). The reason is that anthropogenic influence should not be considered overwhelming in this respect.

More problematic is the question of human emissions. The human metabolism clearly is associated with anthropogenic activities, and the number of humans on earth (or in Europe) clearly is out of its natural boundary. Nevertheless it seems ethically not correct to submit this type of emissions to those which are effectively controllable by man. Human control in that respect, i.e. regulation of the number of people on earth for the sake of limiting emissions to the atmosphere, can not be acceptable. Therefore also these emissions should be considered "natural".

15 SUPPLEMENTARY DOCUMENTS

16 VERIFICATION PROCEDURES**17 REFERENCES**

- [1] Sutton M.A., Place C.J., Eager M., Fowler D., Smith R.I. (1995). Atmos.Environ. 29, 1393-1411.
- [2] Bouwman A.F., Lee D.S., Asman W.A.H., Dentener F., Van Der Hoek K.W., Olivier J.G.J. (1997). A Global High Resolution Emission Inventory for Ammonia. Global Biogeochemical Cycles 11, 561.
- [3] Crutzen P.J., Aselmann I., Seiler W. (1986). Tellus 38B, 271-284.
- [4] Graedel T.E., Eisner T. (1988). Tellus 40B, 335-339.
- [5] Jelinek A. (1997). In: Emission Inventories of Air Pollutants Project, Final Report, Annex 2. Czech Hydrometeorological Institute, Prague.
- [5] Chapter 10.4 (Enteric Fermentation), V2.0, this handbook
- [6] Niethammer J., Krapp F., Eds. (1986). Handbuch der Säugetiere Europas, Band 2/II, Paarhufer - Artiodactyla (Suidae, Cervidae, Bovidae). Aula, Wiesbaden.
- [7] Bundesamt für Umweltschutz, Emissions of air polluting substances from natural sources in Switzerland (in German). Schriftenreihe Umweltschutz 75, Berne (CH), November 1987.

18 BIBLIOGRAPHY**19 RELEASE VERSION, DATE AND SOURCE**

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SOURCE ACTIVITY TITLE: OTHER SOURCES AND SINKS
Volcanoes

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1 ACTIVITIES INCLUDED

The current chapter includes emissions from geothermal activities, both eruptive and non-eruptive. Sources include volcanoes, but also fumaroles, geysers, metamorphic degassing or other activities related to molten magma in the earth's crust. Heated magma under pressure contains gases like sulfur dioxide, carbon dioxide, hydrogen sulfide, mercury and chlorine. These gases may be released when magma gets close to the surface and the pressure may be discharged.

With respect to the different sources, non-eruptive volcanoes that outgas at relatively constant rates seem to be more important than those from sporadic eruptions, both for CO₂ [1] and SO₂ [2]. However the sporadic emissions are much more difficult to assess

Some of the emissions may also be considered anthropogenic, when produced at geothermal power plants where artificial holes are drilled to obtain hot water from the earth's interior. These emissions however are treated in SNAP 0507 and are assumed to be rather small.

2 CONTRIBUTIONS TO TOTAL EMISSIONS

The emissions from volcanoes show great regional and temporal variation. Most affected are volcanic areas, and also volcanic activity tends to be highly variable. The number of active subaerial volcanoes per year based on a 5-year running average is approximately 60 [2, 3]

In Europe, significant volcanic emissions are currently limited to Italy and to Iceland. For Italy, the SO₂ emissions from Mt. Etna have been estimate to amount to 1.5+/-0.3 Mt per year [4], while globally for all non-eruptive volcanoes 9 Mt have been reported [2, 5]. On a global scale and including the highly variable annual contribution of eruptive volcanism of about additional 4 Mt per year, SO₂ from volcanoes is estimated to account for about 5-10 % of the anthropogenic flux [5] (in Japan about 50 % of the total [6]). H₂S Emissions are considered to be quickly oxidized to SO₂ in the atmosphere [7] and have been assumed negligible by some authors [7, 8]. In contrast, a very recent compilation [2] estimates the global emissions of S from H₂S and other species at about 3.5 Mt/yr additionally. Then the volcanic sulfur flux is about 13 % of the anthropogenic flux.

For CO₂, emissions from subaerial volcanoes are considered greater than those from the submarine ones (mostly mid-oceanic ridge system) and are in the range of

0.01-0.05 x 10¹² mol/yr (0.44-2.2 Mt/yr) for one major volcano [9] and are globally at 65 Mt/yr [10], clearly two orders of magnitude lower than the anthropogenic output of CO₂ [1]. Values for the Etna plume have been measured at about 13 Mt/yr, with a similar amount of diffusive emissions [4]. Possibly due to the low solubility of CO₂ in silicate melts at upper crustal depths, the annual quiescent release of CO₂ from all active volcanoes seems to be more than an order of magnitude greater than that annually emitted directly from all forms of erupting lava.

Considerable emissions of aerosols are present in most volcanic plumes [11]. Aerosol emissions are however not subject of the current guidebook. Emissions of Hg and Cl₂ or F₂ have been measured occasionally, but are very difficult to generalize. [7, 12]

This activity is not believed to be a significant source of PM_{2.5} (as of December 2006).

3 GENERAL

3.1 Description

Heated rocks in the earth's crust may be chemically transformed such that gases are released. Carbonates may thus release CO₂, and Sulfates SO₂. These gases may be dissolved at a high pressure in the molten magma. Reaching the surface (either at the sea floor for submarine volcanoes, or at the atmosphere) the pressure decreases and the gases are emitted into the atmosphere.

3.2 Definitions

Volcano: Site where molten magma / lava occasionally reaches the surface

Non-arc volcano: Volcano on a hot spot or rift zone - erupts more frequently, total number is smaller

Arc volcano: Volcano at a subduction zone - eruptions are more violent

Fumarole: Gas vent caused by leaks from magma underneath

Geysir: Water fountain driven by venting gas due to hot magma

3.3 Techniques

A differentiation of techniques is not applicable to natural emission sources. However different source categories exist. Volcanoes are sources that have magma outflow. By contrast, fumaroles and other sources only vent gases through cracks in the rocks.

There are significantly different emission patterns also among volcanoes. Outgassing may occur continuously (globally the larger portion of emissions), or are episodic in the course of an eruption. Differentiation can also be made among eruptive emissions: Eruptions in an arc tectonic regime tend to be more violent, but seem to have a more predictable pattern of explosivity strength vs. SO₂ emissions.

The different types of volcanoes are well known and data are available. Generally, continuous flow volcanoes have a low viscosity magma and also for that reason have flat slopes, while eruptive volcanoes are comparatively steep.

3.4 Emissions

Volcanic activities release gases from the minerals being heated to form magma. Most important emissions are SO₂ and CO₂, but also H₂S. Trace constituents include Hg (mostly as sulfur complexes, Cl₂ and F₂).

3.5 Controls

There is no controls to natural emissions by definition.

4 SIMPLER METHODOLOGY

Primary source of geothermal emissions are active volcanoes. These volcanoes are well known and geologically described. Emissions from explosive volcanism can be assessed based on the Volcanic Explosivity Index (VEI) of volcanoes. The Smithsonian Global Volcanism Network catalogues each eruption during the past 200 years. Differentiation is to be made between arc-volcanoes and non-arc volcanoes. CO₂ emissions may be derived from SO₂ emissions, considering the additional uncertainties Emissions from many continuous emitting volcanoes have been listed [2], other volcanoes should be scaled to one of those listed.

The secondary sources (fumaroles, geysers) are hardly ever significant sources. Emissions should be estimated from approximations of the number of sources, the volume gas flow and the concentrations.

5 DETAILED METHODOLOGY

Emissions from specific volcanoes can be assessed using spectrometric data [13] from ground assessments also in combination with available satellite data [14]. Evaluations may take advantage of the existing dataset of the Total Ozone Mapping Spectrometer (TOMS) aboard NASA satellite Nimbus 7, which allows evaluation of SO₂ emissions [5], or the SBUV/2 instrument carried by NOAA-11 [15].

6 RELEVANT ACTIVITY STATISTICS

There are no statistical data available. Instead, geological information needs to be obtained from the respective national geological survey.

Satellite data can in principle be obtained from NASA or NOAA, respectively. The exact procedures however have not been checked.

7 POINT SOURCE CRITERIA

Each active volcano is to be considered a point source.

8 EMISSION FACTORS, QUALITY CODES AND REFERENCES

SO₂:

Explosive emissions:

Arc volcanoes: $\log E = -0.25 + 0.76 \text{ VEI}$ [5]

where E is emission of SO₂ (kt)

VEI is the volcanic explosivity index

Non arc-volcanoes: emitted SO₂ is typically much higher and less dependent on the VEI. An order of magnitude higher emissions should be assumed for eruptions of non-arc volcanoes, using the same formula as for arc volcanoes. The uncertainty is very high however. Global emissions are considered to be around 4 Mt/yr [5].

Non-explosive emissions:

9 Mt/yr globally for non-explosive volcanoes (Etna-type), which emit at steady state. More specific information has been compiled [2]. Data given specifically therein should be applied.

CO₂-emissions are in the range of $0.01-0.05 \times 10^{12}$ mol/yr (0.44-2.2 Mt/yr) for one major volcano [9], but occasionally much higher (Mt. Etna: plume emissions and diffusive emissions combined 25 Mt/yr). The ratio of CO₂/SO₂ is considered to be around 1.5 for arc emissions, but globally up to 4 or 5 on the molar scale [9], part of the difference caused by the unusually high CO₂ emissions from Mt. Etna. These figures may be taken if no other information is available.

F and Cl emission data are available for Mt. Erebus, Antarctica, which has a very uncommon alkaline magma, rich in halogens and various trace metals. These data therefore need to be seen as an indication of an upper boundary rather than as an emission factor as such. The average F/S ratio (by weight) in Erebus gas reported is 0.69, for Cl/S it is 0.55 [7]. Similarly Hekla (Iceland) is renowned for its high concentration of F and Cl during eruption. However these results should only be taken if specific information is available, as they are known to be on the upper end.

Emissions of 4-20 mg Hg / kg fumarole vapors have been reported and may be applied [11]. According to [2], the ratio of SO₂-S to S in other sulfur species is about 2:1, with 71 % of the sulfur contained in H₂S. The mass ratio of H₂S/SO₂ is 0.21 and may be applied for estimating H₂S emissions.

CO₂ Emissions from geothermal fields have been reported in [16]. Strictly these emissions are anthropogenic and should be reported in connection with power generation (SNAP 1)

9 SPECIES PROFILES

Table 9.1: Profiles for sulfur compounds in % S (from [2])

SO ₂	63
H ₂ S	24.5
CS ₂	2.4
OCS	1.5
SO ₄ ²⁻	1.4
particulate S	0.8
other:	6.6

10 UNCERTAINTY ESTIMATES

The measured variability of diffusive emission fluxes may be in the order of 20 % (relative standard deviation [4]). The uncertainty with the emission factors however is assumed to be in the range of one order of magnitude.

11 WEAKEST ASPECTS/PRIORITY AREAS FOR IMPROVEMENT IN CURRENT METHODOLOGY

Surrogate parameters to establish emission factors are rather weak. Other surrogates than the Volcanic eruptivity index, which are linked more closely to the emissions, need to be identified. Validation of TOMS data for assessing eruptive emissions is needed.

12 SPATIAL DISSAGGREGATION CRITERIA FOR AREA SOURCES

Emission areas should be limited to geologically active areas, like calderas.

13 TEMPORAL DISSAGGREGATION CRITERIA

No generalization possible; temporal disaggregation may be performed for past periods according to available records of volcanic activity.

14 ADDITIONAL COMMENTS

Volcanic emissions are the typical example of emissions not at all influenced by man. Nevertheless there may be cases where this is not quite true, especially in connection with geothermal power use or other deep drills. These emissions however are to be taken into account elsewhere.

15 SUPPLEMENTARY DOCUMENTS

16 VERIFICATION PROCEDURES

Emissions from volcanic sources may be estimated from spectroscopic measurements (correlation spectrometer, COSPEC: [14]; LIDAR [12]).

17 REFERENCES

- [1] Gerlach-Terrence-M. (1990) Natural sources of greenhouse gases: carbon dioxide emissions from volcanoes. Transactions - 1990 International Symposium on Geothermal Energy, Kailua-Kona, HI, USA - Geothermal Resources Council v14 pt 1. Publ by Geothermal Resources Council, Davis, CA, USA. p639-641
- [2] Andres R.J. and Kasgnoc A.D. (1997). A Time-averaged Inventory of Subaerial Volcanic Sulfur Emissions; submitted to J.Geophys.Res. <http://blueskies.sprl.umich.edu/geia/emits/volcano.html>.
- [3] Simkin T. Siebert L. (1984). Explosive eruptions in space and time: Durations, intervals, and a comparison of the world's active volcanic belts. In: Explosive Volcanism: Inception, Evaluation, and Hazards, pp. 110-121, National Academy Press, Washington, D.C.
- [4] Allard P., Carbonnelle J., Dajlevic D., LeBronec J., Morel P., Robe M.C., Maurenas J.M., Faivre-Pierret R., Martin D., Sabroux J.C., Zettwog P.(1991). Eruptive and diffusive Emissions of CO₂ from Mount Etna; Nature, 351, 387-391
- [5] Bluth-G-J-S, Schnetzler-C-C, Krueger-A-J, Walter-L-S. (1993). The contribution of explosive volcanism to global atmospheric sulphur dioxide concentrations. Nature 366 (6453), 327-329.
- [6] Fujita-S. (1993). Volcanic activity: some effects of the emissions on the acidification of the environment. Journal of Japan Society of Air Pollution; vol 28; no 2; pp 72-90.
- [7] Zreda-Gostynska-G, Kyle-P-R, Finnegan-D-L. (1993). Chlorine, fluorine, and sulfur emissions from Mount Erebus, Antarctica and estimated contributions to the Antarctic atmosphere. Geophysical Research Letters 20 (18), 1959-1962.
- [8] Stoiber R.E., Williams S.N., Huebert B. (1987). Annual Contribution of Sulfur Dioxide to the Atmosphere by Volcanoes. J. Volcan.geotherm.Res.33, 1-8.
- [9] Gerlach-T-M. (1991). Present-Day CO₂ Emissions from Volcanos. EOS Trans (American-Geophysical-Union), Jun 4, 91, 72 (23), 249 (3).
- [10] Williams S.N., Schaefer S.J., Calvache V. M. L., Lopez D.(1992). Global carbon dioxide emission to the atmosphere by volcanoes. Geochimica et Cosmochimica Acta, 56, 1765-1770.
- [11] Ammann-M; Burtscher-H; Siegmann-H-C. (1990). Monitoring volcanic activity by characterization of ultrafine aerosol. Journal of Aerosol Science 21 (Supp 1), 275-278.

- [12] Ferrara-R, Maserti-B-E, De-L-A, Cioni-R, Raco-B, Taddeucci-G, Edner- H. (1994). Atmospheric Mercury Emission at Solfatara Volcano (Pozzuoli, Phlegraen Fields-Italy). *Chemosphere*, 29 (7) 1421(8).
- [13] Hoff-R-M, Gallant-A-J. (1980). Sulfur dioxide emissions from La Soufriere Volcano, St. Vincent, West Indies. *Science*, 209(4459), 923-924.
- [14] Gerlach-TM; McGee-KA (1994). Total sulfur dioxide emissions and pre-eruption vapor-saturated magma at Mount St. Helens, 1980-88. *Geophysical Research Letters* 21 (25) 2833-2836.
- [15] McPeters Richard D (1993). The atmospheric budget for Pinatobu derived from NOAA-11 SBUV/2 spectral data. *Geophys.Res.Lett.* 20(18), 1971-74.
- [16] Haraden J. (1989). CO₂ production rates for geothermal energy and fossil fuels. *Energy* 14, 867-873.

18 BIBLIOGRAPHY

19 RELEASE VERSION, DATE AND SOURCE

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1 ACTIVITIES INCLUDED

Natural gas as a product of microbial degradation derives from a variety of geological periods and therefore is stored under very different conditions in the earth's crust. While the very old storage chambers tend to be very well sealed (otherwise they would have been emptied in the time since formation), seeping of natural gas may occur at storage sites of more recent times, like from the glaciation periods. Several of such gas seeping sites have been observed, most easily under water when methane bubbles rise from a lake or the sea floor.

Gas seeps from natural gas reservoirs can not easily be distinguished from gas being developed from organic material buried in the sediment. However differentiation should be made to methane production in soils, which is described in the "wetlands" chapter (proposed SNAP 1105). The difference between these two sources of freshly produced methane is primarily in the age of the organic material to be processed, which may be thousands of years old in the first case, but is fresh material (also with respect of ^{14}C content) in the latter case. Also, the amount of material can be assessed from vegetation density in the latter case.

2 CONTRIBUTIONS TO TOTAL EMISSIONS

Due to the irregular emission patterns, only rough estimations are possible. A compilation of reports on gas seepages indicates that globally emissions may be between 8 and 65 Tg CH_4 per year [1] (approx. 2-13 % of global annual CH_4 emissions).

This activity is not believed to be a significant source of $\text{PM}_{2.5}$ (as of December 2006).

3 GENERAL

3.1 Description

Gas seeps from natural gas reservoirs or from reservoirs of organic matter occur both under shallow sea surfaces as well as at land surfaces. However only seeps under water are easily identified due to formation of gas bubbles.

Natural gas reservoirs may start seeping gas after seismic activities (earthquakes) or also depending on the outside temperature. Emissions depend on the emission rate and the size of the seep area. While a number of gas seeps have been observed, in general they seem to be highly limited in extent.

In order to be able to quantify emissions from this source, steady flow and homogeneous composition of the seeps need to be assumed. This is however not the real situation. In fact, outbursts of activity have been observed that indicate some methane formation is still active and the reservoirs are being emptied periodically. This would indicate that maximum observed emission rates should not be considered typical [2].

Submarine emissions may occur also from the deep ocean. Here only seeps from the continental shelves are considered, as most of deep ocean emissions will be dissolved in sea water long before reaching the surface.

3.2 Definitions

3.3 Techniques

Emissions from underground gas storage would be expected to be steady flow. Observed gas flow however is strongly variable, in some instances taking place primarily during summer and fall, and preferably during low tide [3]. In these cases, methane is assumed to be produced by on-going processes from buried organic material. ^{14}C depletion in the resulting methane emission proves that this material is from fossil origin [1]. In addition to bacterial production, thermogenic production is assumed, with the associated emissions being accompanied by oil seeps.

While gas formation, especially for bacterial production, may depend on the ambient temperature and decrease considerably during wintertime, gas release may also depend on other parameters. High pressure (as at high tide) or a low degree of filling of the underground storage (after a previous outburst) may halt emissions for some time.

3.4 Emissions

Natural gas emissions are primarily methane (CH_4), but at a lower rate also other alkanes may be released.

3.5 Controls

There is no control to natural emissions by definition.

4 SIMPLER METHODOLOGY

Size and location of seeping areas need to be obtained from geological offices, research institutions or petrol companies. The emissions are then calculated for each of these areas separately by:

$$E = F \times A$$

A is the area in m^2 , F the average flux per m^2 (see section 8).

5 DETAILED METHODOLOGY

There is no state-of-the-art methodology.

6 RELEVANT ACTIVITY STATISTICS

There are no statistical data available. The activity (number and size of fields where seeping is taking place) may be obtained from research institutions, geological offices, or petrol companies.

7 POINT SOURCE CRITERIA

No point sources.

8 EMISSION FACTORS, QUALITY CODES AND REFERENCES

As indicated earlier, emission factors are highly uncertain due to the irregularity of the emissions taking place. Emission rates of 10 l/h from one single seep hole have been reported near the California South Coast [4], but also the area-based value of 1 l/h/m² in the Danish coastal waters [5]. These values need to be considered absolute maxima, with typical values at least a factor of 20 lower [1, 3].

According to a compilation of all known source areas [1], the highest overall seepage rate is given for the California South Coast (Santa Barbara Channel) at 400 g/yr/m². As a typical emission factor, the numbers given for the Gulf of Mexico, the North Carolina coast and the Danish Kattegat and Skagerrak should be used: 50 g/yr/m². These factors refer to active areas only. Whenever available local information should be used however, as this number can only give a rough guidance for orientation.

For those emissions taking place at larger depths, the dissolution of methane in sea water from rising gas bubbles needs to be considered. This uptake has been discussed in more detail [1] with respect to bubble size. As a first guideline, it may be assumed that only 50 % of the emissions at 100 m depth will reach the surface.

9 SPECIES PROFILES

A typical profile for gas seeps is suggested in [4], at 75 % methane, 7 % each propane and n-butane and 6 % ethane (by weight).

10 UNCERTAINTY ESTIMATES

The uncertainty with the emission factors is assumed to be in the range of at least one order of magnitude, data quality E.

11 WEAKEST ASPECTS/PRIORITY AREAS FOR IMPROVEMENT IN CURRENT METHODOLOGY

More information on activity rates have to be obtained. Especially flux measurements are needed in areas where these emissions are known to take place. Ambient methane concentrations near potential emission areas should be observed for a prolonged period of time in order to determine duration and strength of possible active and quiet periods.

12 SPATIAL DISSAGGREGATION CRITERIA FOR AREA SOURCES

Even distribution within each field.

13 TEMPORAL DISSAGGREGATION CRITERIA

Constant emission flux is assumed, as otherwise temporal disaggregation would have to be based on observations, event based.

14 ADDITIONAL COMMENTS

Natural gas seeps are not at all influenced by humans.

15 SUPPLEMENTARY DOCUMENTS

16 VERIFICATION PROCEDURES

17 REFERENCES

- [1] Hovland M., Judd A.G., Burke R.A. (1993). The global flux of methane from shallow submarine sediments. *Chemosphere* 26, 559-578.
- [2] Iversen N. (1997). Aalborg University, DK, personal information.
- [3] Martens C.S., Klump J.V. (1980), Biogeochemical cycling in a organic-rich coastal marine basin. 1. Methane sediment-water exchange processes. *Geochim. Cosmochim. Acta* 44, 471-490.
- [4] Radian Corporation (1996). EIIP Volume 5, biogenic sources preferred methods. Final report to the Area Sources Committee, Emission Inventory Improvement Program, May 1996.
- [5] Fenger J., Fenhann J., Kilde N. (1990). Danish budget for greenhouse gases. Nord 1990:97, Nordic council of ministers, Copenhagen, 1990.

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SNAP CODE:	111000
SOURCE ACTIVITY TITLE:	OTHER SOURCES AND SINKS <i>Lightning</i>
NOSE CODE:	301.10.01
NFR CODE:	N/A

1 ACTIVITIES INCLUDED

Lightning and corona discharge during thunderstorm events cause atmospheric chemical reactions to take place at high voltages and high temperatures. These reactions cause the production of NO_x in the atmosphere. Such production processes are, strictly speaking, not real emissions as the compounds involved (primarily N₂ and O₂) are not injected into the atmosphere but are present anyway. However as these processes can not adequately be described by conventional atmospheric models on one hand, and their impact is eventually identical to those from (anthropogenic) emissions on the other hand, they are easy to be compared on the emission level and thus are frequently treated as such.

2 CONTRIBUTIONS TO TOTAL EMISSIONS

Global NO_x production by lightning has been estimated in the range of 3-5 TgN/yr [1]. For the U.S., 40% of the yearly lightning-generated NO is estimated to be produced during the summer months [2]. Other estimates using the calculation schemes given below indicate that the lightning NO comprises only 3% of the total NO_x emissions budget, with a maximum contribution of 24% at the maximum hour with respect to anthropogenic emissions at a definite period in summer for the U.S. North-East [3].

These figures apply to emissions within the whole troposphere. Emissions in the boundary-layer (circa lowest 1 km) are obviously considerably less. In reporting emissions under the joint EMEP/CORINAIR system, care must be taken to report only emissions between the ground and 1 km - see section 8.

This activity is not believed to be a significant source of PM_{2.5} (as of December 2006).

3 GENERAL

3.1 Description

The electrical discharge of lightning creates plasma channels in the atmosphere characterized by the high fraction of ionic loads and high temperatures. Major compounds of the atmosphere, notably nitrogen, oxygen, and water, may be ionized and then undergo chemical transformation. While the exact pathway of such transformations is largely unknown, a few assessments have been made [4, 5]. A major species to be produced is nitric oxide (NO), but also other compounds containing nitrogen, oxygen and hydrogen atoms are being formed. Crucial for the formation is the high temperature during the flash (up to 30000 K) and the

subsequent rapid cooling below 1500 K which prevents the freshly formed NO from immediate destruction.

3.2 Definitions

- Lightning: Atmospheric discharge during thunderstorm events
- CG - discharge: Flash starting in the cloud, bringing several coulomb of negative charge to earth within about 0.5 s (negative discharge)
- IC - discharge: Flash that does not connect to earth: intracloud, intercloud and cloud-to-air flashes

3.3 Techniques

While lightning exhibits different characteristics depending on whether it is cloud-to-ground (CG), cloud-to-cloud or within-cloud (inter- and intra-cloud, IC), emission estimation techniques have not been resolved to this level of detail. It has been reported that IC discharges may be up to ten times less efficient in producing NO_x than the CG discharges [4]. However, newer information suggests that these discharges may be nearly equal [1, 5]. The amount and distribution of NO produced is believed to be dependent on the energy and the frequency of lightning strokes, which in turn is dependent on cloud temperatures and cloud heights. IC lightning is known to be more frequent than CG lightning. The ratio has been correlated to the cold cloud thickness (cold cloud = below freezing), representing the size of the electric field involved that may determine the number of IC flashes [6]. Despite of generally large variations of this ratio, a dependence on geographical latitude has been found using cold cloud thickness as a parameter.

3.4 Emissions

Out of the compounds being formed in lightning discharges, numbers are given only for NO and NO₂ as NO_x. These emissions seem to be the most relevant.

Differentiation needs to be made for IC and CG lightning, especially with respect to their injections into the atmosphere as relevant for models. IC lightning occurs at altitudes above about 5 km and may be neglected in some boundary layer models, while CG lightning is expected to reach from the ground to about 7 km high (north of 30 deg. Latitude) or 10 km high (south of 30 deg. Latitude). The NO formed is distributed decreasing with height as a function of air density [7]. For a 7 km flash, about 20% of the emissions would then occur in the lowest 1000m, and 80% between ground level and 5 km.

In [3] the IC component -- here only calculated in the detailed methodology -- is assumed to add an extra 21% NO at 60 deg. N and 61% at the equator. All IC flashes are assumed to occur above a height of 5 km.

3.5 Controls

There are no controls to natural emissions by definition.

4 SIMPLER METHODOLOGY

The number of lightning flashes can be obtained from measurements (see section 5), or from estimations. In the latter case, the flashes are estimated from meteorological data on thunderstorm occurrence and from the geographical latitude of the area considered. Support for these estimations may be given by satellite data [8]. Emissions are then calculated according to [3] :

$$CG_{NO} = E \times M$$

where CG_{NO} is the NO produced by the cloud-to-ground part of the lightning flash

$E = 4 \times 10^8$ J per cloud-to-ground flash and

$M = 9 \times 10^{16}$ molecules NO / J

Calculated in mass units, this yields 2.75 kg NO_x (calculated as NO_2) per flash of lightning. About 20% of this amount is assumed to be emitted below 1 km altitude, 80 % below 5 km altitude.

5 DETAILED METHODOLOGY

The difference of the detailed methodology is that the number of lightning flashes is actually counted and cloud-to-cloud flashes are included. Data for the U.S. are available from the East Coast lightning detection network, or from the lightning strike data archive from Global Atmospheric, Inc. in Tucson, AZ. In many European countries, especially in Western Europe, national networks are operative and may be accessible through the respective national meteorological service. These networks do not include cloud-to-cloud (IC) flashes however.

Emissions now are calculated as [9]:

$$LNO = (N_{CG} \cdot EF_{CG} / \epsilon_{CG}) + [(N_{CG} / \epsilon_{CG}) \cdot (10/(1 + (\Phi/30)^2) - 1)] \cdot EF_{IC}$$

where:

LNO = NO emissions for lightning flashes in study area, molecules NO

N_{CG} = Number of cloud-to-ground (CG) flashes recorded

ϵ_{CG} = Efficiency of the CG network

EF_{CG} = Emission factor for NO for each CG lightning flash

Φ = Latitude of the study area in degrees

EF_{IC} = Emission factor for NO for each inter- or intra - cloud (IC) lightning flash

The emission factors needed for calculation are given in section 8. For the U.S. East Coast, the efficiency has been reported to be 0.7 [9]. The equation takes care of the fact that IC lightning is, dependent on the latitude, about four times as frequent as CG lightning. While recent investigations [6] indicate a less pronounced latitude dependence than the one given here, the results are virtually identical at 40 deg. Latitude.

Emissions from IC flashes are assumed to take place above 5 km altitude only. In contrast, 80% of the CG-lightning emissions are expected at altitudes below 5 km, and 20% even below 1 km.

6 RELEVANT ACTIVITY STATISTICS

Meteorological data on lightning frequencies need to be obtained. Lightning climatologies are being produced or are available in the meteorological offices of many countries. The data may either derive from reporting thunderstorm events, from observation networks, or from satellite information.

7 POINT SOURCE CRITERIA

No point sources.

8 EMISSION FACTORS, QUALITY CODES AND REFERENCES

A large variety of emission factors is given in the literature from laboratory as well as field investigations. According to [10], the low, median and high end of these estimates may be given at 0.36×10^{26} , 4×10^{26} , and 30×10^{26} molecules NO per flash. As indicated in [10] and also discussed in [3], the highest of these figures (from [11]) is not supported by studies modelling nitrate deposition. [1] estimates global NO_x production from a best fit approximation between a global model and observations from regions where lightning is expected to be a major source. This study yields results close to the lowest of the three factors given, thus we recommend to apply the factor of 0.36×10^{26} molecules NO (2.75 kg NO_x) for each flash of lightning as EF_{CG} . Only part of these emissions should be reported: See Table 8.1.

Literature data [4, 9] suggest an emission factor for IC lightning of an order of magnitude lower than for CG lightning. We tentatively are recommending that EF_{IC} be set equal to 0.36×10^{25} molecules NO (275 g NO_x). Recent theoretical considerations [5] however indicate that such a low emission factor might not be realistic. The total energy dissipated in an IC flash should then be at least as high as in GC flashes. Even considering the decreased NO formation at high altitude the EF_{IC} should be considerably higher then (maybe a factor 5). However quantification is missing, and the emissions only concern altitudes above 5 km anyway (where they are relevant primarily on the global scale). Thus an update will only be given at a later stage, when new evidence emerges.

All recommended emission factors are compiled in the following table for the respective altitudes. Note that reporting will only be necessary for lightning emissions up to one km at this stage. The upper layer emissions may be needed at a later stage only.

Table 8.1: Recommended emission factors per flash of lightning in molecules NO and kg NO_x (calculated as NO₂), respectively.

ALTITUDE	EF _{CG} (molecules)	EF _{CG} (kg)	EF _{IC} (molecules)	EF _{IC} (kg)	COMMENT
below 1 km	0.72x10 ²⁵	0.55	0	0	report
1km to 5 km	2.16x10 ²⁵	1.65	0	0	do not report !
above 5 km	0.72x10 ²⁵	0.55	0.36x10 ²⁵	0.275	do not report !
total	3.6 x10 ²⁵	2.75	0.36x10 ²⁵	0.275	do not report !

Because of the uncertainty in the NO production factors, we assume a quality code of D. Additional information on uncertainties can be obtained from [1] and [12].

9 SPECIES PROFILES

Emissions are given for NO_x. While virtually all of the oxidation product is originally NO, a considerable part is transformed to NO₂ very quickly (depending on ozone availability etc.). This part may be in the order of 25% of the original NO [11].

10 UNCERTAINTY ESTIMATES

The uncertainty with the emission factors has been estimated a factor of three, however the validity of these results have to be checked with respect to those literature estimates giving results different by up to an order of magnitude (see [5] and [10]).

11 WEAKEST ASPECTS/PRIORITY AREAS FOR IMPROVEMENT IN CURRENT METHODOLOGY

Depending on the methodology of assessing the emission factors, there are still large discrepancies. These have to be settled before any more detailed estimations can be performed. In addition, the chemical conversion processes in lightning need to be better understood, especially with regard to IC lightning.

12 SPATIAL DISAGGREGATION CRITERIA FOR AREA SOURCES

Spatial disaggregation can be performed according to the distribution of lightning and thunderstorm events.

13 TEMPORAL DISAGGREGATION CRITERIA

Temporal disaggregation should be done according to diurnal and annual cycle of thunderstorm / lightning activity.

14 ADDITIONAL COMMENTS

Lightning is not known to be influenced by humans at all, thus it should be considered as a purely natural source.

15 SUPPLEMENTARY DOCUMENTS**16 VERIFICATION PROCEDURES****17 REFERENCES**

- [1] Levy H., Moxim W., Kasibhatla P. (1996). A global three-dimensional time-dependent lightning source of tropospheric NO_x. *J.Geophys.Res.*, 101, 22911-22922.
- [2] Placet M., Battye R., Fehsenfeld F., Basset G. (1990). Emissions involved in acid deposition processes. NAPAP State of Science and Technology Report 1, Chapter 5, pp. 9-19.
- [3] Novak J., Pierce T. (1993). Natural emissions of oxidant precursors. *Water, Air and Soil Pollution*, 67, 57-77.
- [4] Sisterson D., Liaw Y. (1990). An Evaluation of Lightning and Corona Discharge on Thunderstorm Air and Precipitation Chemistry. *Journal of Atmospheric Chemistry*, 10 (1) 83-96.
- [5] Gallardo L., Cooray V. (1996). Cloud cloud-to-cloud discharges be as effective as cloud-to-ground discharges in producing NO_x? *Tellus*, 48B, 641-651.
- [6] Price C., Rind D. (1993). What determines the cloud-to-ground lightning fraction in thunderstorms? *Geophys.Res.Lett.* 20, 463-466.
- [7] Pierce T., Novak J. (1991). Estimating Natural Emissions for EPA's Regional Oxidant Model. Presented at Environmental Protection Agency/Air and Waste Management Association International Specialty Conference on Emission Inventory Issues in the 1990's, Durham, NC., September 9-12, 1991. 14p.
- [8] Turman B., Edgar B. (1982). Global lightning distribution at dawn and dusk. *J. Geophys. Res.* 87, 1191-1206.
- [9] RADIANT Corp. (1996). EIIP Volume 5, Biogenic sources preferred methods. Final report to the Area Sources Committee, Emission Inventory Improvement Program, May 1996.
- [10] Biazar A, McNider R. (1995). Regional estimates of lightning production of nitrogen oxides. *J.Geophys.Res.* 100, 22861-22874.
- [11] Franzblau E., Popp C. (1989). Nitrogen oxides produced from lightning. *J. Geophys. Res.* 94, 11089-11104.

- [12] Lawrence M., Chameides W., Kasibhatla P., Levy H., Moxim W. (1995) Lightning and atmospheric chemistry: the rate of atmospheric NO production. In: Handbook of Atmospheric Electrodynamics, vol. 1, edited by H. Volland, pp. 189-202, CRC Press, Boca Raton, Florida, USA.

18 BIBLIOGRAPHY

19 RELEASE VERSION, DATE AND SOURCE

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December 2006

20 POINT OF ENQUIRY

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SNAP CODE:	112101
	112102
	112103
	112104
	112105

SOURCE ACTIVITY TITLE:	OTHER SOURCES AND SINKS:
	CHANGES IN FOREST AND OTHER WOODY BIOMASS STOCKS
	<i>Tropical Forests</i>
	<i>Temperate Forests</i>
	<i>Boreal Forests</i>
	<i>Grassland/tundra</i>
	<i>Other</i>

NOSE CODE:	301.21.01
	301.21.02
	301.21.03
	301.21.04
	301.21.05

NFR CODE:	N/A
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A specific methodology for these activities has not been prepared yet as this is a new sub group. It will be investigated this year.

The expert panel leaders for this sub group are listed below.

These activities are not believed to be a significant source of PM_{2.5} (as of December 2006).

Updated with particulate matter details by:

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