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Atmospheric Chemistry and Physics Discussions

Mass spectrometric measurement of hydrogen isotope fractionation for the reactions of chloromethane with OH and Cl

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Abstract

Chloromethane (CH_3Cl) is an important provider of chlorine to the stratosphere but yet lacks detailed knowledge of its budget. Stable isotope analysis is potentially a powerful tool to constrain CH_3Cl flux estimates. The largest degree of isotope fractionation is expected to occur for deuterium in CH_3Cl in the hydrogen abstraction reactions with its main sink reactant tropospheric OH and its minor sink reactant Cl atoms. We determined the isotope fractionation by stable hydrogen isotope analysis of the fraction of CH_3Cl remaining after reaction with hydroxyl and chlorine radicals in a 3.5 m^3 Teflon smog-chamber at 293 ± 1K. We measured the increasing stable hydrogen isotope values of the unreacted CH_3Cl using compound specific thermal conversion isotope ratio mass spectrometry. The isotope fractionations of CH_3Cl for the reactions with hydroxyl and chlorine radicals were found to be -242 ± 7 mUr (or ‰) and -280 ± 11 mUr, respectively. For comparison, we performed

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similar experiments using methane (CH_4) as the target compound with OH and obtained a fractionation constant of -205 ± 6 mUr which is in good agreement with values previously reported. The observed large kinetic isotope effects are helpful when employing isotopic analyses of CH_3Cl in the atmosphere to improve our knowledge of its atmospheric budget.

1 Introduction

Chloromethane (often named methyl chloride) is the most abundant chlorine containing trace gas in the Earth's atmosphere, currently with a global mean mixing ratio of ~540 ± 5 parts per trillion by volume (pptv) and an atmospheric lifetime of 1.0–1.2 years (Carpenter et al., 2014). The global emissions of CH_3Cl have been estimated to be in the range of 4 to 5 Tg yr^{-1} (1 Tg = 10^{12} g) stemming from predominantly natural but also anthropogenic sources (Montzka and Fraser, 2003; WMO, 2011; Carpenter et al., 2014). However, current estimates of the CH_3Cl global budget and the apportionment between sources and sinks are still highly uncertain. Known natural sources of CH_3Cl include tropical plants (Yokouchi et al., 2002; Yokouchi et al., 2007; Umezawa et al., 2015), wood-rotting fungi (Harper, 1985), oceans (Moore et al., 1996), plants of salt marshes (Rhew et al., 2003; Rhew et al., 2000), aerated and flooded soil (Redeker et al., 2000; Keppler et al., 2000), senescent leaves and leaf litter (Hamilton et al., 2003; Derendorp et al., 2011) and wild fires. Anthropogenic CH_3Cl release to the atmosphere comes from the combustion of coal and biomass with minor emissions from cattle (Williams et al., 1999) and humans (Keppler et al., 2017). In addition, it has been reported that emissions from industrial sources, particularly in China, might be much higher than previously assumed (Li et al., 2016).

The dominant sink for atmospheric CH_3Cl results from the reaction with photochemically-produced hydroxyl radicals (OH), currently estimated at about 2.8 Tg yr^{-1} (Carpenter et al., 2014). Furthermore, in the marine boundary layer the reaction of CH_3Cl with chlorine radicals (Cl) represents another sink estimated to account for up to 0.4 Tg yr^{-1} (Khalil et al., 1999; Montzka and Fraser, 2003). Microbial CH_3Cl degradation in soils may be a relevant additional global sink (McAnulla et al., 2001; Harper et al., 2003; Miller et al., 2004; Jaeger et al., 2018a) but its impact on the global CH_3Cl budget is still highly uncertain. The microbial CH_3Cl soil sink strength has been estimated to range from 0.1 to 1.6 Tg yr^{-1} (Harper et al., 2003; Keppler et al., 2005; Carpenter et al., 2014). Moreover, small proportions of tropospheric CH_3Cl are lost to the stratosphere (146 Gg yr^{-1}, 1 Gg = 10^9 g) and

to cold polar oceans (370 Gg yr^{-1}) though oceans in total are a net source (Carpenter et al., 2014).

A potentially powerful tool in the investigation of the budgets of atmospheric volatile organic compounds is the use of stable isotope ratios (Brenninkmeijer et al., 2003; Gensch et al., 2014). Stable isotope analysis, when used in combination with CH_3Cl flux measurements, has the potential to better constrain the atmospheric CH_3Cl budget as suggested by Keppler et al. (2005) and Saito & Yokouchi (2008). The isotopic composition of tropospheric CH_3Cl depends on the isotopic source signatures and the kinetic isotope effects (KIE) of the sinks. Several studies have investigated the stable carbon isotope source signature of CH_3Cl produced via biotic and abiotic processes, however, for a more detailed overview we refer readers to the studies reported by Keppler et al. (2005) and Saito & Yokouchi (2008). Moreover, a few studies have measured the KIE of stable carbon isotopes of CH_3Cl during oxidation or biodegradation by bacterial isolates (Miller et al., 2001; Nadalig et al., 2013; Nadalig et al., 2014) or in soils under laboratory conditions (Miller et al., 2004; Jaeger et al., 2018a). The first, and so far, only available analysis of the KIE for reaction of CH_3Cl with OH has been reported by Gola et al. (2005) and revealed an unexpectedly large stable carbon isotope fractionation. The experiments were carried out in a smog chamber using long path Fourier-transform infrared spectroscopy (FTIR) detection. However, in view of the unexpected isotope fractionation we consider it important to confirm this result using other measurement methods such as stable isotope ratio mass spectrometry (IRMS).

So far isotopic investigations of CH_3Cl have predominantly focused on stable carbon isotope measurements. Stable hydrogen isotope measurements including both sources and sinks of CH_3Cl have become available only recently (Greule et al., 2012; Nadalig et al., 2014; Nadalig et al., 2013; Jaeger et al., 2018b; Jaeger et al., 2018a). Moreover, relative rate experiments have been carried out for three isotopologues of CH_3Cl and their reactions with Cl and OH. The OH and Cl reaction rates of CH_2DCl were measured by long-path FTIR spectroscopy relative to CH_3Cl at 298 ± 2 K and 1 atm (Sellevåg et al., 2006) (Table 1).

In this manuscript, using a 3.5 m^3 Teflon smog chamber and IRMS measurements, we present results from kinetic studies of the hydrogen isotope fractionation in the atmospheric OH and Cl loss processes of CH_3Cl. Furthermore, we also measured the isotope fractionation for the reaction between methane (CH_4) and OH using a similar experimental design and compared this value with those from previous studies.

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2 Materials and Methods

2.1 Smog chamber experiments with chloromethane

The isotope fractionation experiments were performed in a 3.5 m^3 Teflon smog-chamber (fluorinated ethylene propylene, FEP 200A, DuPont, Wilmington, DE, USA) with initial CH_3Cl mixing ratio of about 10 parts per million by volume (ppmv). Atomic chlorine were generated via photolysis of molecular chlorine (Cl_2) (Rießner Gase, 0.971% Cl_2 in N_2) by a solar simulator with an actinic flux comparable to the sun in mid-summer in Germany (Bleicher et al., 2014). Hydroxyl radicals were generated via the photolysis of ozone (O_3) at 253.7 nm in the presence of water vapor (RH = 70%) (produced by double-distilled water in a three-neck bottle humidifier) and/or H_2. To obtain efficient OH formation, a Philips TUV lamp (55 W) was welded in Teflon film (FEP 200) and mounted inside the smog chamber. O_3 was monitored by a chemiluminescence analyzer (UPK 8001). The chamber was continuously flushed with purified, hydrocarbon-free zero air (zero-air-generator, cmc instruments, <1 ppbv of O_3, <500 pptv NO_X, <100 ppbv of CH_4) at a rate of 4 L min^{-1} to maintain a slight overpressure of 0.5-1 Pa logged with a differential pressure sensor (Kalinsky Elektronik DS1). The quality of the air inside the chamber in terms of possible contamination was controlled by monitoring NO and NO_X (EcoPhysics CLD 88p, coupled with a photolytic converter, EcoPhysics PLC 860). Perfluorohexane (PFH) with an initial mixing ratio of ~25 parts per billion by volume (ppbv) was used as an internal standard to correct the resulting concentrations for dilution. The temperature was set to 20±1°C and monitored, together with the relative humidity, by a Teflon-cased sensor (Rotronic, HC2-IC102). To guarantee constant mixing and small temperature gradients, a Teflon fan was mounted and operated inside the chamber. More detailed specification of the smog chamber can be found elsewhere (e.g. Wittmer et al., 2015). The mixing ratios of CH_3Cl and PFH were monitored by gas chromatography-mass spectrometry (GC-MS, Agilent Technologies, Palo Alto, CA) with a time resolution of 15 minutes throughout the experiments. Aliquots (5 ml) were withdrawn from the chamber with a gas tight syringe, injected into a stream of He (30 ml min-1) and directed to a pre-concentration unit that was attached to the GC-MS. The pre-concentration unit consisted of a simple 8 port valve (VICI Valco) equipped with two cryotraps made of fused silica, which were immersed in liquid nitrogen for trapping the analytes. Prior to each sample measurement, a gaseous standard (5 ml of 100 ppmv CH_3Cl in N_2) was measured. Figure 1 shows the design of the smog chamber used in our experiments.

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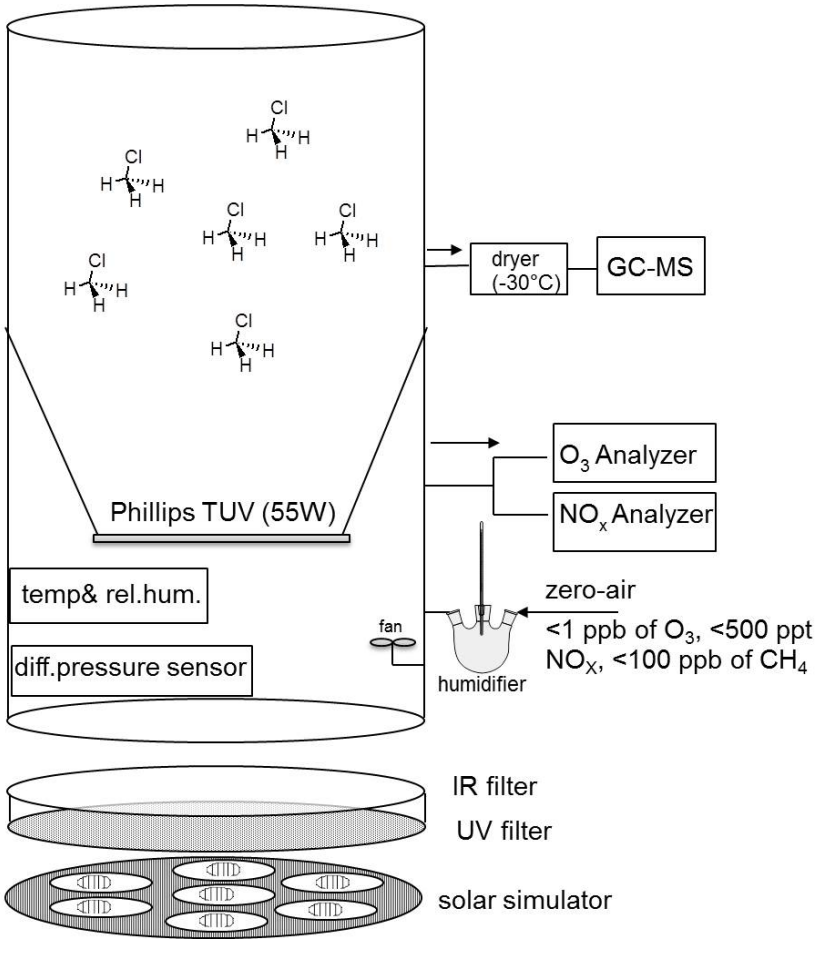
In the CH_3Cl and OH experiments (1 and 2) 2000 ppmv H_2 was used to scavenge chlorine atoms originating from the photolysis or oxidation of formyl chloride (HCOCl), which forms as an intermediate in the reaction cascade. Under the experimental conditions typically more than 70% of the CH_3Cl was degraded within 7 to 10 h. From each experiment (CH_3Cl + OH and CH_3Cl + Cl) 10 to 15 canister samples (2 L stainless steel, evacuated $<10^{-4}$ mbar) were collected at regular time intervals for subsequent stable hydrogen isotope measurements at Heidelberg University.

Figure 1: Scheme of the experimental smog chamber



2.2 Smog chamber degradation experiments with methane

The CH_4 degradation experiments were carried under the same conditions as the CH_3Cl degradation experiments but without PFA as an internal standard. Instead we used the flushing flow rate of zero air to account for the dilution during the experiment. The initial CH_4 mixing ratio was 6 ppmv. Throughout these experiments CH_4 mixing ratios were monitored with a Picarro G225i cavity ring down spectrometer directly connected to the chamber.

2.3 Stable hydrogen isotope analysis using isotope ratio mass spectrometry

2.3.1 Chloromethane

Stable hydrogen isotope ratios of CH_3Cl were measured by an in-house built cryogenic pre-concentration unit coupled to a Hewlett Packard HP 6890 gas chromatograph (Agilent Technologies, Palo Alto, CA) and an isotope ratio mass spectrometer (IRMS) (Isoprime, Manchester, UK) as described in detail by Greule et al. (2012). Diverging from the method of Greule et al. (2012) a ceramic tube reactor without chromium pellets at 1450°C was instead used for high-temperature conversion (HTC). A tank of high-purity H_2 (Alphagaz 2, hydrogen 6.0, Air Liquide, Düsseldorf, Germany) with a δ^2H value of ~-250 mUr (milliurey = 0.001 = 1‰, cf. section below) versus VSMOV was used as the working gas. All measured sample δ^2H values were monitored for their relative trueness by analyzing an in-house working standard of known δ^2H value. The CH_3Cl working standard was calibrated against IAEA standards NBS 22, LVSEC (carbon), VSMOW and SLAP (hydrogen) using TC/EA-IRMS (elemental analyser-isotopic ratio mass spectrometer, IsoLab, Max Planck Institute for Biogeochemistry, Jena, Germany) resulting in a δ^2H value of -140.1 ± 1.0 mUr vs. VSMOW ($n = 10$, 1σ). The H_3^+ factor, determined daily during this investigation (two different measurement periods), was in the range of 5.75 - 6.16 (first period) and 8.90 - 9.21 (second period). The mean precision based on replicate measurements ($n = 6$) of the CH_3Cl working standard was 2.1 mUr and 3.8 mUr for the first and second measurement periods, respectively. Samples were analyzed three times ($n = 3$), and the standard deviations (SD) of the measurements were in the range of 1.2 to 103.8 mUr. Lowest SD were observed for samples with lowest δ^2H values (~-140 mUr) and highest mixing ratios and higher SD for samples with highest δ^2H values (~+800 mUr) and lowest mixing ratios.

To comply with International System of Units (SI) guidelines, we follow the proposal of Brand and Coplen (2012) and use the symbol Ur, after H.C. Urey (Urey, 1948), as the isotope delta (Coplen, 2011) value unit. Thus, an isotope-delta value expressed traditionally as −50 ‰ can be written −50 mUr. Similarly as for the δ^2H values, throughout the manuscript we also report the isotope enrichment factor ε in mUr.

Please note that the above described 1-point calibration of the δ^2H data might be affected by an additional error ("scale compression") and particularly might affect the uncertainties of the very positive δ^2H values. Unfortunately CH_3Cl working standards with distinct isotopic signatures spanning the full range of measured δ^2H values (-150 to ~+800 mUr) are not currently available to eliminate or minimize such an error.

2.3.2 Methane

Stable hydrogen isotope ratios of CH_4 were analyzed using an in-house built cryogenic pre-concentration unit coupled to a Hewlett Packard HP 6890 gas chromatograph (Agilent Technologies, Palo Alto, CA) and an isotope ratio mass spectrometer (DeltaPlus XL, ThermoQuest Finnigan, Bremen, Germany). The working gas was the same as that used for δ^2H analysis of CH_3Cl (c.f. section 2.3.1.).

All δ^2H values obtained from analysis of CH_4 were corrected using two CH_4 working standards (isometric instruments, Victoria, Canada) calibrated against IAEA and NIST reference substances (not specified by the company). The calibrated δ^2H values of the working standard in mUr vs. V-SMOW were -144 ± 4mUr and -138 ± 4 mUr.

The H_3^+ factor determined daily during the two week measurement period was in the range 2.38–2.43. The daily average precision based on replicate measurements of the CH_4 working standard was 4.9 mUr (n = 7). Samples were analyzed 3 times (n = 3), and the SD of the measurements were in the range of 1.4 to 40.9 mUr. Lowest SD were observed for samples with lowest δ^2H values (~-140 mUr) and highest mixing ratios and higher SD for samples with highest δ^2H values (~+800 mUr) and lowest mixing ratios.

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2.4 Kinetic isotope effect, fractionation constant α and the isotope enrichment constant ε

The isotopic composition of atmospheric compounds might be altered by the kinetic isotope effects of physical, chemical or biological loss processes. The kinetic isotope effect (KIE) is usually defined as:

$$KIE = \frac{k_1}{k_2} \quad (1)$$

where k_1 and k_2 are the reaction rate constants for loss of the lighter and the heavier isotopologues, respectively. The KIE is typically expressed as isotope fractionation ε (also termed isotope enrichment constant) or isotope fractionation constant α.

In this study the isotope fractionation constant α and the isotope enrichment constant ε are derived from the slope of the Rayleigh plot according to (Clark and Fritz, 1997; Elsner et al., 2005) and equation 2:

$$\ln\frac{R_t}{R_0} = ln\left(\frac{\delta^2H_t+1000}{\delta^2H_0+1000}\right) = ln\frac{(\delta^2H_0+\Delta\delta^2H+1)}{(\delta^2H_0+1)} \cong (\alpha-1)\cdot lnf = \varepsilon\cdot lnf \quad (2)$$

Where R_t and R_0 are the $^2H/^1H$ ratios in CH_3Cl or CH_4 at the different time points and time zero, respectively, and f is the remaining CH_3Cl or CH_4 fraction at the different time points. Negative values of ε indicates that the remaining CH_3Cl or CH_4 is enriched in the heavier isotope and corresponds to a $\alpha<1$, meaning that over the entire experiment, the heavier CH_3Cl or CH_4 react by this factor more slowly than the lighter CH_3Cl or CH_4.

The kinetic isotope effect is then calculated as:

$$KIE = \frac{1}{\alpha} \quad (3)$$

To correct for ongoing analyte dilution the remaining fraction f has been calculated as follows

$$f = c_{xT}*c_{i0}/(c_{x0}*c_{iT}) \quad (4)$$

where c_{x0} and c_{xT} are the mixing ratios of CH_3Cl at time zero and time t and c_{iO} and c_{iT} are the respective concentrations of the internal standard PFH.

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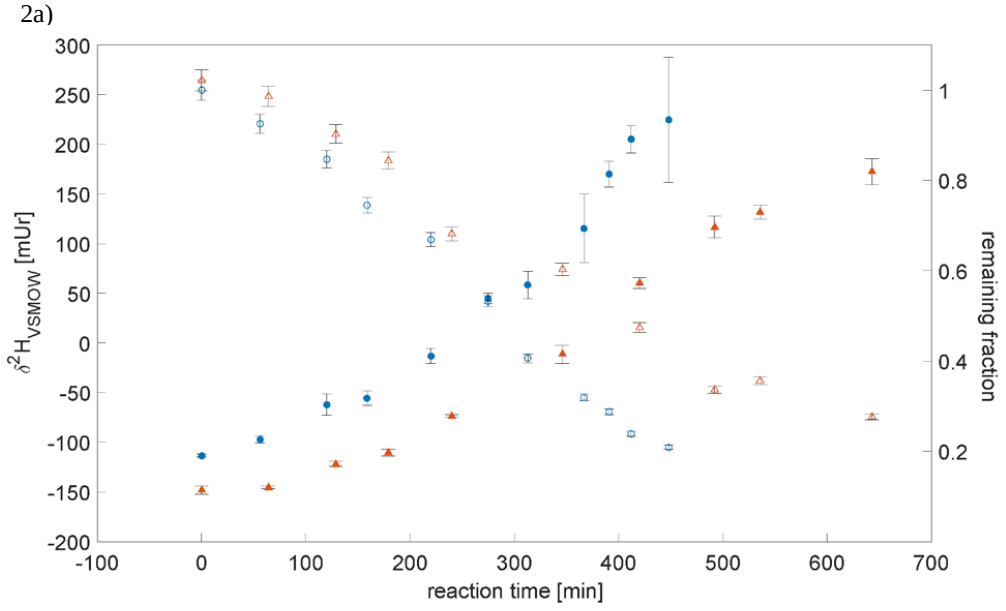
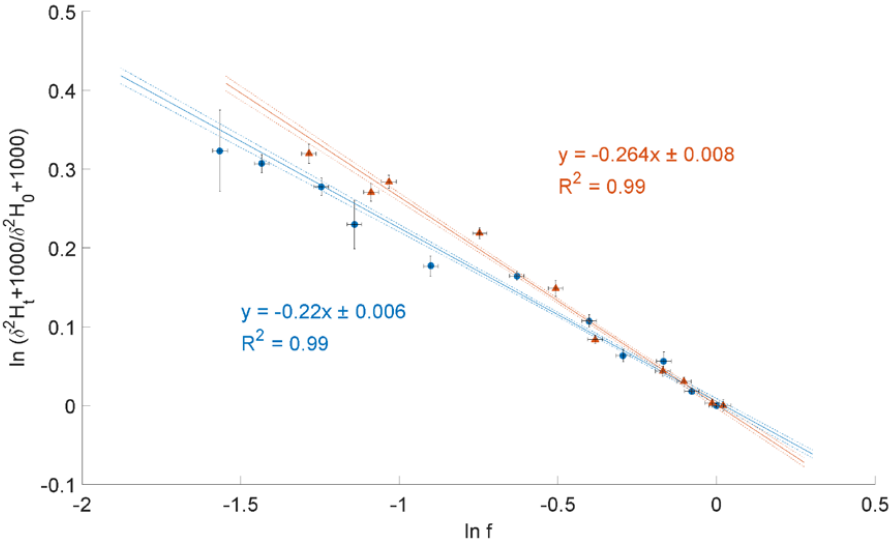
3 Results

The first experiment of CH_3Cl degradation with OH was performed on the 25/02/2014 and repeated under similar conditions on the 03/02/2015. Under the experimental conditions (see methods section) more than 70% of the CH_3Cl was degraded within 7 to 10 h. The results from these two experiments are shown in Figure 2. Both the trend of changes in δ^2H values of CH_3Cl as well as the remaining fraction of CH_3Cl observed in the two independent experiments are in good agreement (Figure 2a). The calculated ε values for experiments 1 and 2 are -264 ± 8 mUr and -220 ± 6 mUr, respectively (Figure 2b), with a correlation coefficient R^2 of the slope of the regression line of 0.99 for both experiments.

Figure 2: Reaction of CH_3Cl and OH. Two independent experiments (triangles and dots) were carried out using an initial mixing ratio of ~10 ppmv CH_3Cl. More than 70% of the CH_3Cl was degraded within 8 to 10 h. (a) Measured δ^2H values (filled circles and triangles) of CH_3Cl versus residual fraction (open circles and triangles) of CH_3Cl (calculated from changes of CH_3Cl and PFH). Error bars of δ^2H value of CH_3Cl indicate the standard deviation (SD) of the mean of three replicate measurements. Some error bars lie within the symbol. (b) Rayleigh plot (equation 2). Error bars were calculated by error propagation including uncertainties in δ^2H values of CH_3Cl and the remaining fraction. Dashed lines represent 95% confidence intervals of the linear regressions (bold lines).



2b)



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249 The CH_3Cl degradation with Cl experiment was conducted on the 18/02/2014. Here, over
250 90% of CH_3Cl was degraded during reaction with Cl radicals within 7 to 8 hours (Figure 3a).
251 The calculated ε of experiment 3 is -280 ± 11 mUr (Figure 3b) with a correlation coefficient
252 of the slope of the regression line of 0.99. Due to limited analytical resources it was not
253 possible to repeat this experiment.

254

255 **Figure 3:** Reaction of CH_3Cl and Cl. Initial mixing ratio of CH_3Cl was ~10 ppmv. More than
256 90% of the CH_3Cl was degraded within 7 to 8 h. (a) Measured δ^2H values (filled circles) of
257 CH_3Cl versus residual fraction (open diamonds) CH_3Cl. Error bars of δ^2H values of CH_3Cl
258 indicate the standard deviation (SD) of the mean of three replicate measurements. Some error
259 bars lie within the symbol. (b) Rayleigh plot (equation 2). Data are expressed as the
260 mean ± standard error of the mean, n = 3. Error bars were calculated by error propagation
261 including uncertainties in δ^2H values of CH_3Cl. Dashed lines represent 95% confidence
262 intervals of the linear regressions (bold line).

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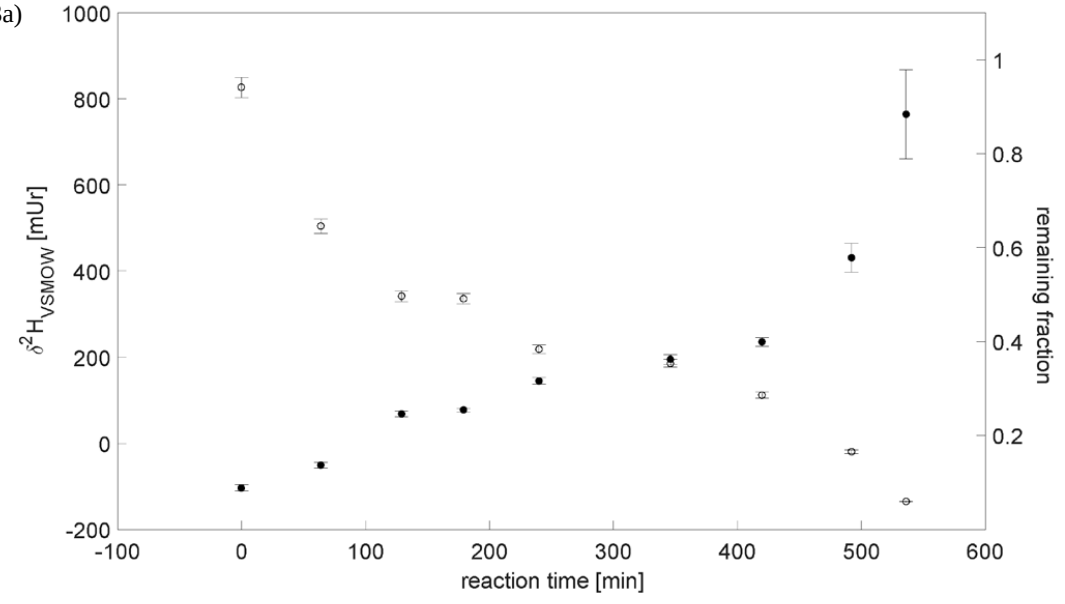
3a)



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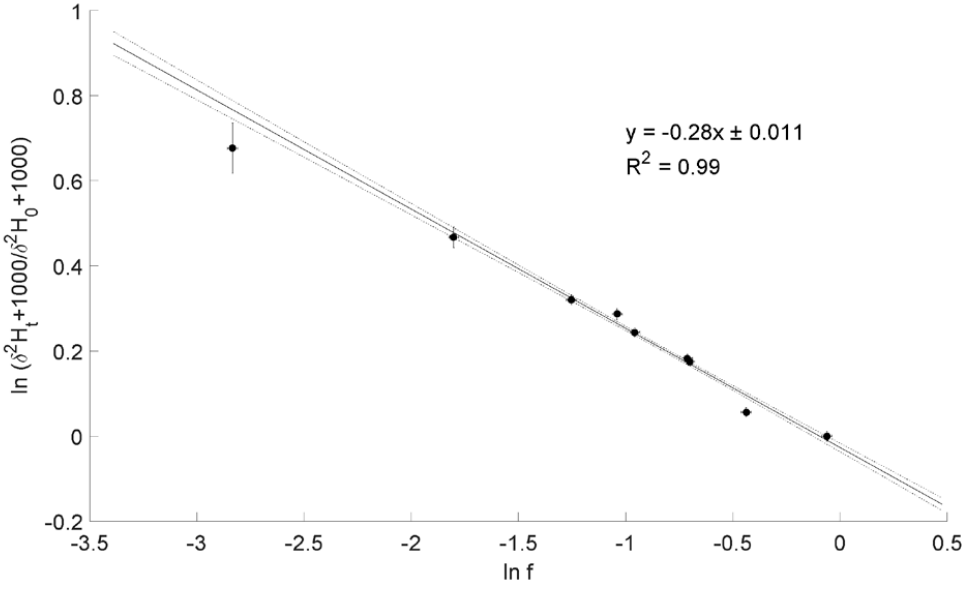
264 3b



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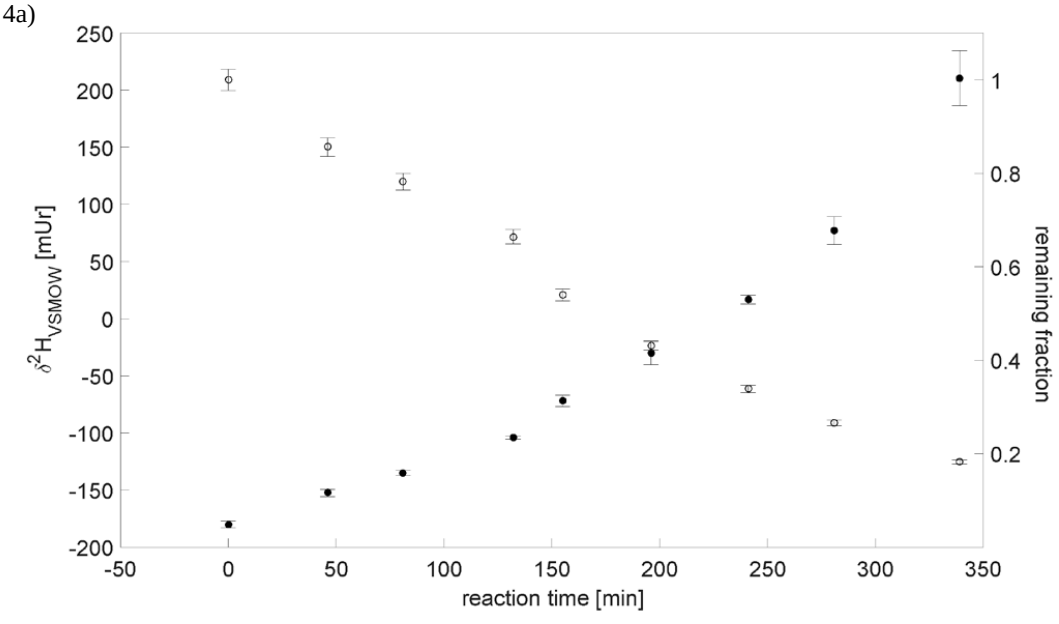
267

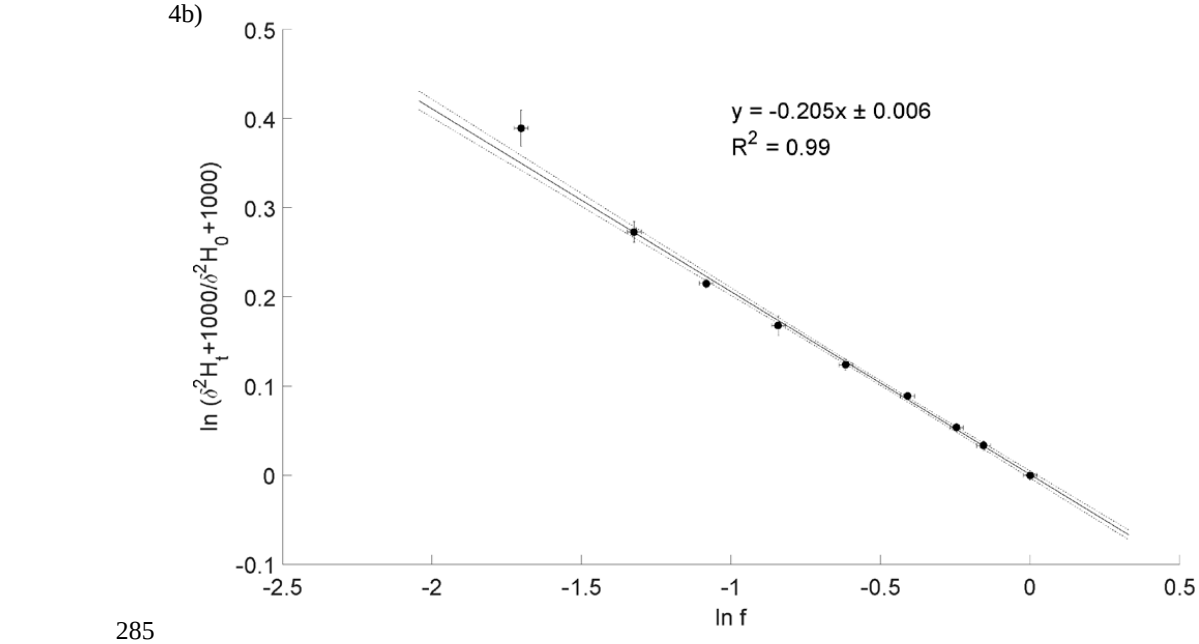
268 The experiment to determine the isotope enrichment constant of the degradation of CH_4 by
269 hydroxyl radicals was conducted on the 02/02/2015. Over 80% of CH_4 was degraded during

reaction with OH radicals within 7 hours (Figure 4a). The calculated ε of experiment 4 is -205 ± 6 mUr (Figure 4b) with a correlation coefficient of the slope of the regression line of 0.99.

Figure 4: Reaction of CH_4 and OH. Initial mixing ratio of CH_4 was ~6 ppmv. More than 80% of the CH_4 was degraded within 7 h. (a) Measured δ^2H values of CH_4 versus residual fraction of CH_4. Error bars of δ^2H values of CH_4 indicate the standard deviation (SD) of the mean of three replicate measurements. Some error bars lie within the symbol. (b) Rayleigh plot (equation 2). Error bars were calculated by error propagation including uncertainties in δ^2H values of CH_4 and the remaining fraction. Dashed lines represent 95% confidence intervals of the linear regressions (bold line).

4a)





4 Discussion

Chloromethane reacts with both hydroxyl and chlorine radicals in the atmosphere. The first degradation step of CH_3Cl in both reactions is the abstraction of a hydrogen atom to yield CH_2Cl and H_2O or HCl, respectively (Spence et al., 1976; Khalil and Rasmussen, 1999). In both reactions hydrogen is directly present in the reacting bond, and thus influenced by the so-called primary isotope effect (Elsner et al., 2005). Particularly for hydrogen these primary kinetic isotope effects are in general large as they involve a large change in relative mass of the atoms being abstracted. In the following we would like to discuss and compare our results with (i) previous work conducted by (Sellevåg et al., 2006), (ii) with OH degradation experiments of CH_4 and (iii) with the very recent report of biochemical degradation of CH_3Cl in soils and plants (Jaeger et al., 2018b; Jaeger et al., 2018a).

Although our experimental results show relatively large hydrogen isotope fractionations with ε values of -242 (mean result from two independent experiments) and -280 mUr, for reaction of CH_3Cl with OH and Cl radicals, respectively, they are smaller than the isotope fractionations previously measured and theoretically calculated by (Sellevåg et al., 2006) (Table 1). These researchers employed smog chamber experiments at 298 K and used FTIR measurements to determine the stable hydrogen isotope fractionation of CH_3Cl and reported ε values of -410 and -420 mUr for the reaction of CH_3Cl with OH and Cl radicals, respectively.

They also performed theoretical calculations of ε for the reactions of CH_2DCl with OH and Cl radicals and reported ε values in the range of -330 to -430 and -540 to -590 mUr, respectively (Table 1). Whilst we do not know the reasons for the large discrepancies in the experimental ε values observed here and those reported by Sellevåg et al. (2006), we suggest that they may be due to differences in the experimental smog chamber set-up or the different measurement techniques employed in each of the studies. However, we also conducted similar smog chamber experiments for the degradation of CH_4 with hydroxyl radicals (see methods section and Figure 4) and calculated an ε value of -205 ± 6 mUr for the reaction of CH_4 with OH radicals at a temperature of 293 ± 1 K. In Table 1 we compare our results with those from a number of previous studies (Saueressig et al., 2001; Sellevåg et al., 2006; DeMore, 1993; Gierczak et al., 1997; Xiao et al., 1993), which were conducted at temperatures ranging from 277 to 298 K (Table 1). The ε values for the reaction of CH_4 with OH radicals from all studies ranged from -145 to -294 mUr with a mean value of -229 ± 44 mUr with the most negative ε value of -294 ± 18 mUr reported by Sellavag and coworkers (2006). The ε value found in this study (-205 ± 6 mUr) was in good agreement with previous experimentally reported values conducted at similar temperatures. This finding gave us confidence that our experimental design and the measurements made using GC-IRMS were reliable.

Compared to primary isotope effects, changes in bonding are much smaller in the case of secondary isotope effects, where positions adjacent to the reacting bond are only slightly affected by the proximity to the reaction centre (Elsner et al., 2005; Kirsch, 1977). It was suggested that for the same element, secondary isotope effects are generally at least 1 order of magnitude smaller than primary isotope effects (Kirsch, 1977; Westaway, 1987; Merrigan et al., 1999).

We therefore compared our results from chemical degradation experiments with those from recently reported biochemical degradation experiments (Jaeger et al., 2018a; Jaeger et al., 2018b). So far, the only known pathway for biochemical consumption of CH_3Cl is corrinoid- and tetrahydrofolate-dependent and is termed cmu (abbreviation for chloromethane utilization). This pathway was characterized in detail for the aerobic facultative methylotrophic strain *Methylobacterium extorquens* CM4 (Vannelli et al., 1999) and involves genes that were also detected in several other chloromethane-degrading strains (Schafer et al., 2007; Nadalig et al., 2013; Nadalig et al., 2011). During degradation of CH_3Cl the methyl group is transferred to a corrinoid cofactor by the protein CmuA. In this case the carbon-

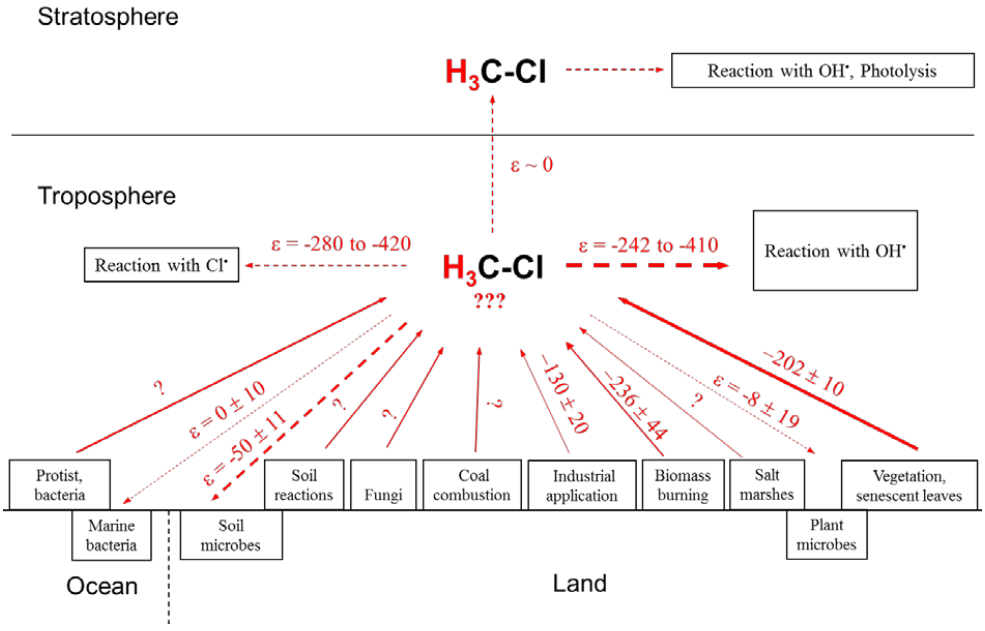
chlorine bond of CH_3Cl is broken and thus since the hydrogen atoms are adjacent to the reacting bond only a secondary isotope effect would be expected. Indeed, the first ε values reported (Jaeger et al., 2018a; Jaeger et al., 2018b) for CH_3Cl biodegradation by different soils and plants (ferns) are in the range of -50 ± 13 mUr and -8 ± 19 mUr, respectively, and thus showing considerably smaller kinetic isotope effects than for chemical degradation of CH_3Cl by OH and Cl radicals measured in either this study or reported by Sellevåg et al. (2006).

5 Conclusions and future perspectives

We have performed experiments to measure the hydrogen isotope fractionation of the remaining unreacted CH_3Cl following its degradation by hydroxyl and chlorine radicals in a 3.5 m^3 Teflon smog-chamber at 293 ± 1K. δ^2H values of CH_3Cl were measured using GC-IRMS. The calculated isotope fractionations of CH_3Cl for the reactions with hydroxyl and with chlorine radicals were found to be smaller than either the experimentally measured (by FTIR) or theoretical values reported by Sellevåg et al. (2006). We also performed degradation experiments of CH_4 using the same smog-chamber facilities yielding an isotope enrichment constant for the reaction of CH_4 with hydroxyl radicals of -205 ± 6 mUr which is in excellent agreement with previous reported results. Although stable hydrogen isotope measurements of CH_3Cl sources are still scarce, some recent studies have reported first data on δ^2H values of CH_3Cl sources and ε values on sinks (Greule et al., 2012; Jaeger et al., 2018a; Jaeger et al., 2018b; Nadalig et al., 2014; Nadalig et al., 2013).

We have summarized all available information regarding δ^2H values of environmental CH_3Cl sources in Table 2. Furthermore, the known CH_3Cl sinks and their associated isotope enrichment constants are presented in Table 3. Eventually Figure 5 displays the global CH_3Cl budget showing the known hydrogen isotope signatures of sources and isotope enrichment constants associated with sinks.

Figure 5. Scheme of major sources and sinks involved in the global CH_3Cl cycle (modified after Keppler et al., 2005) with known (experimentally determined) corresponding δ^2H values and isotope enrichment constants, respectively. Red straight and dashed lines of arrows indicate sources and sinks of CH_3Cl, respectively. Questions marks indicate where currently no data exist. All values are given in mUr.



Our results suggest that stable hydrogen isotope measurements of both sources and sinks of CH_3Cl and particularly the observed large kinetic isotope effect of the atmospheric CH_3Cl sinks might strongly assist with the refinement of current models of the global atmospheric CH_3Cl budget. In contrast to the large hydrogen fractionation of CH_3Cl by chemical degradation of OH and Cl radicals, the isotope fractionation of CH_3Cl biodegradation are in the range of an order of magnitude lower. This therefore holds the opportunity to improve our understanding of the global CH_3Cl budget once the δ^2H value of atmospheric CH_3Cl has been measured. The stable hydrogen isotopic composition of tropospheric CH_3Cl depends on the isotopic source signatures and the kinetic isotope effects of the sinks, primarily the reaction with OH and consumption by soils and potentially plants.

Several attempts at modelling the global CH_3Cl budget using stable carbon isotope ratios have already been made (Harper et al., 2001; Harper et al., 2003; Thompson et al., 2002; Keppler et al., 2005; Saito and Yokouchi, 2008) but there are still major uncertainties regarding source and sink strengths as well as the respective stable isotope signatures. Therefore, we now suggest combining our knowledge of stable carbon and hydrogen isotopes of CH_3Cl in the environment. Such a two dimensional (2D) stable isotope approach of hydrogen and carbon can be used to better understand the processes of CH_3Cl biodegradation and formation. Furthermore, when this approach is combined with CH_3Cl flux estimates it

could help to better constrain the strength of CH_3Cl sinks and sources within the global CH_3Cl budget (Nadalig et al., 2014; Jaeger et al. 2018b)

We would highlight that currently no data is available for the δ^2H value of atmospheric CH_3Cl. Although it will be a massive analytical challenge to obtain this value, we strongly consider that it would likely lead to a better refined isotopic mass balance for atmospheric CH_3Cl and thus to our better understanding of the global CH_3Cl budget.

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562 Table 1: Reported hydrogen isotope enrichment constants for the reaction of CH_3Cl with OH
563 radicals and with Cl atoms and the reaction of CH_4 with OH radicals.

Reaction	**ε / mUr**	**Method and remarks**	**Reference**
CH_3Cl + OH	-242 mean	experimental: 3.5 m^3 smog-chamber at 293 ± 1 K; IRMS	Exp. 1 & 2, this study
CH_3Cl + OH	-410 ± 50	experimental: smog-chamber, long-path FTIR spectroscopy relative to CH_3Cl at 298 ± 2 K	Sellevåg et al. 2006
CH_3Cl + OH	-330 to -430	theoretical calculations	Sellavåg et al. 2006
CH_3Cl + Cl	-280 ± 11	experimental: 3.5 m^3 smog-chamber at 293 ± 1 K; IRMS	Exp. 3, this study
CH_3Cl + Cl	-420 ± 40	experimental: smog-chamber, long-path FTIR spectroscopy relative to CH_3Cl at 298 ± 2 K	Sellavåg et al. 2006
CH_3Cl + Cl	-540 to -590	theoretical calculations	Sellavåg et al. 2006
CH_4 + OH	-205 ± 6	experimental: 3.5 m^3 smog-chamber at 293 ± 1 K; IRMS	Exp. 4, this study
CH_4 + OH	-227 ± 11	experimental: at 296 K, IRMS and tunable diode laser absorption spectroscopy	Saueressig et al. 2001
CH_4 + OH	-231 ± 45	experimental: at 277 K	Gierczak et al., 1997
CH_4 + OH	-251 ± 10	ab initio at 298 K	Xiao et al., 1993
CH_4 + OH	-145 ± 30	experimental: at 298 K	DeMore et al., 1993
CH_4 + OH	-294 ± 18	experimental: smog-chamber, long-path FTIR spectroscopy relative to CH_3Cl at 298 ± 2 K	Sellavåg et al. 2006
CH_4 + OH	-60 to -270	theoretical at 298 K	Sellavåg et al. 2006

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Table 2. Known sources of tropospheric CH_3Cl and corresponding δ^2H values.

Sources	Source (best estimate)[a] (Gg yr^{-1})	Source (full range)[a] (Gg yr^{-1})	Mean δ^2H value mUr vs VSMOV	Uncertainty δ^2H value ± mUr
Open field biomass burning	355	142 to 569	-236[b]	44
Biomass burning indoor	113	56 to 169	-236[b]	44
Tropical and subtropical plants	2040	1430 to 2650	–202[c]	10
Fungi	145	128 to 162	?	
Salt marshes	85	1.1 to 170	?	
Coal combustion	162	29 to 295	?	
Industrial chemical production[d]	363	278 to 448	–130[e]	20
Oceans	700	510 to 910	?	
Others[f]	~58	27 to 86	?	
Total sources	3658 (4021)	2601 to 5459		

[a] Values for source (best estimate) and source (full range) were taken from Carpenter and Reimann (2014), except for emissions associated with chemical production by the industry which are from Li et al. (2016). Value shown for total sources in brackets includes chemical production by the industry.

[b] Greule et al. 2012; please note that all values provided for CH_3Cl released from dried plants at elevated temperatures have been corrected by -23 mUr due to recalibration of the reference gas.

[c] Jaeger et al. (2018b)

[d] Li et al. (2016)

[e] taken from Greule et al. (2012), Nadalig et al. (2013) and Jaeger et al. (2018a & 2018b); please note that values provided by Greule et al. (2012) and Nadalig et al. (2013) for CH_3Cl from sources of the chemical industry have been corrected by -23 mUr due to recalibration of the reference gas

[f] including mangroves, wetlands, rice paddies and shrublands

? denotes that no value has been provided

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Table 3. Known sinks of tropospheric CH_3Cl and the mean isotope enrichment constant ε reported for each.

Sinks	Sink (best estimate)[a] (Gg yr^{-1})	Sink (full range)[a] (Gg yr^{-1})	Isotope enrichment constant ε / mUr	Uncertainty ε ± mUr
Reaction with OH in troposphere	2832	2470 to 3420	-242[b] -410[c]	7[b] 50[c]
Loss to stratosphere	146	?	0[c]	?
Reaction with Cl in marine boundary layer	370[d]	180 to 550[d]	-280[b] -420[c]	11[b] 40[c]
Microbial degradation in soil	1058	664 to 1482	-50[e]	13[e]
Loss in ocean	370	296 to 445	0[f]	10[f]
Microbial degradation in plants[g]	?	?	-8[g]	19[g]
Total sinks	4406 (4776)			

[a] Values for sink strength (best estimate and full range) were taken from Carpenter and Reimann (2014), except for the value of the reaction with Cl-radicals in marine boundary layer and for total sinks shown in brackets which includes the potential sink strength by Cl-radicals in marine boundary layer (Montzka and Fraser, 2003).

[b] this study, mean value of two experiments

[c] Sellevåg et al. (2006)

[d] Thompson et al. (2002) and discussion in this manuscript

[e] Jaeger et al. (2018a)

[f] Nadalig et al. (2014)

[f] Jaeger et al. (2018b)

? denotes that no value has been provided

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