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L. J. Carpenter, D. J. Wevill, S. O'Doherty, G. Spain, P. G. Simmonds. Atmospheric bromoform at Mace Head, Ireland: Evidence for a peatland source. *Atmospheric Chemistry and Physics Discussions*, European Geosciences Union, 2005, 5 (4), pp.5935-5955. hal-00301694

HAL Id: hal-00301694

<https://hal.archives-ouvertes.fr/hal-00301694>

Submitted on 15 Aug 2005

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Atmospheric bromoform at Mace Head, Ireland: Evidence for a peatland source

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Received: 21 June 2005 – Accepted: 11 July 2005 – Published: 15 August 2005

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In situ atmospheric observations of bromoform (CHBr_3) made over a 2.5 year period at Mace Head, Ireland from May 2001–December 2003, including during the NAMBLEX (North Atlantic Marine Boundary Layer Experiment) campaign, show broad maxima from spring until autumn and winter minima, with mixing ratios of 5.3 ± 1.0 pptv (mid March–mid October) and 1.8 ± 0.8 pptv (December–February). This indicates that, unlike CHCl_3 , which has a summer minimum and winter maximum at Mace Head, local biological sources of CHBr_3 have a greater influence on the atmospheric data than photochemical decay during long-range transport. The emission sources are predominantly macroalgal, but we find evidence for a small terrestrial flux from peatland ecosystems, which so far has not been accounted for in the CHBr_3 budget. Sharp increases in CHCl_3 and CHBr_3 concentrations and decreases in O_3 concentrations occurred at night when the wind direction switched from an ocean- to a land-based sector (land breeze) and the wind speed dropped to below 5 ms^{-1} . These observations infer a shallow atmospheric boundary layer with increased O_3 deposition and concentration of local emissions of both CHCl_3 and CHBr_3 . The ratio of $\Delta\text{CHCl}_3/\Delta\text{CHBr}_3$ varied strongly according to the prevailing wind direction; from 0.6 ± 0.1 in south-easterly ($100\text{--}170^\circ$) air to 1.9 ± 0.8 in north-easterly ($40\text{--}70^\circ$) air. Of these land-sectors, the south-easterly air masses are likely to be strongly influenced by macroalgal beds along the coast and the emission ratios probably reflect those from seaweeds in addition to land sources. The north-easterly airmasses however have a fetch predominantly over land, which locally is comprised of coastal peatland ecosystems (peat bogs and coastal conifer plantations), previously identified as being strong sources of atmospheric CHCl_3 under these conditions. Although we cannot entirely rule out other local land or coastal sources, our observations also suggest peatland ecosystem emissions of CHBr_3 . We use correlations between CHCl_3 and CHBr_3 during the land breeze events in conjunction with previous estimates of local wetland CHCl_3 release to tentatively deduce a global wetland CHBr_3 source of 26.9 ($0.5\text{--}1247$) Gg yr^{-1} , which is approximately 10% of the total

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1. Introduction

Bromoform is ubiquitous in the marine environment and is the major natural carrier of organic bromine to the atmosphere. It has a relatively short tropospheric lifetime of ~ 3 weeks and is believed to make a significant contribution to inorganic bromine in the upper troposphere and lower stratosphere (Sturges et al., 2000; Nielsen and Douglass, 2001; von Glasow et al., 2004). The major sources of CHBr_3 have been identified as marine in origin, namely macroalgae, ocean phytoplankton and water chlorination (Gschwend et al., 1985; Carpenter and Liss, 2000; Quack and Wallace, 2003 and references therein). To date, there are only limited and inconclusive studies of production by terrestrial ecosystems. CHBr_3 has been found in the air of upper soil of a pristine spruce forest (Haselmann et al., 2000). However, in soil air of rural areas, Hoekstra et al. (1998) found no detectable concentrations of CHBr_3 , except after in-situ enrichment by KBr . This suggests the potential of soils containing elevated bromide, e.g. coastal soils/peatlands, to produce brominated trihalomethanes. Peat bogs cover approximately 3% of the total continental land mass. They are comprised exclusively of organic matter and in additional coastal bogs are subject to deposition of halides from sea spray, making such sites potentially significant organohalogen producers. Biester et al. (2004) found that the majority of halogens in peat exist in an organically bound form, with concentrations dependent upon peat decomposition processes. Speciated organohalogens were however not identified in this study.

Chloroform concentrations at Mace Head can be substantially influenced by local biological emissions from the pine forests and peat bogs, which cover large parts of Ireland (Dimmer et al., 2001; O'Doherty et al., 2001). Extrapolating from chamber studies made on Irish peatland ecosystems, Dimmer et al. (2001) estimated global annual fluxes of 4.7 (0.1–151.9) Gg yr^{-1} of CHCl_3 from such systems and 24.1 Gg yr^{-1} from total wetlands. CHBr_3 measurements were not made in the latter study. Here, we

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present over 2.5 years of CHBr_3 data measured at Mace Head and use the tracer-ratio technique, in conjunction with the data from Dimmer et al., to estimate the contribution of peatland and wetlands to global CHBr_3 emissions.

2. Experimental

2.1. Site description

Mace Head is located on the remote western coast of County Galway, Ireland ($53^\circ 19' \text{ N}$, $9^\circ 54' \text{ W}$). The site is well known for background air measurements and receives relatively clean marine air from the prevailing westerly sector associated with the easterly tracking cyclonic systems of the North Atlantic. The surrounding area is sparsely populated and contains a large expanse of peat bogs. The nearest large population centre is Galway, ~ 62 miles to the east. Read et al. (2005)¹ give a full description of the site.

2.2. Instrumentation

The York halocarbon measurements were made during the NAMBLEX campaign (2 August–3 September 2002) with a Perkin Elmer Turbomass GC-MS system with electron impact ionisation and operating in selective ion recording mode. Air sampling was from a height of 20 m through a stainless steel $3/4''$ manifold, pumped at $\sim 30 \text{ L min}^{-1}$. Sample volumes of 3 L were pre-concentrated on a two stage carbon-based adsorbent trap held at -10°C before being transferred to the GC by rapid heating to 400°C . Halocarbons were separated on a SGE BPX5 capillary column ($50 \text{ m} \times 0.32 \text{ mm i.d.} \times 3 \mu \text{m d.f.}$). Detection limits are between 0.02 and 0.12 pptv for approximately hourly samples of CHCl_3 , CH_3I , $\text{C}_2\text{H}_5\text{I}$, $1\text{-C}_3\text{H}_7\text{I}$, $2\text{-C}_3\text{H}_7\text{I}$, CH_2ClI , CH_2Br_2 ,

¹Read, K. A., Heard, E., Methven, J., and The NAMBLEX team: The North Atlantic Marine Boundary Layer Experiment (NAMBLEX). Overview of the campaign held at Mace Head, Ireland, in summer 2002, Atmos. Chem. Phys. Discuss., submitted, 2005.

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CHBr₂Cl, CH₂Brl, CHBr₃ and CH₂I₂ with a precision of 3–8% (5% for CHBr₃) and complete recovery of all compounds. Our calibration system utilises fixed volume (50 μl) loop injections of the output of thermostatted in-house permeation tubes filled with pure liquids (≥97% purity, Aldrich, UK) into a 100 ml min⁻¹ stream of purified nitrogen gas in order to dilute parts per million by volume (ppmv) mixing ratios into low parts per trillion by volume (pptv). The calibration is completely automated, allowing multi-point calibrations during routine operation. The overall accuracy of the measurements is estimated to be ±15%. A full description of the calibration and GC-MS methodology is given in Wevill and Carpenter (2004).

Routine AGAGE (Advanced Global Atmospheric Gases Experiment) GC-MS observations, made by Bristol University, of a range of halocarbons commenced at Mace Head in 1994. A full description of the instrumentation can be found in Prinn et al. (2000) and Simmonds et al. (1995). A nominal, precisely repeatable two liters of ambient air or calibration standard are trapped onto a 3-stage mixed adsorbent contained in a microtrap. The microtrap is located between two thermoelectric coolers maintained at -50°C during sampling. This low temperature enables the small mass of adsorbent material to have the capacity to quantitatively trap the compounds of interest. The trapped sample is purged briefly with clean helium then thermally desorbed (245°C) directly onto a chromatographic capillary column (CPSil-5, 100 m×0.32 mm i.d. ×5 μm d.f.), interfaced to an Agilent benchtop MS. The individual trace components are separated and identified using both their chromatographic retention times, and their characteristic ion masses. Each component is quantified by integrating the response of a selected target ion and where possible, two qualifier ions are used to ensure compound specific ion ratios are maintained. Custom software developed at Scripps Institution of Oceanography (SIO), enables constant monitoring of ion ratios, peak width ratios and many other diagnostic parameters. This enables rapid identification and determination of instrumental problems. At present standard and air analysis are alternated every two hours to provide six calibrated measurements per day. The method of standard preparation and propagation is described in detail in O'Doherty et al. (2004). Briefly,

all working standards are 35l stainless steel electropolished tanks (Essex Cryogenics, Missouri, USA) filled with either Mace Head “real-air”, under baseline conditions, and have a working life of 3–4 months. Calibration of ambient air measurements is achieved by means of bracketing them with measurements from a working standard tank. At present we do not assign a CHBr_3 calibration for the ADS-GC-MS data at Mace Head.

2.3. Comparison of York/Bristol measurements and conversion of Bristol CHBr_3 relative response to pptv values

An “organic” x-y correlation (i.e. assuming equal errors in x and y data) of York/Bristol CHCl_3 data taken within the same 40-min period during the NAMBLEX campaign showed that on average the York measurements were 13% higher than the AGAGE data (with zero offset), with an R^2 value of 0.95 (Fig. 1a). The fact that the instruments were not sampling for the same time periods may explain some or most of the scatter, however there is clearly also a calibration difference although this is within the estimated error of our measurements.

The same procedure for the correlation of York CHBr_3 mixing ratios against the Bristol CHBr_3 relative response to a Mace Head reference air sample contained in a 36L electropolished Essex canister) revealed more variance in the linear regression (R^2 value of 0.74) and an offset in the Bristol data (Fig. 1b). Because the York instrument has the same precision for CHBr_3 and CHCl_3 measurements (Wevill and Carpenter, 2004), we attribute the increased scatter of the correlation compared to Fig. 1a to the shorter lifetime and hence greater natural variability of atmospheric CHBr_3 . The offset of the Bristol data may be caused by incomplete trapping or recovery of high concentrations of CHBr_3 in the microtrap, which was designed for quantitative measurements of the more volatile Halons and CFCs rather than CHBr_3 . A comparison of the York data with Bristol CHBr_3 mixing ratios converted using both the organic regression values and a standard linear least squares regression (with York data on the x axis) is shown in Fig. 2 and reveals the higher dynamic range of the York data. Although the Bristol

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instrument appears to underestimate the highest mixing ratios of CHBr_3 , nevertheless the average concentrations derived using this approach are equal to the York average concentrations. We chose to use a standard correlation to “correct” the Bristol data to mixing ratios because the organic correlation under-predicted low values and in fact

5 resulted in occasional negative mixing ratios in winter.

2.4. Data selection for local land breeze events

Land breeze events, where the wind direction changes from westerly (ocean sector) in the day to easterly (land sector) at night, are quite commonly observed at Mace Head (e.g. O’Doherty et al., 2001; Biraud et al., 2002). They are accompanied by a

10 rapid decrease in O_3 , due to increased deposition over land, and increase in CHCl_3 , CO_2 and ^{212}Pb concentrations during low wind speeds due to local emissions (Biraud et al., 2002). Here, we select events within the land sector ($20\text{--}140^\circ$) with a $>$ than 40% maximum decrease in $[\text{O}_3]$ and $>100\%$ change in $[\text{CHCl}_3]$. We define the fetch of the land breeze events as the product of their time duration (typically 5–7 h) and the

15 average wind speed recorded during the event (typically $1\text{--}5\text{ ms}^{-1}$) and select events with a fetch of $<50\text{ km}$, i.e. dominated by local continental emissions.

2.5. Tracer-ratio approach

The tracer-ratio technique has been used, in various forms, to estimate trace gas emissions by utilising correlations obtained with a reference compound with a known emission, in this case CHCl_3 (e.g. Dunse et al., 2001; Biraud et al., 2002; Carpenter et al.,

20 2003). Visual inspection of the data from Mace Head clearly showed the strong covariance of CHCl_3 , CHBr_3 and O_3 concentrations during land breeze events. For each event, the maximum change in concentration ΔX (where $X=\text{CHCl}_3$ or CHBr_3) was calculated (the maxima in both species were always coincident temporally), both in units

25 of pptv, yielding one data point for each event. We then use the simple approach of

assuming a linear relationship between the enhancements in mixing ratio,

$$\Delta\text{CHCl}_3/\Delta\text{CHBr}_3 = E \quad (1)$$

where E is the emission ratio, deduced from the slope of the x-y graph. Here we omit the intercept term, a function of the relative initial concentrations of the trace gases (e.g. Carpenter et al., 2003). The overall approach assumes no atmospheric decay along the trajectory, which in this case should apply, given the short fetches.

3. Results and discussion

3.1. General trends and seasonality

The complete Bristol data set, fixed to the York calibration scale as described in Sect. 2.3, is shown in Fig. 3. Broad spring-autumn maxima, with occasional peaks of over 25 pptv but an average of ~5 pptv, and winter minima are apparent. Figure 4 shows an averaged annual cycle for CHBr₃, comprising 2-weekly average data points, with mixing ratios of 5.3±1.0 pptv (mid March–mid October) and 1.8±0.8 pptv (December–February). Also shown is an average UVB (290–320 nm) annual cycle computed for Mace Head. The only other published annual measurements of CHBr₃ have been made at the free tropospheric site of Mauna Loa (Atlas and Ridley, 1996, discussed in Nielsen and Douglass, 2001) and at Alert, in the Arctic boundary layer (Yokouchi et al., 1996). Both these studies showed winter maxima and summer minima, suggesting higher reactive loss in summer during long-range transport, but could give little information on the seasonal variation of sources. Thus our data gives a unique insight into the seasonal variation of coastal CHBr₃ emissions.

CHBr₃ appears to be the most abundant organic halogen compound emitted from macroalgae (e.g. Nightingale et al., 1995; Mtolera et al., 1996; Carpenter et al., 2000) and these emissions are significant on a global scale, comprising probably the dominant source of atmospheric CHBr₃ (Carpenter and Liss, 2000). Halocarbon formation

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in macroalgae is initiated by oxidative stress, resulting in H_2O_2 production as part of a defense mechanism (Theiler et al., 1978; Pedersén et al., 1996 and references therein). The formation of polyhalomethanes are initiated by haloperoxidase enzymes produced in the cells, which catalyse the reaction of accumulated halide ions with H_2O_2 , forming a hypohalous acid. Halogenated species such as $CHBr_3$ may be subsequently formed in the algal cells and transferred across the cell membrane to seawater (Gschwend et al., 1985). Alternatively, formation may be initiated outside the cell by reaction of the released hypohalous acid with organic matter in seawater (Wever et al., 1991).

Macroalgae exhibit quite constant cover over the course of the year, resulting from the continual recruitment of young plants and the prolific output of spores. Our data indicate that $CHBr_3$ emission occurs all year, with only a short winter period of minimal levels (November–February), when the algal population of the intertidal region is at its lowest. Goodwin et al. (1997) and Klick (1993) reported that $CHBr_3$ release from brown macroalgae maximises in mid summer, probably due to enhanced tissue decay during this period. Further, a number of studies have shown that $CHBr_3$ release from macroalgae is stimulated by light (e.g. Klick, 1993; Nightingale et al., 1995; Mtolera et al., 1996; Goodwin et al., 1997; Carpenter et al., 2000). Light-induced stress, especially at high light intensities, results in increased cellular levels of activated oxygen species including hydrogen peroxide, as by-products of oxygenic photosynthesis, which generally increase the emission of halocarbons. Seaweed incubation experiments reveal an enhancement in $CHBr_3$ release by factors of 2–10 in daylight compared to the dark. Thus, variation in light intensities and/or tissue decay, in conjunction with higher algal coverage from early spring through to late autumn, could explain the seasonal variation of $CHBr_3$ at Mace Head.

It is well established that trace gas emissions from phytoplankton tend to peak in spring (at this latitude around early May), however the contribution of temperate microalgae to atmospheric $CHBr_3$ levels at Mace Head is probably minimal (Carpenter et al., 2003). The highest $CHBr_3$ concentrations are consistently found along coastlines containing macroalgal beds, with coastal and shelf seawater concentrations of $CHBr_3$

of up to 100 times greater than open ocean concentrations (Quack and Wallace, 2003).

3.2. Land breeze events

In total, 28 land breeze events as defined by the selection criteria described in Sect. 2.4 were identified during mid 2001–2003, all occurring during the period 23:00–04:00.

5 These events were characterised by sharp changes in wind direction (from sea to land), [O₃], [CHCl₃] and [CHBr₃], as typified by the event shown in Fig. 5. Only on the 29/30 August 2002 (during the NAMBLEX campaign) were sufficient Bristol and York data available for comparison; the excellent agreement of the (corrected) Bristol and York $\Delta\text{CHCl}_3/\Delta\text{CHBr}_3$ ratios of 0.62 and 0.61, respectively gives some confidence that both
10 data sets can be used to give an estimate of the emission ratio E in Eq. (1).

Figure 6 shows the ΔCHCl_3 vs. ΔCHBr_3 correlations, separated into events occurring in either south-easterly (100–170°) or north-easterly (40–70°) winds. South-easterly winds travel across the Irish shoreline and are expected to be strongly influenced by local macroalgal emissions. During such events, the CHCl₃/CHBr₃ ratio was relatively
15 invariant at ~0.6 ($R^2=0.73$), indicating that average emission ratios of CHCl₃/CHBr₃ from the local coastline are also relatively constant. Both CHCl₃ and CHBr₃ are known products of macroalgae (e.g. Nightingale et al., 1995; Carpenter et al., 2000 and references therein), with CHBr₃ consistently produced in higher amounts across many different genera.

20 In north-easterly winds, with a fetch inland, the CHCl₃/CHBr₃ ratio was much higher at 1.9 and exhibited much greater variance ($R^2=0.17$ and relative standard deviation (RSD) of 42%). The large variance in the regression may be due to a number of factors including different starting concentrations of CHCl₃ and CHBr₃ along the trajectories during the various events, and different emission ratios from local terrestrial
25 sources, e.g. peat bogs, inland marshes and coastal forests. Although CHBr₃ was not measured, Dimmer et al. (2001) found strikingly different emission ratios of CHCl₃ to CH₃Br in peatland compared to conifer forest in the vicinity of Mace Head, with greater

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variation still within vegetation types. Thus, although our measurements show strong evidence of peatland ecosystem sources of CHBr_3 at Mace Head, an absolute emission ratio compared to CHCl_3 should be treated with caution, not only because of variation within source, but also because of the short atmospheric lifetimes of CHCl_3 and CHBr_3 , and the instrumental issues potentially affecting the Bristol ratios, as discussed in Sect. 2.3. Nevertheless, it is instructive to approximate source strengths from peatland emissions in order to compare with other known source strengths (which are also highly uncertain). Dimmer et al. (2001) estimated a global CHCl_3 flux from peatland soil of 4.7 (0.1 – 151.9) Gg yr^{-1} and from total wetlands of 24.1 Gg yr^{-1} , calculated simply by extrapolating the measured Irish fluxes to estimated global coverages; 1.04×10^6 km^2 for peatbog (Olson, 1992) and 5.3×10^6 km^2 for wetlands (Matthews and Fung, 1987). We use the 1σ limits of the $\text{CHCl}_3/\text{CHBr}_3$ ratio determined here of 1.9 (1.1 – 2.7), together with CHCl_3 source estimates to deduce a total global CHBr_3 peatland source of 5.2 (0.1 – 241) Gg yr^{-1} and wetlands source of 26.9 (0.5 – 1247) Gg yr^{-1} . Any non-peatland ecosystem sources of either CHCl_3 or CHBr_3 over the north-easterly fetch will add error to our peatland emission estimates. However, despite the large uncertainties involved, it is worth comparing our peatland and wetland source estimates with current estimates for CHBr_3 emission of 220 Gg yr^{-1} from macroalgae (Carpenter and Liss, 2000) and a total global source of 300 Gg yr^{-1} (Dvortsov et al., 1999). Thus, it is possible that peatland and wetland emissions of CHBr_3 together are a small ($\sim 10\%$) but significant unaccounted term in the CHBr_3 budget.

4. Conclusions

So far, global modelling studies of tropospheric and stratospheric bromine that have specifically included CHBr_3 (Dvortsov et al., 1999; Nielsen and Douglass, 2001; von Glasow et al., 2004) have assumed a seasonally invariant ocean source of CHBr_3 . Our study indicates a much stronger source of CHBr_3 in summer than in winter, possibly linked to light levels and macroalgal coverage, and points to an additional coastal ter-

restrial source in the CHBr_3 budget of (tentatively) around 10% of the global source. The dual roles of such ecosystems in conjunction with macroalgae indicate the dominance of coastal regions in providing CHBr_3 to the atmosphere. Changes in wetland emissions, through man's impact on land use, and in ultraviolet radiation at the earth's surface due to the effect of ozone-depleting substances on stratospheric ozone, show the possibility of anthropogenic effects on the release of CHBr_3 , which itself is an ozone-depleting substance.

Acknowledgements. We specifically acknowledge the cooperation and efforts of the station operators and support staff: G. Spain and D. Brown at Mace Head. We also thank the Physics Department, National University of Ireland, Galway, for making the research facilities and meteorological data at Mace Head available. The operation of the Mace Head station was supported by the National Aeronautic and Space Administration (NASA) (grants NAGW-732, NAG1-1805, NAG5-3974 NAG-12099, to MIT; grants NAGW-2034, NAG5-4023 to SIO), the Department of the Environment, Food and Rural Affairs (UK) (contracts PECD 1/1/130 and 7/10/154, EPG 1/1/82 and EPG 1/1/130 to International Science Consultants). The halocarbon component of the NAMBLEX campaign was funded by NERC grant NER/A/S/2001/01064. D. J. Wevill thanks the EPSRC for his studentship.

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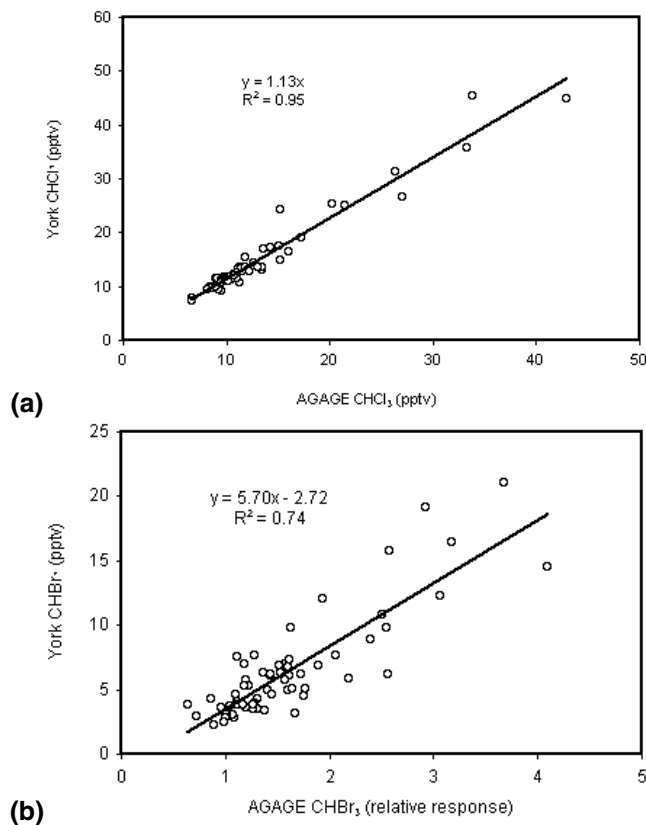


Fig. 1. (a) Organic correlation of York and Bristol CHCl_3 mixing ratios measured during the NAMBLEX campaign, using data points taken within the same 40-min period. (b) Organic correlation of York CHBr_3 mixing ratios and Bristol CHBr_3 relative response data, using data points taken within same 40-min period.

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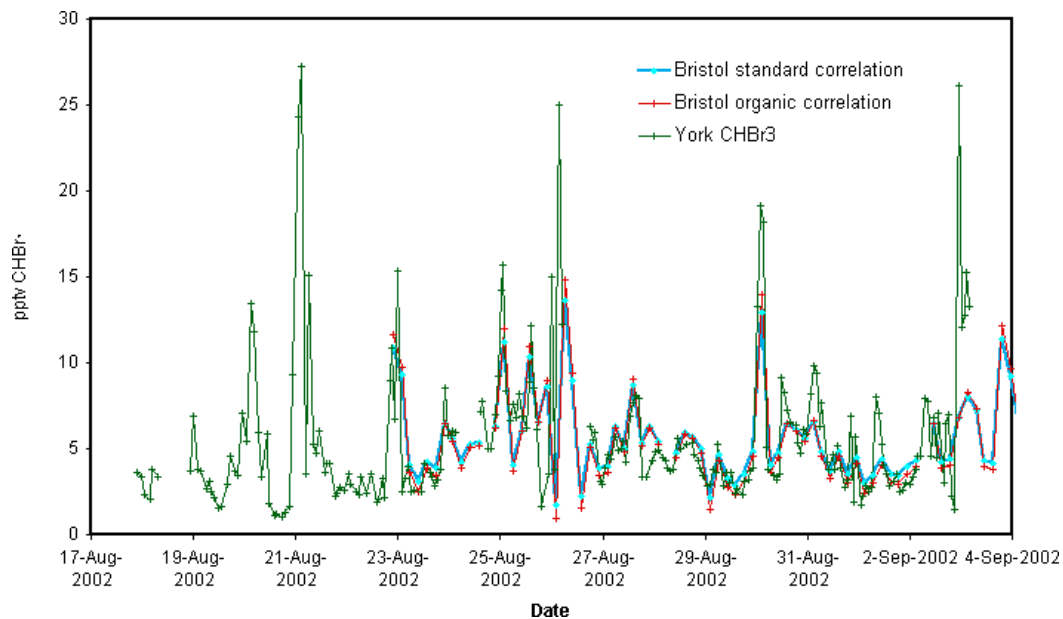


Fig. 2. Comparison of York and Bristol CHBr₃ data, the latter converted to mixing ratios using both the organic correlation of Fig. 1b and a standard correlation.

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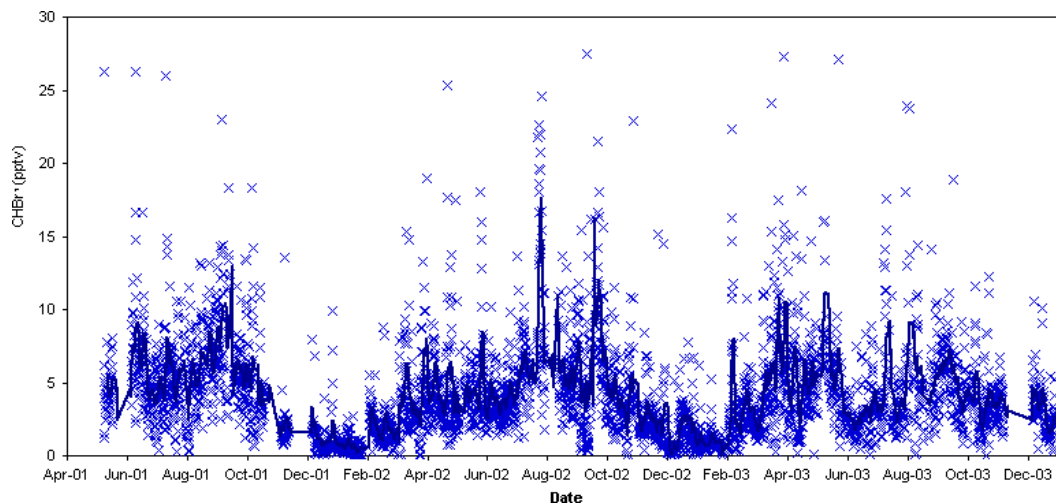


Fig. 3. Bristol CHBr_3 data fixed to the York calibration scale. The crosses are individual (approx. 4-hourly) data points and the solid line is a 20-point moving average.

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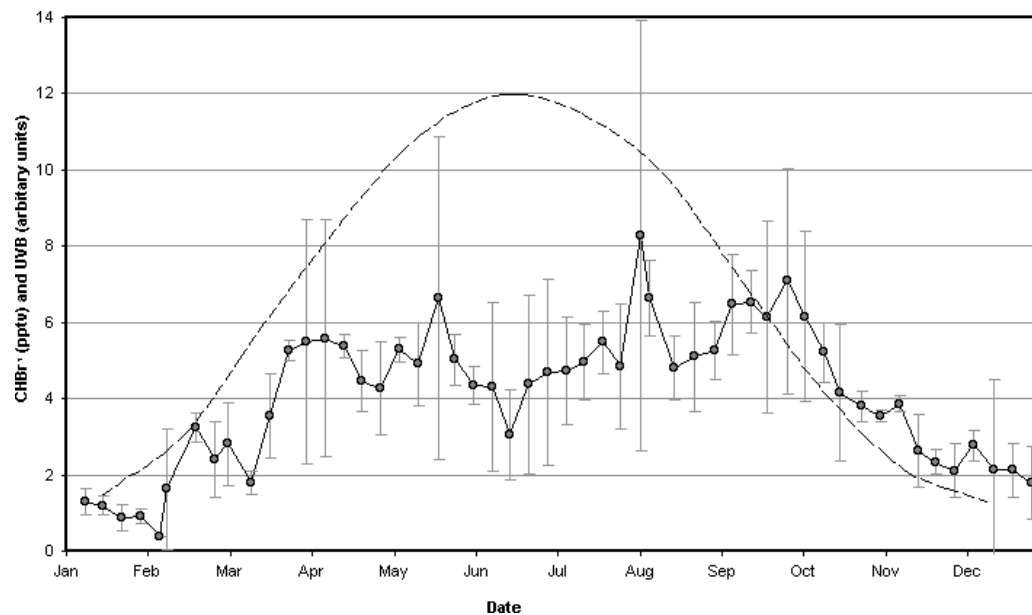


Fig. 4. Mean annual cycle of CHBr₃ and UVB at Mace Head. The error bars on the CHBr₃ data are the RSDs of the 2-weekly average data points.

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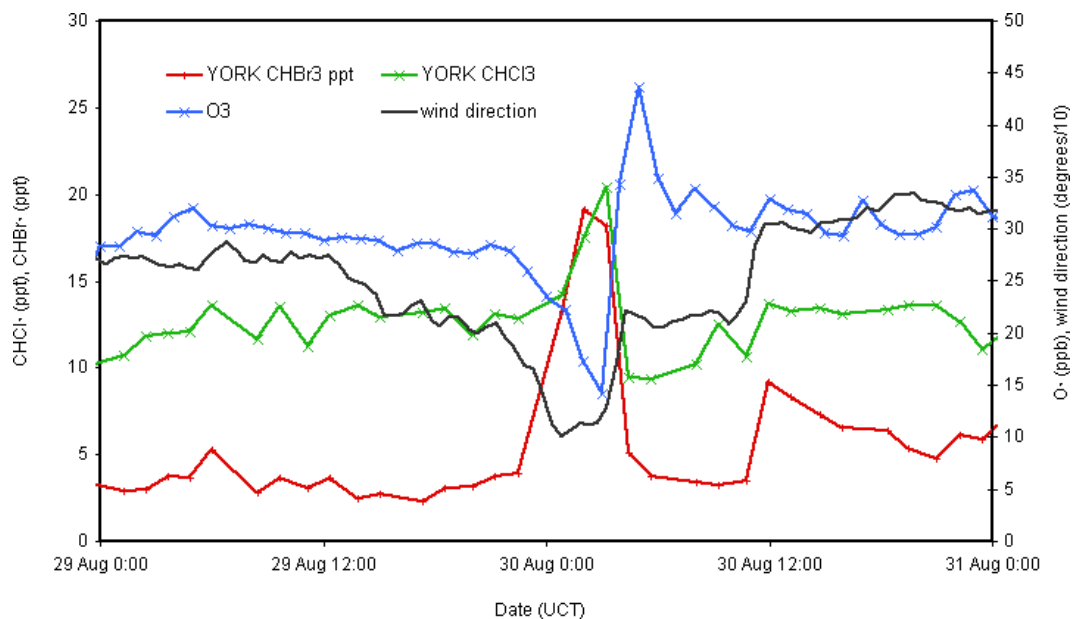


Fig. 5. Land breeze event during the night of 29/30 August 2002.

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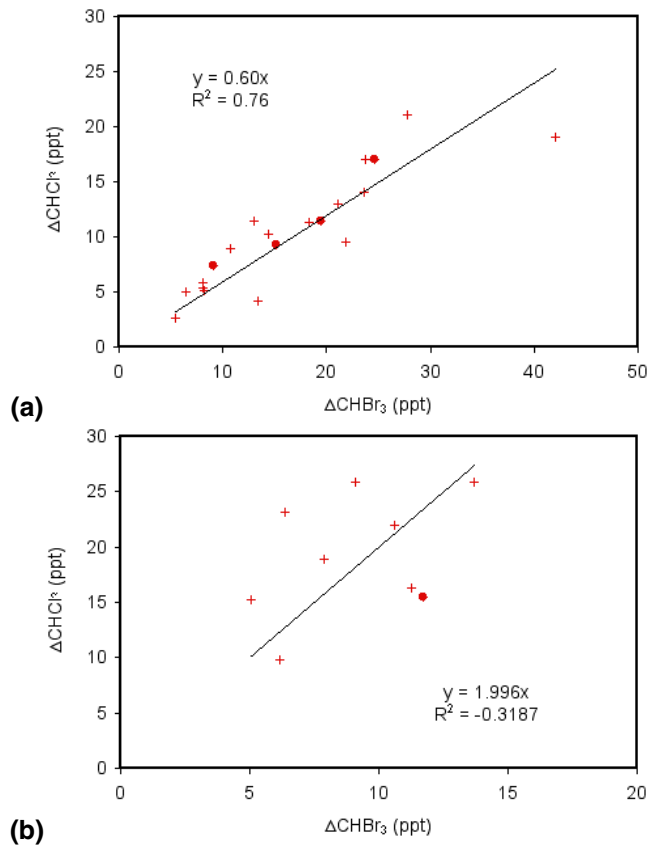


Fig. 6. $\Delta\text{CHCl}_3/\Delta\text{CHBr}_3$ ratios obtained using both Bristol (crosses) and York (filled circle) data, separated into (a) south-easterly ($100\text{--}170^\circ$) and (b) north-easterly ($40\text{--}70^\circ$) winds.

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