Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-235, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 30 March 2016

Atmospheric Chemistry and Physics Discussions

Carbon isotopic signature of coal-derived methane emissions to atmosphere: from coalification to alteration

Giulia Zazzeri[1], Dave Lowry[1], Rebecca E. Fisher[1], James L. France[1,2], Mathias Lanoisellé[1], Bryce F.J. Kelly[4], Jaroslaw M. Necki[3], Charlotte P. Iverach[4], Elisa Ginty[4], Miroslaw Zimnoch[3], Alina Jasek[3] and Euan G. Nisbet[1].

[1]Royal Holloway University of London, Egham Hill, Egham, Surrey TW20 0EX

[2]University of East Anglia, Norwich Research Park, Norwich, Norfolk NR4 7TJ

[3]AGH-University of Science and Technology, Al.Mickiewicza 30 Kraków, Poland

[4]Connected Waters Initiative Research Centre, UNSW Australia

Correspondence to: Dr Giulia Zazzeri (Giulia.Zazzeri.2011@live.rhul.ac.uk)
Prof Euan Nisbet (e.nisbet@es.rhul.ac.uk)

Abstract. Currently, the atmospheric methane burden is rising rapidly, but the extent to which shifts in coal production contribute to this rise is not known. Coalbed methane emissions into the atmosphere are poorly characterised, and this study provides representative $\delta^{13}C_{CH_4}$ signatures to be used in regional and global models in order to allow better apportionment of fossil fuel emissions. Integrated methane emissions from both underground and opencast coal mines in the UK, Australia and Poland were sampled and isotopically characterised. Progression in coal rank and secondary biogenic production of methane due to incursion of water are suggested as the processes affecting the isotopic composition of coal-derived methane. An averaged value of -65 ‰ has been assigned to bituminous coal exploited in open cast mines and of -55 ‰ in deep mines, whereas values of -40 ‰ and -30 ‰ can be allocated to anthracite opencast and deep mines respectively. However, the isotopic signatures that are included in global atmospheric modelling of coal emissions should be region or nation specific, as greater detail is needed, given the wide global variation in coal type.

1 Introduction

Methane emissions from the energy sector have been driven in recent years by the impact of a shift from coal to natural gas, in the US, in the UK and Eastern Europe, whereas in China coal production has increased in this century. Currently, the atmospheric methane burden is rising rapidly (Nisbet et al., 2014), but the extent to which shifts in coal production contribute to this rise is not known. Coalbed methane emissions into the atmosphere are poorly characterised, as they are dispersed over large areas and continue even after the mines' closure (IPCC, 2006). Methane is emitted in coal processing (crashing and pulverisation) and during the initial removal of the overburden; it can be diluted and emitted through ventilation shafts in underground coal mines, or directly emitted to atmosphere from open-cut coal mining, where releases may occur as a result of deterioration of the coal seam.

For the UN Framework Convention on Climate Change, national emissions are estimated by a "bottom-up" approach, based upon a general equation where the coal production data are multiplied by an emission factor that takes into account the mine's gassiness, which in turn is related to the depth of the mine and the coal rank (i.e. carbon content of coal) (U.S. EPA, 2013). These modelled estimates are often calculated without an error

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-235, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 30 March 2016

assessment, and therefore the level of accuracy of the emissions is not known. "Top-down" assessment of methane emissions can be made by chemical transport models constrained by atmospheric measurements (Bousquet et al., 2006; Locatelli et al., 2013). However, this top-down approach provides the total amount of methane emissions into the atmosphere, which has to be distributed among the different methane sources in order to quantify each source contribution.

For methane emissions from fossil fuels (coal and natural gas), the source partitioning is mainly "bottom-up", based on energy use statistics and local inventories, which might be highly uncertain. Conversely, the "top-down" study of the carbon isotopic composition of methane, which is indicative of the methane origin, provides a valuable constraint on the budget appraisal, allowing different sources in a source mix to be distinguished and their individual strength to be evaluated (Liptay et al., 1998; Lowry et al., 2001; Townsend-Small et al., 2012).

Measurements of methane mole fractions can be complemented in atmospheric models by typical $\delta^{13}CCH_4$ signatures of the main methane sources in order to estimate global and regional methane emissions and assess emissions scenarios throughout the past years (Fung et al., 1991; Miller, 2004; Whiticar and Schaefer, 2007; Bousquet et al., 2006; Monteil et al., 2011; Mikaloff Fletcher et al., 2014). However, even though isotopic values are fairly distinctive for specific methanogenic processes, the variety of production pathways and local environmental conditions that discriminate the methane formation process leads to a wide range of $\delta^{13}CCH_4$ values. The global isotopic range for coal is very large, from -80 to -17 ‰ (Rice, 1993), but it can be narrowed down when a specific basin is studied. While there are several studies of isotopic composition of methane generated from coal in Australia, U.S.A. and China (Smith and Rigby, 1981; Dai et al., 1987; Rice et al.; 1989; Aravena et al., 2003; Flores et al., 2008; Papendik et al., 2011), there is a significant lack of information about the isotopic characterisation of methane emissions from coal mines in Europe.

The purpose of this study was to determine links between $\delta^{13}CCH_4$ signatures and coal rank and mining setting, and to provide representative ^{13}C signatures to be used in regional and global atmospheric models in order to produce more accurate methane emission estimates for the coal exploitation sector.

1.1 Process of coalification and parameters affecting the $\delta^{13}C$ signature of methane emissions

The process of coalification involves both biochemical and geochemical reactions. The vegetal matter firstly decays anaerobically under water; the simple molecules derived from initial decomposition (i.e. acetate, CO_2, H_2, NH_4^+, HS^-, long chain fatty acids) are metabolised by fermentative archaea, which produce methane via two methanogenic paths, acetoclastic reaction or CO_2 reduction (Whiticar, 1999). Different pathways lead to diverse $\delta^{13}CCH_4$ isotopic signatures - methane from acetate is ^{13}C enriched relative to methane from CO_2 reduction, ranging from -65 ‰ to -50 ‰ and -110 ‰ to -50 ‰ respectively (Levin et al., 1993; Waldron et al., 1998). With increasing burial and temperatures, coal is subjected to thermal maturation, which implicates more geochemical changes.

As coalification proceeds, the carbon content increases, accompanied by a relative depletion in volatile compounds, such as hydrogen and oxygen, emitted in the form of water, methane, carbon dioxide and higher hydrocarbons through decarboxylation and dehydration reactions (Stach and Murchison, 1982). At higher degrees of coalification and temperature, the liquid hydrocarbons formed in previous stages are thermally cracked to methane, increasing the amount of methane produced (Faiz and Hendry, 2006). Peat and brown coal represent the first stage of the coalification process. The vertical pressure exerted by accumulating sediments

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-235, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 30 March 2016

Atmospheric Chemistry and Physics Discussions

converts peat into lignite. The intensification of the pressure and heat results in the transition from lignite to bituminous coal, and eventually to anthracite, the highest rank of coal (O'Keefe et al., 2013).

During peat and brown coal stages, primary biogenic methane is formed and it is mainly dissolved in water or released during burial, as coal is not appropriately structured for gas retention (Kotarba and Rice, 2001). At more mature stages, thermogenic methane is produced by thermal modification of sedimentary organic matter, which occurs at great depths and intensive heat. Following the basin uplift, methane production can be triggered in the shallower sediments by the meteoric water inflow into the coal (secondary biogenic gas) (Rice, 1993; Scott et al., 1994).

The isotopic signature of the methane produced during the coalification process is controlled by the methane origin pathway (Whiticar, 1996). Thermogenic methane is isotopically enriched in ^{13}C (> -50 ‰) compared to biogenic methane, as methanogens preferentially use the lightest isotopes due to the lower bond energy (Rice, 1993). Intermediate isotopic compositions of methane might reflect a mixing between microbial and thermogenic gases or secondary processes. Indeed, many controlling factors co-drive the fractionation process, and several contentions about their leverage still persist in literature. Deines (1980) asserts that no significant trend is observed in the isotopic signature of methane in relation to the degree of coalification. Conversely, Chung et al. (1979) observed that the composition of the parent material does affect the isotopic composition of the methane accumulated. While the link between isotopic composition and coal rank is not that straightforward, studies carried out in different worldwide coal seams confirm a stronger relationship between coal bed gas composition and depth. Rice (1993), using data from Australian, Chinese and German coal beds, shows that shallow coal beds tend to contain relatively isotopically lighter methane when compared to those at greater depths. In the presence of intrusions of meteoric water, secondary biogenic methane, isotopically lighter, can be generated and mixed with the thermogenic gas previously produced. Colombo et al. (1966) documented a distinct depth correlation in the Ruhr Basin coal in Germany, with methane becoming more ^{13}C-depleted towards the surface zone, independently from coalification patterns. This tendency can be explained either by bacterial methanogenesis, or by secondary processes such as absorption-desorption of methane. Also Scott (2002), in a study about coal seams in the Bowen Basin, Australia, ascribes the progressive methane ^{13}C-enrichment with depth to the meteoritic recharge in the shallowest seams, associated with a higher bacterial activity and a preferential stripping of ^{13}C-CH_4 by water flow.

The migration of methane from the primary zone as a consequence of local pressure release can affect the isotopic composition, since $^{12}CH_4$ diffuses and desorbs more readily than $^{13}CH_4$ (Deines, 1980), but the fractionation effect due to migration is less than 1 ‰ (Fuex, 1980). A much larger variation in the isotopic composition is associated with different methanogenic pathways (30 ‰) and thermal maturation stage (25 ‰) (Clayton, 1998). The measurement of Deuterium, coupled with $\delta^{13}CCH_4$ values, would help to distinguish the pathways of secondary biogenic methane generation, acetoclastic reactions or CO_2 reduction (Faiz and Hendry, 2006), but such distinction is beyond the scope of this study.

Overall, the $\delta^{13}C$ values of methane from coal show an extremely wide range and understanding the processes driving the methane isotopic composition needs to focus on the particular set of geological conditions in each sedimentary basin.

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-235, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 30 March 2016

Atmospheric Chemistry and Physics Discussions

Here we analyse the isotopic signatures of methane plumes emitted to atmosphere, from the dominant bituminous and anthracite mines in Europe and Australia, of both deep and open cut type, to test the theories of isotopic change due to coal rank and interaction with meteoric water.

2 Material and Methods

2.1 Coal basins investigated and type of coal exploited

2.1.1 English and Welsh coal mines

The major coalfields of England and Wales belong to the same stage in the regional stratigraphy of northwest Europe, the Westphalian Stage, between roughly 313 and 304 Ma (Upper Carboniferous). Mining has ceased in many areas. Of those remaining some are located in South Yorkshire (Hatfield and Maltby collieries) (Fig. 1), where 50% of the coalfield's output came from the Barnsley seam, which includes soft coal overlaying a semi-anthracitic coal and bituminous coal in the bottom portion. Seams >2m in thickness are common in the southern half of the Yorkshire coalfield (IMC Group Consulted Limited, 2002). Upper Carboniferous coal measures typical of Yorkshire extend into the East Midlands coalfields. The exposed coalfields are found in the west of this belt, where the coal measures outcrop (roughly from Nottingham to Bradford and Leeds via Chesterfield, Sheffield and Barnsley).

The coal exploited in Daw Mill Colliery is part of the Warwickshire Coalfield, from a seam which varies in thickness from 6.6 to 7.5 m (IMC Group Consulted Limited, 2002).

The coal of the South Wales basin exhibits a well-defined regional progression in rank, which varies from highly volatile bituminous coal in the south and east margin to anthracite in the north-west part, and the main coal-bearing units reach 2.75 km in thickness toward the south-west of the coalfield (Alderton et al., 2004). The coal is now preferentially extracted in opencast mines, as the extensive exploitation of the coal has left the accessible resources within highly deformed structures (e.g. thrust overlaps, vertical faults) that cannot be worked by underground mining methods (Frodsham and Gayer, 1999). Emissions from two deep mines, Unity and Aberpergwm, were investigated, where mine shafts reach depths up to approximately 750 m (http://www.wales-underground.org.uk/pit/geology.shtml).

2.1.2 Upper Silesian Coal Basin in Poland

The Upper Silesian Coal Basin extends from Poland to the Czech Republic and is one of the largest coal basins in Europe, with an area of ~7400 km (Jureczka and Kotas, 1995). The Silesian Region of Poland is estimated to be responsible for significant methane emissions, in the range of 450-1350 Gg annually (Patyńska, 2013). The upper Carboniferous coal-bearing strata of this region are associated with gas deposits of both thermogenic and microbial origin, and the methane content and spatial distribution are coal rank related (Kędzior, 2009) - i.e. the sorption capacity of coal in the basin is found to increase with coal rank. Most of the methane generated during the bituminous stage in the coalification process has escaped from the coal source following basin uplift during Paleogene (Kotarba and Rice, 2001) and diffused through fractures and faults occurring in tectonic zones. The late-stage gas generated by microbial reduction of CO_2 in the coal seams at the top of the Carboniferous sequence accumulated under clay deposition in the Miocene (Kędzior, 2009).

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-235, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 30 March 2016

Atmospheric Chemistry and Physics Discussions

2.1.3 Australia: Hunter Coalfield (Sydney Basin)

The Hunter Coalfield is part of the Sydney Basin, on the east coast of New South Wales in Australia, and consists of 3 major coal measures. The deepest is the early Permian Greta Coal Measures, which is overlain by the late Permian Whittingham Coal Measures and upper Newcastle Coal Measures. Throughout the Hunter coalfield all sedimentary strata are gently folded, and the same coal seam can be mined at the ground surface and at depths of several hundred meters. In the region surveyed both the opencast and underground mines are extracting coal from the Whittingham coal measures, which are generally high volatile bituminous coals, although some medium to low bituminous coals are extracted (Ward and Kelly, 2013). The mining operation is on a much larger scale than in the UK - the total coal production for the Hunter Coalfield was 123.63 Mt in 2011, of which 88.24 Mt was saleable (State of New South wales, 2013)-. Several studies have attested to the dispersion of thermogenic methane formed at higher degrees of coalification (i.e. high temperatures and pressures) during the uplift and the subsequent erosion of the basin, followed by the replenishment of the unsaturated basin with more recently formed methane of biogenic (Faiz and Hendry, 2006; Burra, 2010). Gas emplacement is related to sorption capacity of coal strata, in particular to the pore pressure regime, which is influenced by the local geological features (compressional or extensional) of the basin. The south of the Hunter Coalfield is characterized by higher gas content and enhanced permeability than the northern area, with a large potential for methane production, mainly biogenic (Pinetown, 2014).

2.2 Sampling and measurement methodology

For isotopic characterisation of the methane sources, integrated methane emissions were assessed through detection of the offsite downwind plume. In fact, even when emissions are focused on defined locations, such as vent pipes in underground mines, the methane provenance cannot be localised, since most of collieries are not accessible. For sample collection and measurements of methane emissions downwind of coal mines in the UK and Australia the mobile system described by Zazzeri et al. (2015) has been implemented. The system utilises a Picarro G2301 CRDS (Cavity Ring-Down Spectroscopy) within the survey vehicle, for continuous CH_4 and CO_2 mole fraction measurements, and a mobile module including air inlet, sonic anemometer and GPS receiver on the roof of the vehicle. The entire system is controlled by a laptop, which allows methane mole fractions and the methane plume outline to be displayed in real time on a Google Earth platform during the survey to direct plume sampling. When the plume was encountered, the vehicle was stopped and air samples collected in 3L Tedlar bags, using a diaphragm pump connected to the air inlet. Samples were taken at different locations along the plume transect in order to obtain a wide range of methane mole fractions and isotopic signatures in the collected air. The Upper Silesian basin was surveyed with a Picarro 2101-i measuring continuous CO_2 and CH_4 mole fractions and $\delta^{13}CCO_2$ isotopic ratio with a precision of 0.3 ‰ in 5 minutes. Samples were collected on site for analysis of $\delta^{13}CCH_4$ isotopic ratio.

The carbon isotopic ratio ($\delta^{13}C$) of bag samples was measured in the greenhouse gas laboratory at RHUL (Royal Holloway University of London) in triplicate to high precision (±0.05 ‰) by continuous flow gas chromatography isotope ratio mass spectrometry (CF GC-IRMS) (Fisher et al., 2006). CH_4 and CO_2 mole fractions of samples were measured independently in the laboratory with a Picarro G1301 CRDS analyser, calibrated against the NOAA (National Oceanic and Atmospheric Administration) WMO-2004A and WMO-X2007 reference scales respectively. The $\delta^{13}CCH_4$ signature for each emission plume was calculated using the

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-235, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 30 March 2016

Keeling plot approach, according to which the $\delta^{13}C$ isotopic composition of samples and the inverse of the relative mole fractions have a linear relationship, whose intercept represents the isotopic signature of the source (Pataki et al., 2003). Source signatures were provided with the relative uncertainty, computed by the BCES (Bivariate Correlated Errors and intrinsic Scatter) estimator (Akritas and Bershady, 1996), which accounts for correlated errors between two variables and calculates the error on the slope and intercept of the best interpolation line. Mole fraction data and co-located coordinates were used to map the mole fraction variability using the ArcGIS software.

3 Results and Discussion

While most of the emissions from deep mines come specifically from ventilation shafts, which are point sources, emissions from open-cut mines are wide-spread, and difficult to estimate. However, the objective of this study is not the quantification of emissions, but the assessment of the overall signature of methane released into the atmosphere, made through the sampling of integrated emissions from the whole area. Therefore, even though onsite access to collieries was not possible, by driving around the contiguous area, methane emissions could be intercepted and their mole fractions measured. Table 1 summarises $\delta^{13}CCH_4$ signatures of all the coal mines and coal basins surveyed with the Picarro mobile system. The English and Welsh coal mines surveyed are shown in Fig. 1.

3.1 English coal mines

Emissions from Hatfield colliery, one of the few UK deep mines still open at the time of this study, were investigated on 10th July and 26th September 2013. The main seams that have been worked since 1920's are the Barnsley, the Dunsil and the High Hazel seams, at approximate depths of 401-365 m 414-376 and 313-284 m respectively (Hill, 2001). Mole fractions over 7 ppm were recorded during both surveys while transecting the plume (Fig. 2). The methane desorbed from coal heaps within the colliery area and methane vented from shafts might explain the relatively high mole fractions measured. Keeling plots based on the samples collected (4 on the first and 5 on the second survey) give intercept values of -48.3 ±0.2 and -48.8 ±0.3 ‰ (2SD).

Maltby colliery is one of the largest and deepest mines in England, reaching 991 m in depth, and was closed in March 2013. High volatile bituminous coal was extracted from the Barnsley and Parkgate seams (McEvoy et al., 2006). The coal mine methane was extracted and used for electricity, but over 3 ppm mole fractions were detected in ambient air during both surveys in July and September 2013, giving evidence of methane releases from the recovery system and ventilation shafts. The Keeling plot intercept based on the samples collected in July and in September are -45.9 ±0.3 and -45.4 ±0.2 ‰ (2SD) respectively, ~3 ‰ heavier for Hatfield Colliery on the same sampling days. Both source signatures are in good agreement with the value of -44.1 ‰ observed for desorbed methane of the Barnsley coal seam, in a study conducted by Hitchman et al. (1990b).

Kellingley colliery is a still open deep mine situated in North Yorkshire, which has exploited the Beeston and Silkstone coal seams, characterised by a high quality, low sulphur coal used for coke manufacture. A surface drainage plant for electrical power generation has been implemented (Holloway et al., 2005), and methane releases from the power plant might justify the mole fractions peak of 9 ppm that was observed while driving on

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-235, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 30 March 2016

the north side of the coal mine on 10th July 2013. The Keeling plot analysis of the samples collected indicates a source signature of -46.5 ±0.3 ‰ (2SD).

Thoresby mine is located in Nottinghamshire and is currently exploiting the Parkgate seam at about 650 m underground. The highest mole fractions detected approached 5 ppm, and 9 samples were collected, giving a source signature of -51.2 ±0.3 ‰ (2SD).

The Daw Mill colliery, before its closure in March 2013, was Britain's biggest coal mine, working the Warwickshire Thick seam, which lies between 500 and 1000 m in depth. The estimated methane content of this seam is low (typically about 1.7 m^3/tonne) and, furthermore, a ventilation system is implemented, so that the coal mine methane potential is curtailed (Drake, 1983). During the sampling, the highest methane mole fractions (≈5 ppm) were recorded close to the edge of the colliery, whereas, driving downwind of the site at further distances, only background values were measured. Keeling plot analysis reveals a source signature of -51.4 ±0.2 ‰ (2SD).

3.2 Welsh coal mines

Methane emissions from coal mines in Wales were sampled in order to characterise methane releases from a different rank of coal. The area investigated extended from Cwmllynfell to Merthyr Tydfil (Fig. 1). The South Wales coalfield is estimated to have the highest measured seam gas content in the UK (Creedy, 1991).

3.2.1 Deep Mines

The deep mine Aberpergwm, which closed in December 2012, did not operate coal mine methane schemes and methane was vented up to the surface as part of standard operation systems (Holloway et al., 2005). That is consistent with methane mole fractions peaks of 6 ppm observed when approaching the colliery. Air samples were also collected near Unity deep mine, Wales' largest drift mine, which reopened in 2007 and is located in the town of Cwmgwrach, only one mile away from Aberpergwm. Isotopic source signatures of 33.3 ±1.8 ‰ and -30.9 ±1.4 ‰ result from the Keeling plots based on samples collected respectively near Aberpergwm and Unity colliery, both highly ^{13}C enriched relative to all English collieries.

3.2.2 Opencast Mines

The Picarro mobile system was driven around opencast mines at Cwmllynfell and Abercrave, in the Swansea Valley. Up to 3 ppm methane mole fractions were recorded near the two mines, which were closed in the 1960s as drift mines and are currently exploited as opencast mines. Our measurements confirm that they are still emitting methane, albeit at levels which are not significantly above background. ^{13}C signatures between -41.4 ±0.5 and -41.2 ±0.9 ‰ (2SD) result from isotopic analysis of samples collected downwind of the opencast mines, approximately 10 ‰ lower than the isotopic signature characterizing Welsh deep anthracite mines.

3.3 Polish coal mine

A Picarro mobile survey of the Upper Silesian basin took place on 10th June 2013 and 12 air samples were collected for isotopic analysis. All the mines in this area are deep mines, exploiting the coal at depths ranging from 300 to 900 m. The Keeling plot analysis includes 8 samples collected around the area of Radoszowy, downwind of the KWK Wujek deep mine shafts (white stars in Fig. 4). Methane mole fractions in the range of

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-235, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 30 March 2016

Atmospheric Chemistry and Physics Discussions

263 3-5 ppm were measured in the majority of the area of Katowice and over 20 ppm mole fractions were detected
264 when transecting the plume originating from the exhaust shafts, which confirms the high level of methane that
265 the mine contains. A source signature of -50.9 ± 0.6 ‰ (2SD) was calculated by Keeling plot analysis, which is
266 consistent with the values obtained for the deep mined English bituminous coal.

267 **3.4 Australia: Hunter Coalfield**

268 On 12th and 18th March 2014 12 samples in total were collected along the route in the Hunter Coalfield, where
269 the bituminous coal strata of the Sydney Basin are extracted both in opencast and underground mines. The
270 methane plume width was in the range of 70 km (Fig. 5a). A maximum mole fraction of 13.5 ppm was measured
271 near a vent shaft associated with the Ravensworth underground mine (see white star in Fig. 5b), which exploits
272 the Lemington, Pikes Gully, Lidell (Upper and Middle) and Barret Seams, the deepest of which is the Barret
273 seam, with a maximum overburden depth of ~350 m (GSS Environmental, 2012). The source signature
274 calculated by the Keeling plot analysis based on all the samples collected during both surveys (grey markers in
275 Fig. 5c) is -66.4 ±1.3 ‰ (2SD) and this signature likely includes a mixture of methane derived both from
276 underground and opencast mines. Two samples collected downwind of the Ravensworth ventilation shaft (red
277 pushpins in Fig. 5b) fall off the Keeling plot trend for March 2014 and are not included in the calculation,
278 because they are likely dominated by methane from the vent shaft and are not representative of the regional
279 mixed isotopic signature.
280 In January 2016, 10 samples were collected downwind of a ventilation fan in the Bulga mine (second white star
281 in Fig. 5b), aerating underground workings in the Blakefield South Seam, ranging from 130 to 510 m in depth
282 (Bulga Underground Operation mining, 2015). The Keeling plot for these (black circles in Fig. 5c) indicates a
283 $\delta^{13}C$ source signature of -60.8 ±0.3 ‰ (2SD). The samples collected next to the Ravensworth ventilation shaft in
284 2014 fit on this Keeling plot, suggesting that the $\delta^{13}CCH_4$ isotopic signature of emissions from underground
285 works in the Hunter Coalfield is consistent.

286 **3.5 New representative $\delta^{13}CCH_4$ isotopic signatures for coal-derived methane**

287 The $\delta^{13}CCH_4$ isotopic values for coal have been found to be characteristic of single basins, but general
288 assumptions can be made to characterise coal mines worldwide. Table 2 provides the literature $\delta^{13}CCH_4$ isotopic
289 values characteristic of specific coal basins. The isotopic signatures of emissions from English bituminous coal
290 are consistent with the range of -49 to -31 ‰ suggested by Colombo et al. (1970) for in situ coal bed methane in
291 the Ruhr basin in Germany, which contains the most important German bituminous coal of Upper Carboniferous
292 age and low volatile anthracite (Thomas, 2002). Progression in coal rank might explain the value of -50 ‰ for
293 emissions from English underground mined bituminous coal and -30 ‰ for anthracite deep mines, followed by a
294 5-10 ‰ ^{13}C-depletion caused by the incursion of meteoric water in the basin and the subsequent production of
295 secondary biogenic methane, resulting in -40 ‰ for methane plumes from Welsh open-cut anthracite mines.
296 The link between coal rank and $\delta^{13}CCH_4$ isotopic signature is appreciable in the study of UK coal mines, but
297 differences in the $^{13}C/^{12}C$ isotopic ratio within the same coal sequences can be ascribed to other parameters, such
298 as the depth at which the coal is mined and the occurrence of water incursion. In particular, emissions from
299 Thoresby are more ^{13}C depleted than those measured around Maltby, although both mines exploit the Parkgate
300 seam, meaning that different isotopic signatures cannot be entirely linked to the coal rank. Biogenic methane

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-235, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 30 March 2016

produced in a later stage due to water intrusion might have been mixed with the original thermogenic methane formed during the coalification process.

Differences in methane emissions and their isotopic signature between opencast and deep mines have been assessed by surveying both surface and underground mines, in the Welsh anthracite belt, and in the Hunter Coalfield. The shallower deposits have been more exposed to the weathering and meteoric water, most likely associated with the production of some isotopically lighter microbial methane. Mole fractions up to 2.5 ppm were measured around opencast mines in the Hunter Coalfield, in Australia, within a methane plume of more than 70 km width. The highest methane mole fractions were consistently measured downwind of vent shafts in underground mines. The difference in the source isotopic signature for methane emissions between the two types of mining in the Hunter Coalfield (from -61 to -66 ‰) and in Wales (from -31 to -41 ‰) reflects the isotopic shift of 5-10 ‰ that has been attributed to the occurrence of secondary biogenic methane.

The $\delta^{13}C$ signatures for coalbed methane emissions from the Upper Silesian basin are highly variable, with the most ^{13}C depleted methane associated with diffusion processes or secondary microbial methane generation (Kotarba and Rice, 2001), but the value of -51 ‰ measured for methane emissions from the KWK Wujek deep mine is consistent with the value of -50 ‰ inferred for emissions from English bituminous coal extracted in underground mines.

4 Conclusions

By measuring the isotopic signatures of methane plumes from a representative spread of coal types and depths, we show that the $\delta^{13}C$ isotopic value to be included in regional and global atmospheric models for the estimate of methane emissions from the coal sector must be chosen according to coal rank and type of mining (opencast or underground). For low resolution methane modelling studies an averaged value of -65 ‰ is suggested for bituminous coal exploited in open cast mines and of -55 ‰ in deep mines, whereas values of -40 ‰ and -30 ‰ can be assigned to anthracite opencast and deep mines respectively.

Global methane budget models that incorporate isotopes have used a $\delta^{13}C$ signature of -35 ‰ for coal or a value -40 ‰ for total fossil fuels (e.g. Hein et al., 1997; Mikaloff Fletcher et al., 2004; Bousquet et al., 2006; Monteil et al., 2011), but, given the relative rarity of anthracite coal reserves and the dominance of bituminous coal (http://www.transparencymarketresrch.com/anthracite-coal-mining.html), it seems likely that a global average emission from coal mining activities will be lighter, with the -50 ‰ recorded for deep-mined bituminous coal in Europe being a closer estimate. However, for detailed global modelling of atmospheric methane, isotopic signatures of coal emissions should be region or nation specific, as greater detail is needed given the wide global variation. The assignment of an incorrect global mean, or a correct global mean but inappropriate for regional scale modelling, might lead to incorrect emissions estimates or source apportionment. The new scheme gives the possibility for an educated estimate of the $\delta^{13}C$ signature of emission to atmosphere to be made for an individual coal basin or nation, given information on the type of coal being mined and the method of extraction.

In conclusion, high-precision measurements of $\delta^{13}C$ in plumes of methane emitted to atmosphere from a range of coal mining activities have been used to constrain the isotopic range for specific ranks of coal and mine type, offering more representative isotopic signatures for use in methane budget assessment at regional and global scales.

Acknowledgements

Giulia Zazzeri would like to thank Royal Holloway, University of London for provision of a Crossland scholarship and a contribution from the Department of Earth Sciences from 2011 to 2014. Analysis of samples from Poland was funded through the European Community's Seventh Framework Programme (FP7/2007-2013) in the InGOS project under grant agreement n. 284274. Sampling in Australia was possible due to a grant from the Cotton Research and Development Corporation.

References

Akritas, M.G., and Bershady, M.A., 1996, Linear regression for astronomical data with measurement errors and intrinsic scatter: Astrophysical Journal, v. 470, p. 706-714.

Alderton, D., Oxtoby, N., Brice, H., Grassineau, N., and Bevins, R., 2004, The link between fluids and rank variation in the South Wales Coalfield: evidence from fluid inclusions and stable isotopes: Geofluids, v. 4, p. 221-236.

Aravena, R., Harrison, S., Barker, J., Abercrombie, H., and Rudolph, D., 2003, Origin of methane in the Elk Valley coalfield, southeastern British Columbia, Canada: Chemical Geology, v. 195, p. 219-227.

Bousquet, P., Ciais, P., Miller, J., Dlugokencky, E., Hauglustaine, D., Prigent, C., Van der Werf, G., Peylin, P., Brunke, E.-G., and Carouge, C., 2006, Contribution of anthropogenic and natural sources to atmospheric methane variability: Nature, v. 443, p. 439-443.

Bulga Underground Operations Mining: Operation Plan 2015 to 2021, 2015.

Burra, A., 2010, Application of domains in gas-in-place estimation for opencut coal mine fugitive gas emissions reporting, Bowen Basin Symposium, p. 59-64.

Chung, H.M., and Sackett, W.M., 1979, Use of stable carbon isotope compositions of pyrolytically derived methane as maturity indices for carbonaceous materials: Geochimica et Cosmochimica Acta, v. 43, no.12.

Clayton, J., 1998, Geochemistry of coalbed gas–A review: International Journal of Coal Geology, v. 35, p. 159-173.

Colombo U., G.F., Gonfiantini F., Gonfiantini R., Kneuper G., Teichmuller I., Teichmuller R. , 1970, Carbon isotope study on methane from German coal deposits.: Advances in Organic Geochemistry 1966 (eds G. D. HOBSON and G. C. SPEERS), p. 1-26.

Creedy, D.P., 1991, An introduction to geological aspects of methane occurrence and control in British deep coal mines: Quarterly Journal of Engineering Geology, v. 24, p. 209-220.

Dai, J.X., Qi, H.F., Song, Y., and Guan, D.S., 1987, Composition, carbon isotope characteristics and the origin of coal-bed gases in China and their implication: Scientia Sinica Series B-Chemical Biological Agricultural Medical & Earth Sciences, v. 30, p. 1324-1337.

Deines, P., 1980, The isotopic composition of reduced organic carbon: Handbook of environmental isotope geochemistry, p. 329-406.

Drake, D., 1983, Working the Warwickshire Thick Coal, Improved Techniques for the Extraction of Primary Forms of Energy, Springer Netherlands, p. 156-156.

Faiz, M., and Hendry, P., 2006, Significance of microbial activity in Australian coal bed methane reservoirs—a review: Bulletin of Canadian Petroleum Geology, v. 54, p. 261-272.

377 Fisher, R., Lowry, D., Wilkin, O., Sriskantharajah, S., and Nisbet, E.G., 2006, High-precision, automated stable
378 isotope analysis of atmospheric methane and carbon dioxide using continuous-flow isotope-ratio mass
379 spectrometry: Rapid Communications in Mass Spectrometry, v. 20, p. 200-208.
380 Flores, R.M., Rice, C.A., Stricker, G.D., Warden, A., and Ellis, M.S., 2008, Methanogenic pathways of coal-bed
381 gas in the Powder River Basin, United States: the geologic factor: International Journal of Coal Geology,
382 v. 76, p. 52-75.
383 Frodsham, K., and Gayer, R., 1999, The impact of tectonic deformation upon coal seams in the South Wales
384 coalfield, UK: International Journal of Coal Geology, v. 38, p. 297-332.
385 Fuex, A., 1980, Experimental evidence against an appreciable isotopic fractionation of methane during
386 migration: Physics and Chemistry of the Earth, v. 12, p. 725-732.
387 Fung, I., John, J., Lerner, J., Matthews, E., Prather, M., Steele, L., and Fraser, P., 1991, Three-dimensional
388 model synthesis of the global methane cycle: Journal of Geophysical Research, v. 96, no.D7.
389 GSS Environmental, 2012, Ravensworth Underground Mine: Liddell Seam Project.
390 Hein, R., Crutzen, P.J., and Heimann, M., 1997, An inverse modeling approach to investigate the global
391 atmospheric methane cycle: Global Biogeochemical Cycles, v. 11, p. 43-76.
392 Hill, A., 2001, The South Yorkshire Coalfield: A History and Development, Tempus Books, Gloucestershire.
393 Hitchman, S., Darling, W., and Williams, G., 1990a, Stable isotope ratios in methane containing gases in the
394 United Kingdom.
395 Hitchman, S., Darling, W., and Williams, G., 1990b, Stable isotope ratios in methane containing gases in the
396 United Kingdom. Report of the British Geological Survey, Processes Research Group WE/89/30.
397 Holloway, S., Jones, N., Creedy, D., and Garner, K., 2005, Can new technologies be used to exploit the coal
398 resources in the Yorkshire-Nottinghamshire coalfield?: Carboniferous Hydrocarbon Geology: The
399 Southern North Sea Surrounding Areas, v. 7 Occasional publications of the Yorkshire Geological Society,
400 p. 195–208.
401 IMC Group Consulting Limited, 2002, A review of the remaining reserves at deep mines for the Department of
402 Trade and Industry.
403 IPCC, 2006, 2006 IPCC guidelines for national greenhouse gas inventories: industrial processes and product
404 use, Kanagawa, JP: Institute for Global Environmental Strategies.
405 Jureczka, J., and Kotas, A., 1995, Coal deposits—Upper Silesian Coal Basin: The carboniferous system in
406 Poland, v. 148, p. 164-173.
407 Kanduč, T., Grassa, F., Lazar, J., and Zavšek, S., 2015, Geochemical and isotopic characterization of coalbed
408 gases in active excavation fields at Preloge and Pesje (Velenje Basin) mining areas: RMZ–M&G, v. 62, p.
409 21-36.
410 Kędzior, S., 2009, Accumulation of coal-bed methane in the south-west part of the Upper Silesian Coal Basin
411 (southern Poland): International Journal of Coal Geology, v. 80, p. 20-34.
412 Kotarba, M.J., and Rice, D.D., 2001, Composition and origin of coalbed gases in the Lower Silesian basin,
413 southwest Poland: Applied Geochemistry, v. 16, p. 895-910.
414 Levin, I., Bergamaschi, P., Dörr, H., and Trapp, D., 1993, Stable isotopic signature of methane from major
415 sources in Germany: Chemosphere, v. 26, p. 161-177.

Liptay, K., Chanton, J., Czepiel, P., and Mosher, B., 1998, Use of stable isotopes to determine methane oxidation in landfill cover soils: Journal of Geophysical Research: Atmospheres (1984–2012), v. 103, p. 8243-8250.

Locatelli, R., Bousquet, P., Chevallier, F., Fortems-Cheney, A., Szopa, S., Saunois, M., Agusti-Panareda, A., Bergmann, D., Bian, H., and Cameron-Smith, P., 2013, Impact of transport model errors on the global and regional methane emissions estimated by inverse modelling: Atmospheric chemistry and physics, v. 13, p. 9917-9937.

Lowry, D., Holmes, C.W., Rata, N.D., O'Brien, P., and Nisbet, E.G., 2001, London methane emissions: Use of diurnal changes in concentration and $\delta^{13}C$ to identify urban sources and verify inventories: Journal of Geophysical Research: Atmospheres, v. 106, p. 7427-7448.

McEvoy F. M., M.D., Harrison D.J., Cameron D.G., Burke H.F., Spencer N.A., Evans D.J., Lott G.K., Hobbs S.F. and Highley D.E., 2006, Mineral Resource Information in Support of National, Regional and Local Planning: South Yorkshire (comprising Metropolitan Boroughs of Barnsley, Doncaster and Rotherham and City of Sheffield) British Geological Survey Commissioned Report.

Mikaloff Fletcher, S.E., Tans, P.P., Bruhwiler, L.M., Miller, J.B., and Heimann, M., 2004, CH_4 sources estimated from atmospheric observations of CH4 and its $^{13}C/^{12}C$ isotopic ratios: 2. Inverse modeling of CH_4 fluxes from geographical regions: Global Biogeochemical Cycles, v. 18, no.4.

Miller, J.B., 2004, The carbon isotopic composition of atmospheric methane and its constraints on the global methane budget, Stable Isotopes and Biosphere-Atmosphere Interactions: Processes and Biological Controls, Elsevier Academic Press California, p. 288-306.

Monteil, G., Houweling, S., Dlugockenky, E., Maenhout, G., Vaughn, B., White, J., and Rockmann, T., 2011, Interpreting methane variations in the past two decades using measurements of CH_4 mixing ratio and isotopic composition: Atmospheric chemistry and physics, v. 11, p. 9141-9153.

Nisbet, E.G., Dlugokencky, E.J., and Bousquet, P., 2014, Methane on the rise—again: Science, v. 343, p. 493-495.

O'Keefe, J.M., Bechtel, A., Christanis, K., Dai, S., DiMichele, W.A., Eble, C.F., Esterle, J.S., Mastalerz, M., Raymond, A.L., and Valentim, B.V., 2013, On the fundamental difference between coal rank and coal type: International Journal of Coal Geology, v. 118, p. 58-87.

Papendick, S.L., Downs, K.R., Vo, K.D., Hamilton, S.K., Dawson, G.K., Golding, S.D., and Gilcrease, P.C., 2011, Biogenic methane potential for Surat Basin, Queensland coal seams: International Journal of Coal Geology, v. 88, p. 123-134.

Pataki, D.E., Bowling, D.R., and Ehleringer, J.R., 2003, Seasonal cycle of carbon dioxide and its isotopic composition in an urban atmosphere: Anthropogenic and biogenic effects: Journal of Geophysical Research-Atmospheres, v. 108, no.D23.

Patyńska, R., 2013, Prognoza wskaźników emisji metanu z kopalń metanowych węgla kamiennego w Polsce: Polityka Energetyczna, v. 16.

Pinetown, K., 2014, Regional coal seam gas distribution and burial history of the Hunter Coalfield, Sydney Basin: Australian Journal of Earth Sciences, v. 61, p. 409-426.

Qin, S., Tang, X., Song, Y., and Wang, H., 2006, Distribution and fractionation mechanism of stable carbon isotope of coalbed methane: Science in China Series D: Earth Sciences, v. 49, p. 1252-1258.

456 Rice, D.D., 1993, Composition and origins of coalbed gas: Hydrocarbons from coal: AAPG Studies in Geology,
457 v. 38, p. 159-184.
458 Rice, D.D., Clayton, J.L., and Pawlewicz, M.J., 1989, Characterization of coal-derived hydrocarbons and
459 source-rock potential of coal beds, San Juan Basin, New Mexico and Colorado, USA: International Journal
460 of Coal Geology, v. 13, p. 597-626.
461 Scott, A.R., 2002, Hydrogeologic factors affecting gas content distribution in coal beds: International Journal of
462 Coal Geology, v. 50, p. 363-387.
463 Scott, A.R., Kaiser, W., and Ayers Jr, W.B., 1994, Thermogenic and secondary biogenic gases, San Juan Basin,
464 Colorado and New Mexico--implications for coalbed gas producibility: AAPG bulletin, v. 78, p. 1186-
465 1209.
466 Smith, J.W., Gould, K., and Rigby, D., 1981, The stable isotope geochemistry of Australian coals: Organic
467 Geochemistry, v. 3, p. 111-131.
468 Stach, E., and Murchison, D.G., 1982, Stach's Textbook of coal petrology, Borntraeger, Stuttgart (1982), 426 p.
469 State of New South Wales, Department of Trade & Investment, Division of Resources & Energy (2013). 2013
470 New South Wales Coal Industry profile.
471 Thomas, L., 2002, Coal geology, John Wiley & Sons.
472 Townsend-Small, A., Tyler, S.C., Pataki, D.E., Xu, X.M., and Christensen, L.E., 2012, Isotopic measurements
473 of atmospheric methane in Los Angeles, California, USA: Influence of "fugitive" fossil fuel emissions:
474 Journal of Geophysical Research-Atmospheres, v. 117, no.D7.
475 U.S. EPA, 2013, US Environmental Protection Agency, Global Mitigation of Non-CO_2 Greenhouse Gases:
476 2010–2030, USEPA, Washington, DC EPA-430-R-13-011.
477 Waldron, S., Fallick, A., and Hall, A., 1998, Comment on" Spatial distribution of microbial methane production
478 pathways in temperate zone wetland soils: Stable carbon and hydrogen evidence" by ERC Hornibrook, FJ
479 Longstaffe, and WS Fyfe: Geochimica et Cosmochimica Acta, v. 62, p. 369-372.
480 Ward, C., and Kelly, B.F.J., 2013, Background Paper on New South Wales Geology: with a focus on basins
481 containing coal seam gas resources., UNSW Global, The University of New South Wales, a report
482 prepared for the Office of the NSW Chief Scientist and Engineer.
483 Whiticar, M., and Schaefer, H., 2007, Constraining past global tropospheric methane budgets with carbon and
484 hydrogen isotope ratios in ice: Philosophical Transactions of the Royal Society of London A:
485 Mathematical, Physical and Engineering Sciences, v. 365, p. 1793-1828.
486 Whiticar, M.J., 1996, Stable isotope geochemistry of coals, humic kerogens and related natural gases:
487 International Journal of Coal Geology, v. 32, p. 191-215.
488 Whiticar, M.J., 1999, Carbon and hydrogen isotope systematics of bacterial formation and oxidation of methane:
489 Chemical Geology, v. 161, p. 291-314.
490 Zazzeri, G., Lowry, D., Fisher, R., France, J., Lanoisellé, M., and Nisbet, E., 2015, Plume mapping and isotopic
491 characterisation of anthropogenic methane sources: Atmospheric Environment, v. 110, p. 151-162.
492

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-235, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 30 March 2016

Atmospheric Chemistry and Physics Discussions

493

494

Sampling Site	Country	Sampling Date	$\delta^{13}C$ Signatures [‰]	Samples number used in Keeling plots
Kellingley Colliery	North Yorkshire/UK	Sept-2013	-46.5 ±0.3	5
Maltby Colliery	South Yorkshire/UK	Jul-2013	-45.9 ±0.3	3
		Sept-2013	-45.4 ±0.2	4
Hatfield Colliery	South Yorkshire/UK	Jul-2013	-48.3 ±0.2	4
		Sept-2013	-48.8 ±0.3	5
Thoresby Colliery	Nottinghamshire/UK	Nov-2013	-51.2 ±0.3	9
Daw Mill Colliery	Warwickshire/UK	Nov-2013	-51.4 ±0.2	4
Cwmllynfell Colliery	Wales/UK	Oct-2013	-41.2 ±0.9	4
Abercrave Colliery	Wales/UK	Oct-2013	-41.4 ±0.5	5
Aberpergwm Colliery	Wales/UK	Oct-2013	-33.3 ±1.8	3
Unity Colliery	Wales/UK	Oct-2013	-30.9 ±1.4	5
Hunter Coalfield	Australia	Mar-2014	-66.4 ±1.3	12
Bulga Colliery	Australia	Jan-2016	-60.8 ±0.3	10
Boggabri and Tarrawonga Collieries	Australia	Mar-2014	-55.5 ±1.3	5
Upper Silesian Basin	Poland	Jun-2013	-50.9 ± 1.2	8

495 **Table 1 $\delta^{13}CCH_4$ Signatures of all the coal mines and coal basins surveyed with the Picarro mobile system. Errors in**
496 **the $\delta^{13}C$ Signatures are calculated as 2 standard deviations**.

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-235, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 30 March 2016

Atmospheric Chemistry and Physics Discussions

Site	Coal Rank	$\delta^{13}C$ (‰)	Author	$\delta^{13}C$ (‰) measured in this study
Velenje Basin, Slovenia	Lignite	-71.8 to -43.4	Kanduč et al. (2015)	
Australia	From brown coal to low volatile bituminous coal.	-73.0 to -43.5	Smith et al. (1981)	-66.4 ±1.3 to -55.5 ±1.3
Powder River Basin, U.S.A.	Sub-bituminous coal	-68.4 to -59.5	Flores et al. (2008)	
Queensland Basin, Australia	From Sub-bituminous to high volatile bituminous	-57.3 to -54.2	Papendick et al. (2011)	
San Juan basin, New Mexico and Colorado	High-volatile Bituminous coal	-43.6 to -40.5	Rice et al. (1989)	
Elk Valley Coalfield, British Colombia	Bituminous coal	-65.4 to -51.8	Aravena et al. (2003)	
UK, Barnsley seam	Bituminous coal	-44.1	Hitchman et al. (a, b)	-48.8 ±0.3 to -45.4 ±0.2
Upper Silesian Coal Basin, Poland	Sub-bituminous coal to anthracite	-79.9 to -44.5	Kotarba and Rice (2001)	-50.9 ±0.6
Eastern China	Sub-bituminous to anthracite	-66.9 to -24.9	Dai et al. (1987)	
Ruhr basin, Germany	Bituminous coal, Anthracite	-37 (-60 to -14)	Deines (1980)	
Western Germany	High-volatile bituminous to anthracite.	-70.4 to -16.8	Colombo et al. (1970)	
Qinshui, China	Anthracite	-41.4 to -34.0	Qin et al. (2006)	
Wales, UK	Anthracite			-33.3 ±1.8 to -30.9 ±1.4

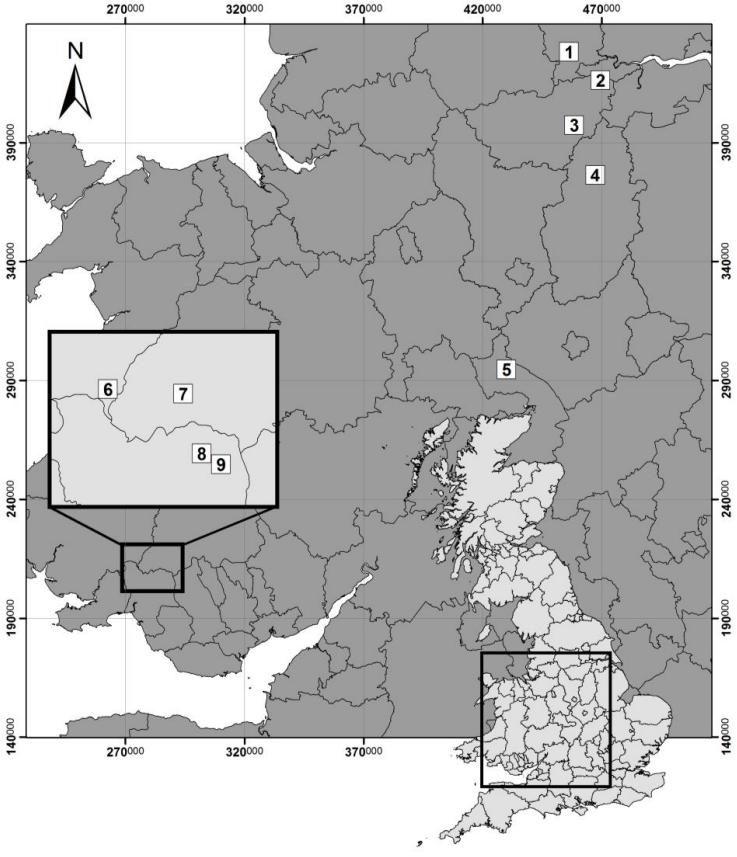
497 **Table 2 Literature isotope values obtained by methane samples from boreholes and coal seams. Errors are not included in the sources.**

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-235, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 30 March 2016

Atmospheric Chemistry and Physics Discussions

498

499



500

501 **Figure 1 Location of English coal mines (1 Kellingley, 2 Hatfield, 3 Maltby, 4 Thoresby, 5 Daw Mill) and Welsh coal**
502 **mines in the anthracite belt (6 Cwmllynfell, 7 Abercrave, 8 Unity, 9 Aberpergwm). Coordinates are displayed in the**
503 **National British Coordinates System.**

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-235, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 30 March 2016

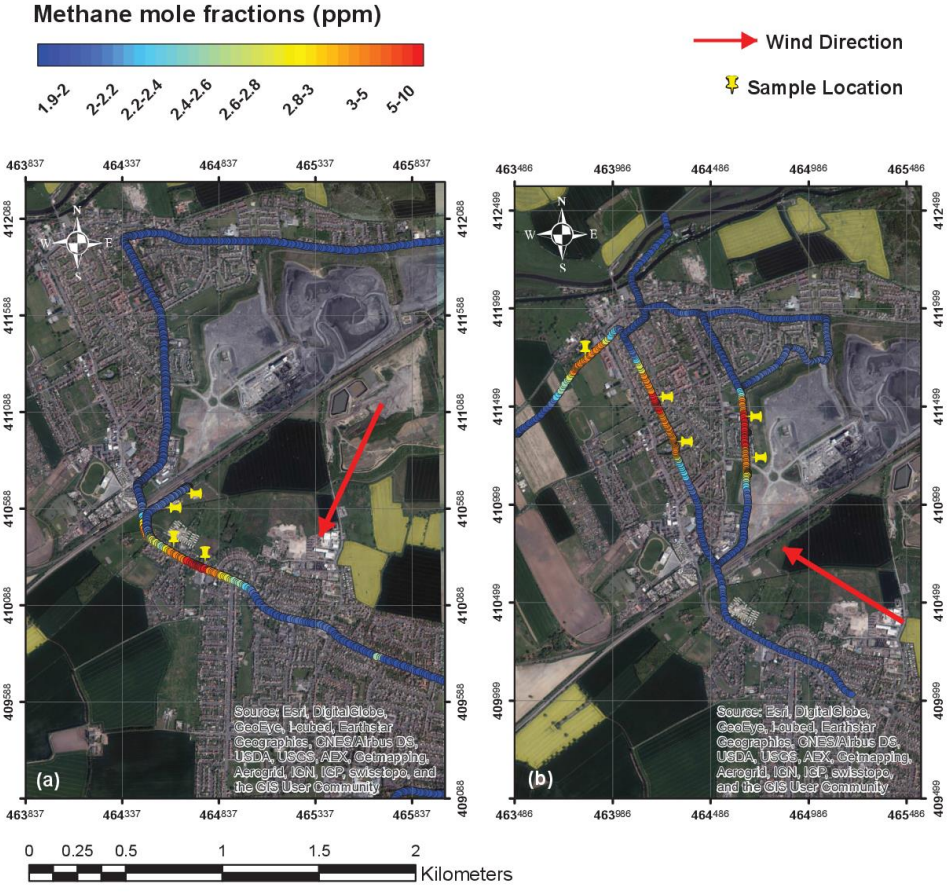


Figure 2 ArcGIS maps of methane mole fractions recorded downwind of Hatfield Colliery on July (a) and September 2013 (b). Grid Coordinates are displayed in the National British Coordinate System.

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-235, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 30 March 2016

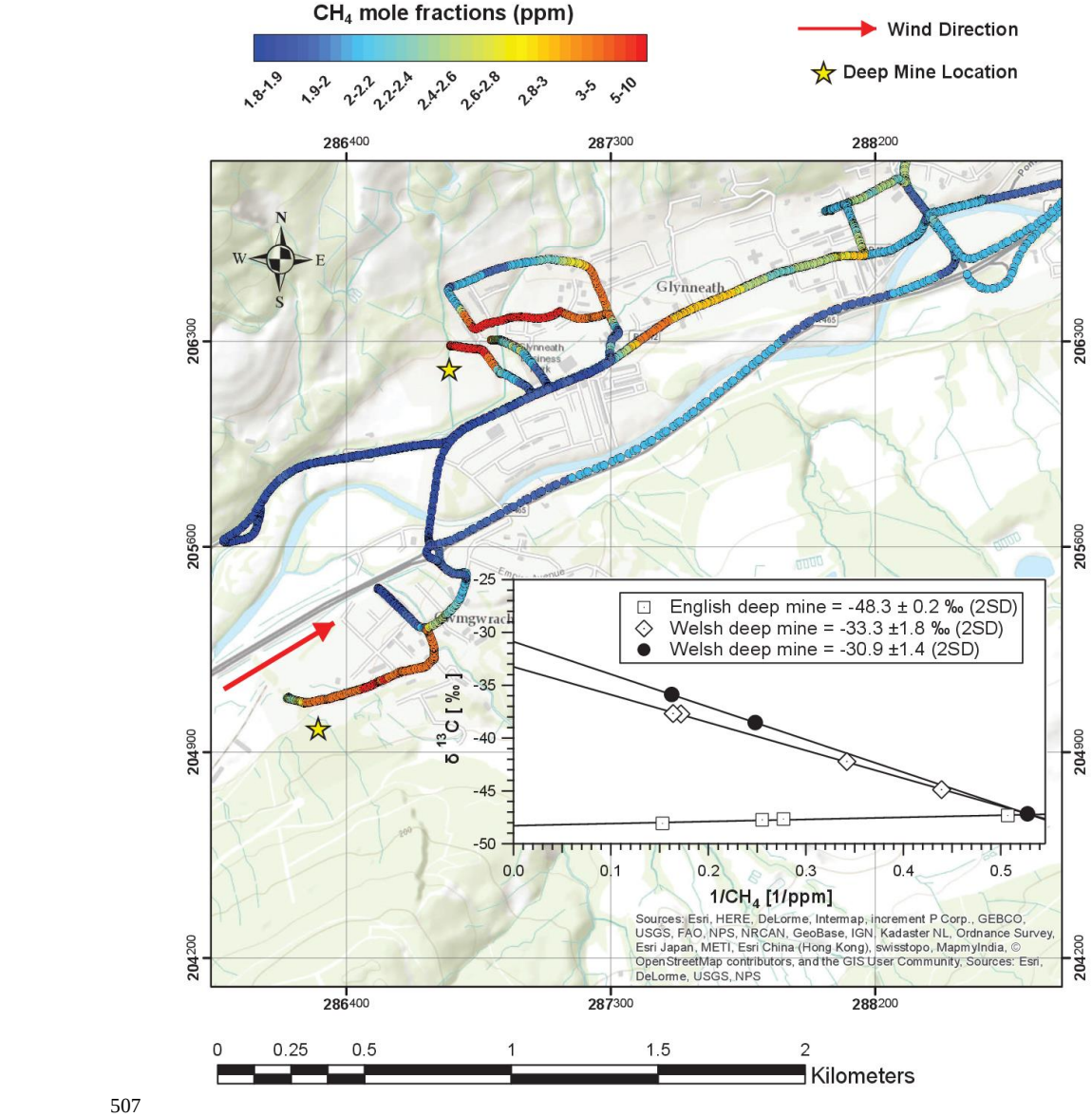


Figure 3 ArcGIS plot of methane mole fractions recorded in October 2013 next to Aberpergwm and Unity deep mines in Wales –yellow stars. Grid Coordinates are displayed in the National British Coordinate System. The embedded graph is the Keeling plot based on samples collected downwind of one English coal mine (Hatfield Colliery) and two Welsh deep mines (Aberpergwm and Unity Colliery). Errors on the y-axis are within 0.05 ‰ and on the x-axis 0.0001 ppm^{-1}, and are not noticeable on the graph.

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-235, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 30 March 2016

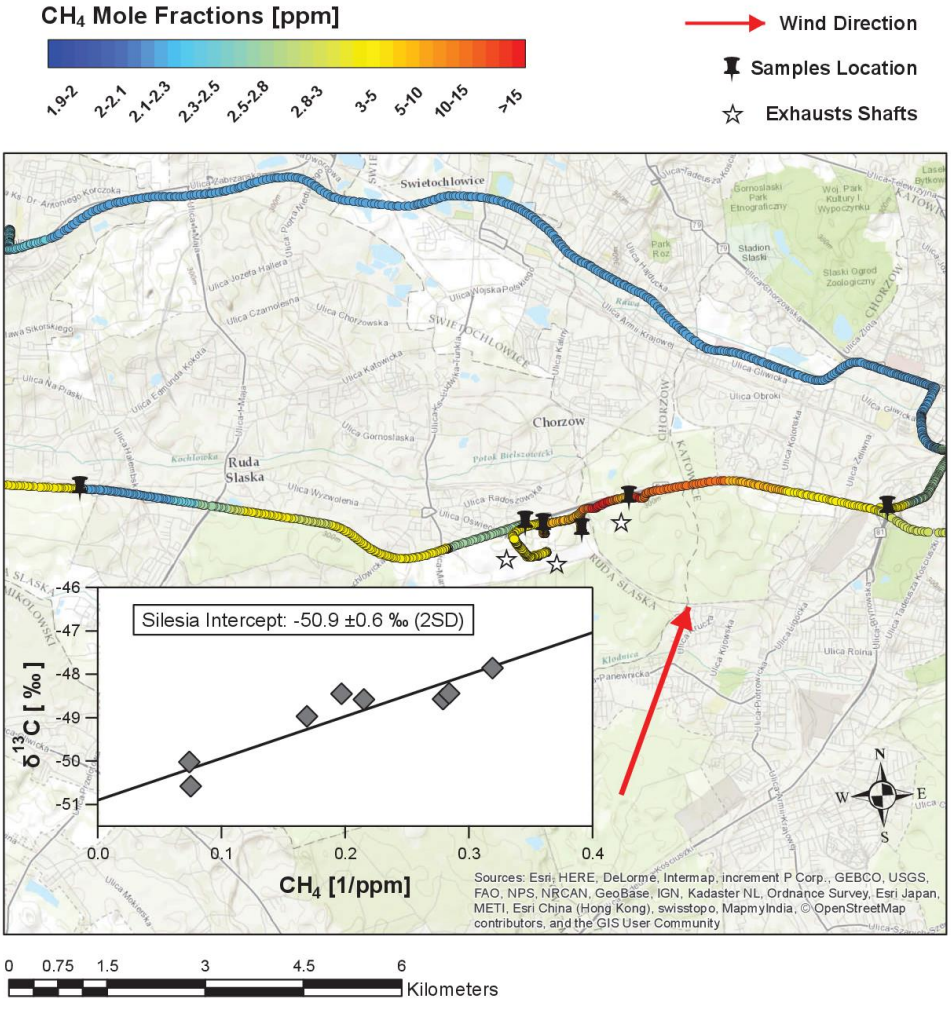
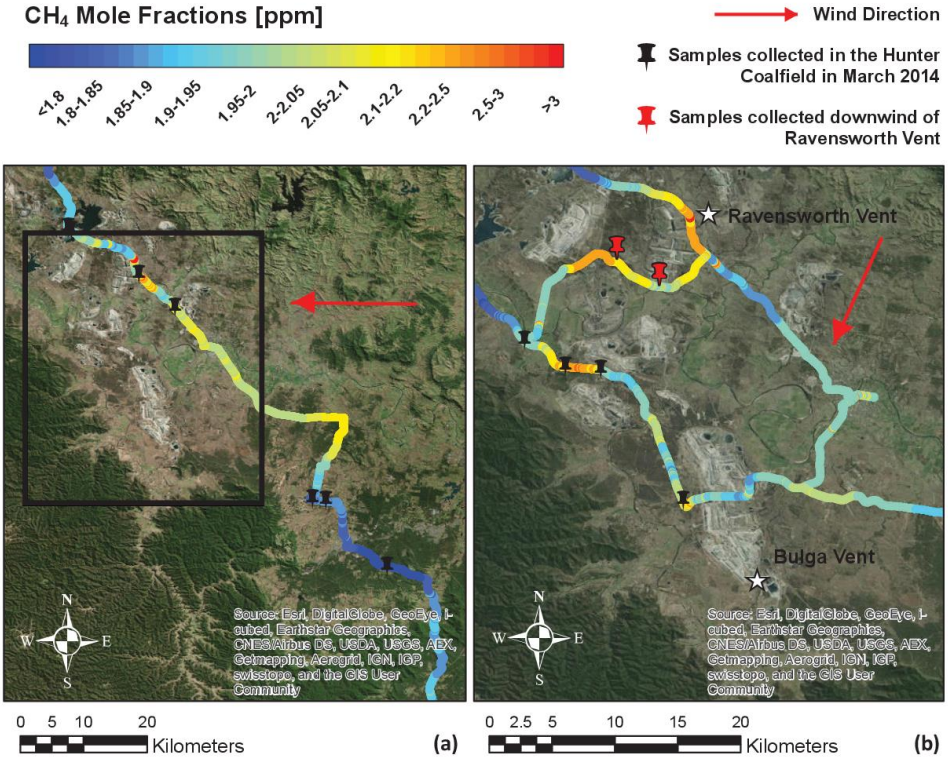


Figure 4 ArcGIS map of methane mole fractions recorded in the Upper Silesian basin in June 2013. White stars represent the KWK Wujek deep mine exhausts shafts. The embedded graph is the Keeling Plot based on samples collected downwind the KWK Wujek deep mine in June 2013. Errors on the y-axis are within 0.05 ‰ and on the x-axis 0.0001 ppm^{-1}, and are not noticeable on the graph.

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-235, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 30 March 2016

Atmospheric Chemistry and Physics Discussions



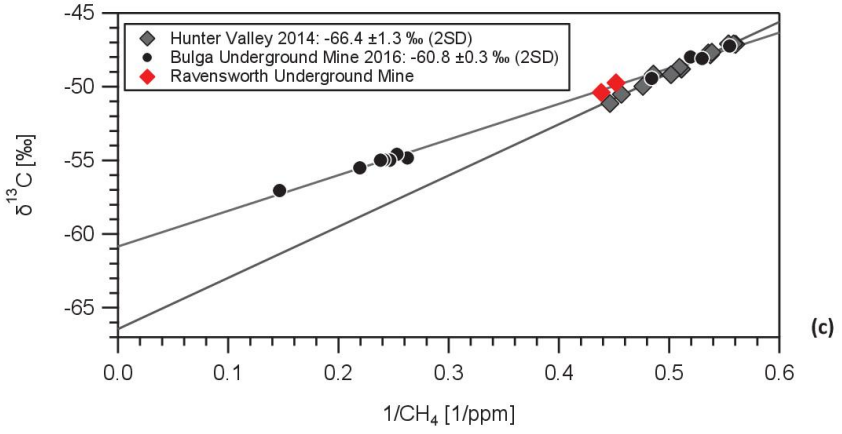


Figure 5 Methane mole fractions recorded in the Hunter Coalfield on 12th March 2014 (a) and during a more detailed survey of the area highlighted by the black square on 18th March 2014 (b). Keeling plot based on samples collected along the route in the Hunter Coalfield in March 2014 (grey and red markers) and near to ventilation in January 2016 (black markers) (c). Errors on the y-axis are within 0.05 ‰ and on the x-axis 0.0001 ppm^{-1}, and are not noticeable on the graph.