We thank the referees for the positive and very constructive comments. We have made the suggested changes accordingly and we have clarified the points that were raised in the comments below. The changes are highlighted in the second version of the manuscript that has been submitted.

Anonymous Referee #1

Minor comments:

- One useful information that I think is missing is the importance of the methane emissions from the regions studied. Some values are provided for the Silesian mines, and some estimates of the coal production is given for the Australian ones, but how much does that represent on the world’s scale, and how important are the British mines? Some literature references would suffice.

Literature values for the relative contribution to global emissions at global scale of the coal basins surveyed have been provided. See Lines 122-123, 144-145, 163-164.

- Table 2 could be extended with information on the mine type (open, deep, active, inactive since, . . . , perhaps also coordinates) and coal type.

Table 2 cannot be extended with such information, as most of the isotopic values reported in literature were obtained by sampling CBM (Coalbed Methane) wells across the basin and no specific mine is reported in these studies. Table 1 has been extended instead, to include information on type of mine and coal.

- I don’t find the map in Figure 1 very interesting: showing just the location of the mines without context information does not bring much to the paper. You could add some simplified geological map, or a map of the estimated distribution of coal mining methane emissions. Also, it is strange not to have the Australian and Silesian mines on a map!

Figure 1 has been removed, and Figure 2 includes now location of the Hunter Coalfield.

- Are you expecting any long or short-term variability of the d13C signature within one coal mine? If you were to re-do the sampling at a different season, or in a few years, would you expect any different result?

After closure coal mines continue to emit methane into the atmosphere through breakages in the coal seam or underground conduits (i.e. wells, vent pipes), until they are entirely flooded due to the groundwater inflow. The rate of methane released after the closure tends to decline over the years. Furthermore, the intrusion of water might trigger biogenic production of methane, which will make emissions lighter (13C-depleted). However, this assumption could not be tested, as surveys were carried out only once for most of coal mines, except for Maltby and Hatfield that were surveyed twice in the same year, for which the isotopic composition of methane emissions was really consistent.

- Many of the readers of this paper won’t be geologists. Although Section 1.1 is helpful and well written, it may help to provide some diagram showing the coal maturation process (when to expect which coal type), to be used as a reference when reading the following sections.

A diagram showing the coal maturation process has been added (Figure 1).
Textual comments:
- Line 26-27: what about Germany

The shift from coal to natural gas involved most of Europe (Line 28)

- Line 67: ... methanogenic path: acetoclastic ... (semicolon missing)

Corrected – see Line 68

- Line 255: the measurements are not significantly above background: is that because the mine emits less, or because the emissions are spread over a larger area in such an open mine?

In surface mines emissions are spread over a wider area as they are not focused on ventilation shafts.

- L298: why would the depth affect the d13C (other than the depth/coal-rank relation?)

The migration of methane towards the surface might lead to a fractionation effect as explained in the introduction. Line 304-305

- L310: There is a 20-35 permil difference between the Hunter Coalfield and the Wales mines, and you are talking of a 5-10 permil shift here ... 

The 5-10 permil difference refers to the difference between emissions from surface and deep mines within the same coalfield.

- L321-323: Do you have any information on the proportion of each coal/mine type in the world?

Added – see Line 334.
1. The results are currently presented in a very descriptive way, giving a lot of detail that makes part of the manuscript hard to read. I feel that a lot of this text, e.g. about locations, setting, would better fit into Table 1.

Most of this text has been moved into Table 1, which now includes the type of coal and the coal seam for each of the mine surveyed.

2. For each campaign, the authors report on the maximum CH4 signal and present detailed figures about the measured concentration distributions. However, this information is not evaluated in any way in the manuscript (e.g. by applying a dispersion model to estimate emission rates). The concentration signals largely depend on the actual meteorological situation at the time of sampling; therefore, a value of e.g. 10 ppm, which certainly confirms that there are indeed releases from the mining area, does not tell anything quantitative about the strength of the emission. I therefore feel that most of the figures are not required. Instead, in order to give the reader a feeling about the homogeneity of the Keeling-plot derived isotopic signature at each individual location and campaign, it would be more appropriate to present, perhaps in one multi-panel figure, individual Keeling plots from ALL campaigns, therewith showing the individual data points and regression lines in a standardized way (e.g. similar as Figure 5c, may be indicating also the maximum concentration in addition to 1/CH4).

A multi-panel figure of Keeling plots of all campaigns has been added (Figure 3). All mole fraction maps were removed except for the map of the Hunter Coalfield, as in this case the location of sampled emissions from deep mines are essential for the interpretation of results.

3. As also indicated by referee #1, I would strongly recommend to prepare an additional figure that summarizes all results, may be including the already published values from Table 2, in a diagram showing the relation between isotopic signature and coal type. This would be of tremendous help for people who want to use these results in their (modelling) studies. Currently, the last sentence in the Abstract leaves the reader/user somehow alone with the problem that rather no representative or at least improved input value for d13C of CH4 from coal is available. It looks to me that with the background knowledge of the authors, they could largely help to improve the current unsatisfactory situation of models using a value, which is definitely not correct/representative, no matter what the scale of the model is.

A diagram showing the process of coalification and the relation between isotopic signature and coal type has been added. (Figure 1).

4. The authors very often claim that secondary processes, e.g. bacterial methane production (or molecular diffusion) have occurred. However, this seems to be a (pure) speculation, because no measures of these processes are presented in the manuscript. These statements need more justification!

The theory of secondary processes leading to more 13C-depleted emissions is based on previous studies of many coal basins that are cited in the introductory section (Line 85-86; 100-103). The correlation of the isotopic composition with the biogenic production triggered by the incursion of meteoric water is an assumption based on measurements found in literature, but fits well with our results. However, such correlation will be presented in this paper only as a hypothetical source of variability in the isotopic composition of emissions from both surface and underground mines in the same coal basin (Line 299).
Specific comments:

Abstract:

*Line 16: Here you promise “... this study provides representative d13C-CH4 signatures to be used in regional and global models ...” is this really the case? See my general comment #3.*

The sentence has been changed (Line 17).

*Line 19: “Progression in coal rank and secondary ...” See my general comment #4*

Secondary processes are indeed suggested as a source of variability in the isotopic composition of coal-derived emissions, even though this assumption is based on the comparison of our results with those of other studies and not on direct measurements of secondary biogenic activities.

Introduction:

*Line 38: “calculated” should perhaps better read “reported”*

Changed – see Line 38

*Line 48: Here the reference “Hein et al., 1997” should be added*

Added – see Line 48

**Sec. 1.1. (there should be a section 1.2 if there is 1.1).**

In general, in particular for a non-specialist, it would be more clear if the processes during coalification (many million years ago) and what the situation is today, when these coal deposits are sampled, would be discussed here separately.

Section 1.1 has been incorporated in the introduction.

The current geological situation differs between coal basins and was included in the description of the single basins in section 2.1. Section 1.1. aims to provide a general understanding of those processes that affect the isotopic composition of methane, that can have occurred directly during the process of coalification itself or in a second stage.

*First paragraph: I am a bit confused here – is it already called “coalification” when fatty acids are converted to CH4 ?*

Yes, it is. The decomposition of organic matter that leads to the peat formation is the first stage of the coalification process.

*Line 87: should be “del13C” not “13C”*

Changed – see Line 88

*Line 108 – 112: Please clarify this sentence, do you mean ranges of 30/25 permil?*

Clarified – see Line 109

*Line 117-118: See general comment #4: Has the theory (interaction with water) really been tested in the current study?*
No and this has been removed from Line 118.

Section 2:

Line 143: Should read “km²” (not km)
Corrected – see Line 144

Line 164: the word “origin” is missing
Added – see Line 167

Line 183: Is it important to mention here the precision of d13C-CO2 measurements?
Precision has been removed. Line 186

Line 185: Please add “CH4”
Added – see Line 188

Line 197: Please give reference to the ArcGIS software (if these maps are not removed, as suggested above).
The source of the GIS map has been added in the caption of Figure 2

Line 204: Please be precise: It is not possible to measure mole fractions of emissions
The sentence has been rephrased. Line 210-215

Section 3: A lot of this detailed information may better go into a table.
Much of this information is now included in Table 1

Line 248: Do you really mean +33.3 permil?
Thank you – corrected on Line 252

Line 255: Please be precise: an emission cannot be “above background”
Corrected – see Line 259

Lines 292 ff and line 310: please see general comment #4
As the likelihood of biogenic secondary methane cannot be proven in the present study the statements on Lines 298, 318 are now only suggestions.
Carbon isotopic signature of coal-derived methane emissions to atmosphere: from coalification to alteration

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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 δ¹³CCH₄ signatures of methane emissions from specific coalfields 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 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 (crushing 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 reported without an error.
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 (Hein et al., 1997; 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 δ¹³CCH₄ 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 δ¹³CCH₄ 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 δ¹³CCH₄ signatures and coal rank and mining setting, and to provide representative ¹³C signatures to be used in atmospheric models in order to produce more accurate methane emission estimates for the coal exploitation sector.

**Process of coalification and parameters affecting the ¹³C 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, CO₂, H₂, NH₄⁺, HS⁻, long chain fatty acids) are metabolised by fermentative archaea, which produce methane via two methanogenic pathways: acetoclastic reaction or CO₂ reduction (Whiticar, 1999). Different pathways lead to diverse δ¹³CCH₄ isotopic signatures - methane from acetate is ¹³C enriched relative to methane from CO₂ 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 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) (see Figure 1).

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 ($\delta^{13}$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 (variation of ~30‰) and thermal maturation stage (variation of ~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.
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 theory 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

Coal-derived methane emissions in the UK are estimated at 66 Gg in 2013 (naei.defra.gov.uk), representing 0.4% of global emissions from mining activity (US EPA, 2012). The major coalfields of England and Wales belong to the same stage in the regional stratigraphy of northwest Europe (Westphalian Stage, Upper Carboniferous). Mining has ceased in many areas. Of those remaining some are located in South Yorkshire (Hatfield and Maltby collieries), where 50% of the coalfield's output came from the Barnsley seam, which includes soft coal overlaying a semi-anthracite 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 km² (Jureczka and Kotas, 1995), contributing 1.3% to global coal-derived methane emissions (US EPA, 2012). Emissions from the Silesian Region of Poland are estimated to be 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 CO₂ in the
coal seams at the top of the Carboniferous sequence accumulated under clay deposition in the Miocene (Kędzior, 2009).

### 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) - and methane emissions are estimated to contribute 4.6% to global coal-derived methane (US EPA, 2012). 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 origin (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 ($\delta^{13}$CCH$_4$) of bag samples was measured in the greenhouse gas laboratory at RHUL (Royal Holloway University of London) in triplicate to high precision ($\pm0.05$ ‰) by continuous flow gas
chromatography isotope ratio mass spectrometry (CF GC-IRMS) (Fisher et al., 2006). CH\textsubscript{4} and CO\textsubscript{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 δ\textsuperscript{13}C\textsubscript{CH\textsubscript{4}} signature for each emission plume was calculated using the Keeling plot approach, according to which the δ\textsuperscript{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 δ\textsuperscript{13}C\textsubscript{CH\textsubscript{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

Methane plumes from Hatfield colliery, one of the few UK deep mines still open at the time of this study, were investigated detected during surveys on 10\textsuperscript{th} July and 26\textsuperscript{th} 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 m 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 (up to 7 ppm). 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, 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 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 ground Thoresby mine, in Nottinghamshire, approached 5 ppm. Nine 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. During the sampling of methane plumes from Daw Mill colliery, 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. Indeed, the estimated methane content of the coal seam exploited in Daw Mill colliery, closed in March 2013, is low (typically about 1.7 m³/tonne) and, furthermore, a ventilation system is implemented, so that the coal mine methane potential is curtailed (Drake, 1983). 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. 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 1.5 km 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 13C enriched relative to all English collieries.

3.2.2 Opencast Mines

The Picarro mobile system was driven around opencast mines at Cwmllynfell and Abercave, 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. 13C 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 Radoszow, downwind of the KWK Wujek deep mine shafts (white stars in Fig. 4). Methane mole fractions in the range of 3-5 ppm were measured in the majority of the area of Katowice and over 20 ppm mole fractions were detected when transecting the plume originating from the exhaust shafts, which confirms the high level of methane that the mine contains. A source signature of -50.9 ± 0.6 ‰ (2SD) was calculated by Keeling plot analysis, which is consistent with the values obtained for the deep mined English bituminous coal.

3.4 Australia: Hunter Coalfield

On 12th and 18th March 2014 12 samples in total were collected along the route in the Hunter Coalfield, where the bituminous coal strata of the Sydney Basin are extracted both in opencast and underground mines. The methane plume width was in the range of 70 km (Fig. 2). A maximum mole fraction of 13.5 ppm was measured near a vent shaft associated with the Ravensworth underground mine (see white star in Fig. 2b), which exploits the Lemington, Pikes Gully, Lidell (Upper and Middle) and Barret Seams, the deepest of which is the Barret seam, with a maximum overburden depth of ~350 m (GSS Environmental, 2012). The source signature calculated by the Keeling plot analysis based on all the samples collected during both surveys (grey markers in Fig. 5a, 5c) is -66.4 ±1.3 ‰ (2SD) and this signature likely includes a mixture of methane derived both from underground and opencast mines. Two samples collected downwind of the Ravensworth ventilation shaft (red pushpins in Fig. 2b) fall off the Keeling plot trend for March 2014 and are not included in the calculation, because they are likely dominated by methane from the vent shaft and are not representative of the regional mixed isotopic signature.

In January 2016, 10 samples were collected downwind of a ventilation fan in the Bulga mine (second white star in Fig. 2b), aerating underground workings in the Blakefield South Seam, ranging from 130 to 510 m in depth (Bulga Underground Operation mining, 2015). The Keeling plot for these (black circles in Fig. 5a, 5c) indicates a δ13C source signature of -60.8 ±0.3 ‰ (2SD). The samples collected next to the Ravensworth ventilation shaft in 2014 fit on this Keeling plot, suggesting that the δ13CCH4 isotopic signature of emissions from underground workings in the Hunter Coalfield is consistent.

3.5 New representative δ13CCH4 isotopic signatures for coal-derived methane

The δ13CCH4 isotopic values for coal have been found to be characteristic of single basins, but general assumptions can be made to characterise coal mines worldwide. Table 2 provides the literature δ13CCH4 isotopic values characteristic of specific coal basins. The isotopic signatures of emissions from English bituminous coal are consistent with the range of -49 to -31 ‰ suggested by Colombo et al. (1970) for in situ coal bed methane in the Ruhr basin in Germany, which contains the most important German bituminous coal of Upper Carboniferous age and low volatile anthracite (Thomas, 2002). Progression in coal rank might explain the value of -50 ‰ for emissions from English underground mined bituminous coal and -30 ‰ for anthracite deep mines, followed by a 5-10 ‰ 13C-depletion likely caused by the incursion of meteoric water in the basin and the subsequent production
of secondary biogenic methane (Scott et al., 1994), resulting in -40 ‰ for methane plumes from Welsh open-cut anthracite mines.

The link between coal rank and δ13CCH4 isotopic signature is appreciable in the study of UK coal mines, but differences in the 13C/12C isotopic ratio within the same coal sequences can be ascribed to other parameters, such as the depth at which the coal is mined - as methane migrates towards the surface, diffusion or microbial fractionation may occur, especially if there is water ingress (see section 1.4). In particular, emissions from Thoresby are more 13C depleted than those measured around Maltby, although both mines exploit the Parkgate seam (see Table 1), meaning that different isotopic signatures cannot be entirely linked to the coal rank. Biogenic methane 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 are 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 δ13C signatures for coalbed methane emissions from the Upper Silesian basin are highly variable, with the most 13C 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 δ13C 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 δ13C 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 global coal reserves and the dominance of bituminous coal (1% and 53% respectively) (World Coal Institute, 2015), 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 δ¹³C 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 δ¹³C 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.

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<table>
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<tr>
<th>Sampling Site</th>
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<th>Coal seam</th>
<th>Coal Type</th>
<th>Mine Type</th>
<th>Sampling Date</th>
<th>$\delta^{13}$C Signatures [‰]</th>
<th>Samples used in Keeling plots</th>
<th>number of Keeling plots</th>
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<tbody>
<tr>
<td>Kellingley Colliery</td>
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<td>Beeston and Silkstone coal seams</td>
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<td>$-48.3 \pm 0.2$</td>
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<td>$-51.4 \pm 0.2$</td>
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<td>Surface Mine</td>
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<td>$-41.2 \pm 0.9$</td>
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<tr>
<td>Aberpergwm Colliery</td>
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<td>$-33.3 \pm 1.8$</td>
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<td>Bulga Colliery</td>
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<td>Blakefield South Seam, from 130 to 510 m in depth (Bulga Underground Operation mining, 2015)</td>
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<td>Upper Silesian Basin</td>
<td>Poland</td>
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<td>Jun-2013</td>
<td>$-50.9 \pm 1.2$</td>
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Table 1 $\delta^{13}$CH$_4$ signatures of all the coal mines and coal basins surveyed with the Picarro mobile system. Errors in the $\delta^{13}$C signatures are calculated as 2 standard deviations. *Only the coalfield is indicated, as the name of the seams exploited could not be retrieved.
<table>
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<tr>
<th>Site</th>
<th>Coal Rank</th>
<th>$\delta^{13}$C (‰)</th>
<th>Author</th>
<th>$\delta^{13}$C (‰) measured in this study</th>
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<td>-66.4 ±1.3 to -55.5 ±1.3</td>
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<td>Elk Valley Coalfield, Canada</td>
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<td>-65.4 to -51.8</td>
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<td>Bowen basin, Australia</td>
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Table 2 Literature isotope values obtained by methane samples CBM (Coalbed Methane) wells across the basin. Errors are not included in the sources.
Figure 1 Coal maturation process.
Figure 3: 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.
Figure 4 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.
Figure 5. 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, and are not noticeable on the graph.
Figure 2 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). Source: Esri, DigitalGlobe, GeoEye, i-cubed, Earthstar Geographics, CNES/Airbus DS, USDA, USGS, AEX, Getmapping, Aerogrid, IGN, IGP, swisstopo and the GIS User Community.
Figure 3 Keeling Plots based on samples collected around English and Welsh coal mine (a), coal mines in Poland (b) and Australia (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.