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

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25 1 Introduction

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2 3

Methane emissions from the energy sector have been driven in recent years by the impact of a shift from coal to 26 27 natural gas, in the US and Europe, whereas in China coal production has increased in this century. Currently, the 28 atmospheric methane burden is rising rapidly (Nisbet et al., 2014), but the extent to which shifts in coal production 29 contribute to this rise is not known. Coalbed methane emissions into the atmosphere are poorly characterised, as 30 they are dispersed over large areas and continue even after the mines' closure (IPCC, 2006). Methane is emitted 31 in coal processing (crashing and pulverisation) and during the initial removal of the overburden; it can be diluted 32 and emitted through ventilation shafts in underground coal mines, or directly emitted to atmosphere from open-33 cut coal mining, where releases may occur as a result of deterioration of the coal seam. 34 For the UN Framework Convention on Climate Change, national emissions are estimated by a "bottom-up" 35 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.

37 carbon content of coal) (U.S. EPA, 2013). These modelled estimates are often 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., 39
- 40 2006; Locatelli et al., 2013). However, this top-down approach provides the total amount of methane emissions
- 41 into the atmosphere, which has to be distributed among the different methane sources in order to quantify each
- 42 source contribution.

43 For methane emissions from fossil fuels (coal and natural gas), the source partitioning is mainly "bottom-up",

44 based on energy use statistics and local inventories, which might be highly uncertain. Conversely, the "top-down" 45

study of the carbon isotopic composition of methane, which is indicative of the methane origin, provides a valuable

46 constraint on the budget appraisal, allowing different sources in a source mix to be distinguished and their

47 individual strength to be evaluated (Hein et al., 1997; Liptay et al., 1998; Lowry et al., 2001; Townsend-Small et 48 al., 2012).

49 Measurements of methane mole fractions can be complemented in atmospheric models by typical δ^{13} CCH₄ 50 signatures of the main methane sources in order to estimate global and regional methane emissions and assess 51 emissions scenarios throughout the past years (Fung et al., 1991; Miller, 2004; Whiticar and Schaefer, 2007; 52 Bousquet et al., 2006; Monteil et al., 2011; Mikaloff Fletcher et al., 2014). However, even though isotopic values 53 are fairly distinctive for specific methanogenic processes, the variety of production pathways and local 54 environmental conditions that discriminate the methane formation process leads to a wide range of δ^{13} CCH₄ 55 values. The global isotopic range for coal is very large, from -80 to -17 ‰ (Rice, 1993), but it can be narrowed 56 down when a specific basin is studied. While there are several studies of isotopic composition of methane 57 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 58 59 isotopic characterisation of methane emissions from coal mines in Europe.

The purpose of this study was to determine links between δ^{13} CCH₄ signatures and coal rank and mining setting, 60

61 and to provide representative ¹³C signatures to be used in atmospheric models in order to produce more accurate

62 methane emission estimates for the coal exploitation sector.

Process of coalification and parameters affecting the δ^{13} C signature of methane emissions 63

64 The process of coalification involves both biochemical and geochemical reactions. The vegetal matter firstly 65 decays anaerobically under water; the simple molecules derived from initial decomposition (i.e. acetate, CO₂, H₂, 66 NH₄⁺, HS⁻, long chain fatty acids) are metabolised by fermentative archaea, which produce methane via two 67 methanogenic paths: acetoclastic reaction or CO₂ reduction (Whiticar, 1999). Different pathways lead to diverse δ^{13} CCH₄ isotopic signatures - methane from acetate is 13 C enriched relative to methane from CO₂ reduction, 68 69 ranging from -65 ‰ to -50 ‰ and -110 ‰ to -50 ‰ respectively (Levin et al., 1993; Waldron et al., 1998). With

70 increasing burial and temperatures, coal is subjected to thermal maturation, which implicates more geochemical

- 71 changes.
- 72 As coalification proceeds, the carbon content increases, accompanied by a relative depletion in volatile

73 compounds, such as hydrogen and oxygen, emitted in the form of water, methane, carbon dioxide and higher

- 74 hydrocarbons through decarboxylation and dehydration reactions (Stach and Murchison, 1982). At higher degrees
- 75 of coalification and temperature, the liquid hydrocarbons formed in previous stages are thermally cracked to
- 76 methane, increasing the amount of methane produced (Faiz and Hendry, 2006). Peat and brown coal represent the
- 77 first stage of the coalification process. The vertical pressure exerted by accumulating sediments converts peat into

78 lignite. The intensification of the pressure and heat results in the transition from lignite to bituminous coal, and

reventually 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).

- 86 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 87 88 to biogenic methane, as methanogens preferentially use the lightest isotopes due to the lower bond energy (Rice, 89 1993). Intermediate isotopic compositions of methane might reflect a mixing between microbial and thermogenic 90 gases or secondary processes. Indeed, many controlling factors co-drive the fractionation process, and several 91 contentions about their leverage still persist in literature. Deines (1980) asserts that no significant trend is observed 92 in the isotopic signature of methane in relation to the degree of coalification. Conversely, Chung et al. (1979) 93 observed that the composition of the parent material does affect the isotopic composition of the methane 94 accumulated. While the link between isotopic composition and coal rank is not that straightforward, studies carried 95 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 96 97 to contain relatively isotopically lighter methane when compared to those at greater depths. In the presence of 98 intrusions of meteoric water, secondary biogenic methane, isotopically lighter, can be generated and mixed with 99 the thermogenic gas previously produced. Colombo et al. (1966) documented a distinct depth correlation in the 100 Ruhr Basin coal in Germany, with methane becoming more ¹³C-depleted towards the surface zone, independently 101 from coalification patterns. This tendency can be explained either by bacterial methanogenesis, or by secondary 102 processes such as absorption-desorption of methane. Also Scott (2002), in a study about coal seams in the Bowen 103 Basin, Australia, ascribes the progressive methane ¹³C-enrichment with depth to the meteoritic recharge in the 104 shallowest seams, associated with a higher bacterial activity and a preferential stripping of ¹³C-CH₄ by water flow. The migration of methane from the primary zone as a consequence of local pressure release can affect the isotopic 105 106 composition, since ¹²CH₄ diffuses and desorbs more readily than ¹³CH₄ (Deines, 1980), but the fractionation effect 107 due to migration is less than 1 ‰ (Fuex, 1980). A much larger variation in the isotopic composition is associated 108 with different methanogenic pathways (variation of ~30 ‰) and thermal maturation stage (variation of ~25 ‰) (Clayton, 1998). The measurement of Deuterium, coupled with δ^{13} CCH₄ values, would help to distinguish the 109 110 pathways of secondary biogenic methane generation, acetoclastic reactions or CO₂ reduction (Faiz and Hendry, 111 2006), but such distinction is beyond the scope of this study.
- 112 Overall, the δ^{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.

115 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
- 117 due to coal rank.

118 2 Material and Methods

119 2.1 Coal basins investigated and type of coal exploited

120 2.1.1 English and Welsh coal mines

121 Coal-derived methane emissions in the UK are estimated at 66 Gg in 2013 (naei.defra.gov.uk), representing 0.4%
122 of global emissions from mining activity (US EPA, 2012). The major coalfields of England and Wales belong to
123 the same stage in the regional stratigraphy of northwest Europe (Westphalian Stage, Upper Carboniferous).
124 Mining has ceased in many areas. Of those remaining some are located in South Yorkshire (Hatfield and Maltby
125 collieries), where 50% of the coalfield's output came from the Barnsley seam, which includes soft coal overlaying
126 a semi-anthracitic coal and bituminous coal in the bottom portion.
127 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-

134 underground.org.uk/pit/geology.shtml).

135 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 136 137 in Europe, with an area of ~7400 km² (Jureczka and Kotas, 1995), contributing 1.3% to global coal-derived 138 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 139 140 associated with gas deposits of both thermogenic and microbial origin, and the methane content and spatial 141 distribution are coal rank related (Kędzior, 2009) - i.e. the sorption capacity of coal in the basin is found to increase 142 with coal rank. Most of the methane generated during the bituminous stage in the coalification process has escaped 143 from the coal source following basin uplift during Paleogene (Kotarba and Rice, 2001) and diffused through 144 fractures and faults occurring in tectonic zones. The late-stage gas generated by microbial reduction of CO₂ in the 145 coal seams at the top of the Carboniferous sequence accumulated under clay deposition in the Miocene (Kędzior, 2009). 146

147 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

155 in the UK - the total coal production for the Hunter Coalfield was 123.63 Mt in 2011, of which 88.24 Mt was 156 saleable (State of New South Wales, 2013) - and methane emissions are estimated to contribute 4.6% to global 157 coal-derived methane (US EPA, 2012). Several studies have attested to the dispersion of thermogenic methane 158 formed at higher degrees of coalification (i.e. high temperatures and pressures) during the uplift and the 159 subsequent erosion of the basin, followed by the replenishment of the unsaturated basin with more recently formed 160 methane of biogenic origin (Faiz and Hendry, 2006; Burra, 2010). Gas emplacement is related to sorption capacity 161 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 characterised by higher gas 162 163 content and enhanced permeability than the northern area, with a large potential for methane production, mainly 164 biogenic (Pinetown, 2014).

165 2.2 Sampling and measurement methodology

For isotopic characterisation of the methane sources, integrated methane emissions were assessed through 166 167 detection of the offsite downwind plume. In fact, even when emissions are focused on defined locations, such as 168 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 169 170 and Australia the mobile system described by Zazzeri et al. (2015) has been implemented. The system utilises a 171 Picarro G2301 CRDS (Cavity Ring-Down Spectroscopy) within the survey vehicle, for continuous CH₄ and CO₂ 172 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 173 174 methane plume outline to be displayed in real time on a Google Earth platform during the survey to direct plume 175 sampling. When the plume was encountered, the vehicle was stopped and air samples collected in 3L Tedlar bags, 176 using a diaphragm pump connected to the air inlet. Samples were taken at different locations along the plume 177 transect in order to obtain a wide range of methane mole fractions and isotopic signatures in the collected air. The 178 Upper Silesian basin was surveyed with a Picarro 2101-i measuring continuous CO₂ and CH₄ mole fractions and 179 δ^{13} CCO₂ isotopic ratio. Samples were collected on site for analysis of δ^{13} CCH₄ isotopic ratio.

180 The carbon isotopic ratio (δ^{13} CCH₄) of bag samples was measured in the greenhouse gas laboratory at RHUL 181 (Royal Holloway University of London) in triplicate to high precision (± 0.05 ‰) by continuous flow gas 182 chromatography isotope ratio mass spectrometry (CF GC-IRMS) (Fisher et al., 2006). CH₄ and CO₂ mole fractions 183 of samples were measured independently in the laboratory with a Picarro G1301 CRDS analyser, calibrated 184 against the NOAA (National Oceanic and Atmospheric Administration) WMO-2004A and WMO-X2007 185 reference scales respectively. The δ^{13} CCH₄ signature for each emission plume was calculated using the Keeling plot approach, according to which the δ^{13} C isotopic composition of samples and the inverse of the relative mole 186 187 fractions have a linear relationship, whose intercept represents the isotopic signature of the source (Pataki et al., 188 2003). Source signatures were provided with the relative uncertainty, computed by the BCES (Bivariate 189 Correlated Errors and intrinsic Scatter) estimator (Akritas and Bershady, 1996), which accounts for correlated 190 errors between two variables and calculates the error on the slope and intercept of the best interpolation line. Mole 191 fraction data and co-located coordinates were used to map the mole fraction variability using the ArcGIS software.

192 **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^{13} CCH₄ signatures of all the coal mines and

199 coal basins surveyed with the Picarro mobile system.

200 **3.1 English coal mines**

- 201 Methane plumes from Hatfield colliery, one of the few UK deep mines still open at the time of this study, were 202 detected during surveys on 10^{th} July and 26^{th} September 2013. The methane desorbed from coal heaps within the 203 colliery area and methane vented from shafts might explain the relatively high mole fractions measured (up to 7 204 ppm). Keeling plots based on the samples collected (4 on the first and 5 on the second survey) give intercept 205 values of -48.3 ±0.2 and -48.8 ±0.3 ‰ (2SD).
- Maltby colliery is one of the largest and deepest mines in England and was closed in March 2013. The coal mine methane was extracted and used for electricity, but over 3 ppm mole fractions were detected in ambient air during
- 207 methane was extracted and used for electricity, but over 3 ppm mole fractions were detected in ambient air during 208 both surveys in July and September 2013, giving evidence of methane releases from the recovery system and
- 209 ventilation shafts. The Keeling plot intercepts based on the samples collected in July and in September are -45.9
- ± 0.3 and $-45.4 \pm 0.2 \%$ (2SD) respectively, $\sim 3 \%$ heavier than the isotopic source signatures calculated for Hatfield
- 211 Colliery on the same sampling days. Both source signatures are in good agreement with the value of -44.1 ‰
- 212 observed for desorbed methane of the Barnsley coal seam, in a study conducted by Hitchman et al. (1990b).
- 213 Kellingley colliery is a still open deep mine situated in North Yorkshire. A surface drainage plant for electrical
- 214 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 10^{th} July 2013. The Keeling plot analysis of the samples collected indicates a source signature of -46.5 ±0.3 ‰ (2SD).
- 217 The highest mole fractions detected around Thoresby mine, in Nottinghamshire, approached 5 ppm. Nine samples
- 218 were collected, giving a source signature of -51.2 ± 0.3 % (2SD).
- 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 $-51.4 \pm 0.2 \%$ (2SD).

225 3.2 Welsh coal mines

226 Methane emissions from coal mines in Wales were sampled in order to characterise methane releases from a

- 227 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).

229 **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
- 232 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
- 235 % result from the Keeling plots based on samples collected respectively near Aberpergwm and Unity colliery,
- both highly ¹³C enriched relative to all English collieries.

237 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, even though the methane mole fractions recorded downwind of the mines were 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 characterising

244 Welsh deep anthracite mines.

245 **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 246 247 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, 248 downwind of the KWK Wujek deep mine shafts. Methane mole fractions in the range of 3-5 ppm were measured 249 250 in the majority of the area of Katowice and over 20 ppm mole fractions were detected when transecting the plume 251 originating from the exhaust shafts, which confirms the high level of methane that the mine contains. A source 252 signature of -50.9 ± 0.6 % (2SD) was calculated by Keeling plot analysis, which is consistent with the values 253 obtained for the deep mined English bituminous coal.

254 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 255 256 bituminous coal strata of the Sydney Basin are extracted both in opencast and underground mines. The methane 257 plume width was in the range of 70 km (Fig. 1). 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. 1b). The source signature 258 259 calculated by the Keeling plot analysis based on all the samples collected during both surveys (grey markers in Fig. 2c) is -66.4 ±1.3 ‰ (2SD) and this signature likely includes a mixture of methane derived both from 260 261 underground and opencast mines. Two samples collected downwind of the Ravensworth ventilation shaft (red 262 pushpins in Fig. 1b) fall off the Keeling plot trend for March 2014 and are not included in the calculation, because 263 they are likely dominated by methane from the vent shaft and are not representative of the regional mixed isotopic signature. 264

- 265 In January 2016, 10 samples were collected downwind of a ventilation fan in the Bulga mine (second white star
- 266 in Fig. 1b). The Keeling plot for these (black circles in Fig. 2c) indicates a δ^{13} C source signature of -60.8 ±0.3 ‰
- 267 (2SD). The samples collected next to the Ravensworth ventilation shaft in 2014 fit on this Keeling plot, suggesting
- 268 that the δ^{13} CCH₄ isotopic signature of emissions from underground works in the Hunter Coalfield is consistent.

269 **3.5** New representative δ^{13} CCH₄ isotopic signatures for coal-derived methane

270 The δ^{13} CCH₄ isotopic values for coal have been found to be characteristic of single basins, but general assumptions 271 can be made to characterise coal mines worldwide. Table 2 provides the literature δ^{13} CCH₄ isotopic values 272 characteristic of specific coal basins. The isotopic signatures of emissions from English bituminous coal are 273 consistent with the range of -49 to -31 ‰ suggested by Colombo et al. (1970) for in situ coal bed methane in the 274 Ruhr basin in Germany, which contains the most important German bituminous coal of Upper Carboniferous age 275 and low volatile anthracite (Thomas, 2002). Progression in coal rank might explain the value of -50 ‰ for 276 emissions from English underground mined bituminous coal and -30 ‰ for anthracite deep mines, followed by a 277 5-10 \% ¹³C-depletion likely caused by the incursion of meteoric water in the basin and the subsequent production 278 of secondary biogenic methane (Scott et al., 1994), resulting in -40 ‰ for methane plumes from Welsh open-cut 279 anthracite mines.

- 280 The link between coal rank and δ^{13} CCH₄ isotopic signature is appreciable in the study of UK coal mines, but differences in the ${}^{13}C/{}^{12}C$ isotopic ratio within the same coal sequences could be ascribed to other parameters, 281 282 such as the depth at which the coal is mined - as methane migrates towards the surface, diffusion or microbial 283 fractionation may occur, especially if there is water ingress (see section 1). In particular, emissions from Thoresby 284 are more ¹³C depleted than those measured around Maltby, although both mines exploit the Parkgate seam (see 285 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 286 formed during the coalification process. 287
- 207 formed during the coameation 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.
- 294 The difference in the source isotopic signature for methane emissions between the two types of mining both in
- the Hunter Coalfield (from -61 to -66 ‰) and in Wales (from -31 to -41 ‰) reflects the isotopic shift of 5-10 ‰
- that might be attributed to the occurrence of secondary biogenic methane.
- 297 The δ^{13} C signatures for coalbed methane emissions from the Upper Silesian basin are highly variable, with the
- 298 most ¹³C depleted methane associated with diffusion processes or secondary microbial methane generation
- 299 (Kotarba and Rice, 2001), but the value of -51 ‰ measured for methane emissions from the KWK Wujek deep
- 300 mine is consistent with the value of -50 ‰ inferred for emissions from English bituminous coal extracted in
- 301 underground mines.
- 302 Figure 3 shows the δ^{13} CCH₄ signatures ranges for each coal type, based on the results of this study and isotopic
- 303 values found in literature.

304 4 Conclusions

305 By measuring the isotopic signatures of methane plumes from a representative spread of coal types and depths, 306 we show that the δ^{13} C isotopic value to be included in regional and global atmospheric models for the estimate of 307 methane emissions from the coal sector must be chosen according to coal rank and type of mining (opencast or 308 underground). For low resolution methane modelling studies, our measurements suggest an averaged value of -65 ‰ for bituminous coal exploited in open cast mines and of -55 ‰ or less negative, in deep mines, whereas 309 310 values of -40 ‰ and -30 ‰ can be assigned to anthracite opencast and deep mines respectively. However, the 311 ranges of isotopic signatures of methane emitted from each of these categories are wide (see Fig. 3), and the 312 isotopic ratios of methane from Chinese coal mines is little studied. Further measurements would be required in 313 other coal mining areas, especially China, to determine appropriate values for global modelling of methane 314 isotopes.

Global methane budget models that incorporate isotopes have used a δ^{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 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

- 321 should be region or nation specific, as greater detail is needed given the wide global variation. The assignment of
- 322 an incorrect global mean, or a correct global mean but inappropriate for regional scale modelling, might lead to
- 323 incorrect emissions estimates or source apportionment. The new scheme gives the possibility for an educated
- 324 estimate of the δ^{13} C signature of emission to atmosphere to be made for an individual coal basin or nation, given
- 325 information on the type of coal being mined and the method of extraction.
- 326 In conclusion, high-precision measurements of δ^{13} C in plumes of methane emitted to atmosphere from a range of
- 327 coal mining activities have been used to constrain the isotopic range for specific ranks of coal and mine type,
- 328 offering more representative isotopic signatures for use in methane budget assessment at regional and global
- 329 scales.

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Sampling Site	Country	Coal seam	Coal Type	Mine Type	Sampling Date	δ ¹³ C Signatures [‰]	Samples number used in Keeling plots
Kellingley Colliery (active)	North Yorkshire/UK	Beeston and Silkstone coal seams	Bituminous coal	Deep Mine	Sept-2013	-46.5 ±0.3	5
Maltby Colliery (closed since March 2013)	South Yorkshire/UK	Barnsley and Parkgate seams, up to 991 m in depth (McEvoy et al., 2006)	Bituminous coal	Deep Mine	Jul-2013 Sept-2013	-45.9 ±0.3 -45.4 ±0.2	3 4
Hatfield Colliery (active)	South Yorkshire/UK	Barnsley (~401-465 m), Dunsil (414-376 m) High Hazel (~313- 284 m) seams (Hill, 2001)	Bituminous coal	Deep Mine	Jul-2013 Sept-2013	-48.3 ±0.2 -48.8 ±0.3	4 5
Thoresby Colliery (active)	Nottinghamshire/UK	Parkgate seam, ~650 m in depth	Bituminous coal	Deep Mine	Nov-2013	-51.2 ±0.3	9
Daw Mill Colliery (closed since March 2013)	Warwickshire/UK	Warwickshire Thick seam, from 500 to 1000 m in depth.	Bituminous coal	Deep Mine	Nov-2013	-51.4 ±0.2	4
Cwmllynfell Colliery (active)	Wales/UK	South Wales Coalfield*	Anthracite	Surface Mine	Oct-2013	-41.2 ±0.9	4
Abercrave Colliery (active)	Wales/UK	South Wales Coalfield	Anthracite	Surface Mine	Oct-2013	-41.4 ±0.5	5
Aberpergwm Colliery (closed since December	Wales/UK	South Wales Coalfield	Anthracite	Deep Mine	Oct-2013	-33.3 ±1.8	5
Unity Colliery (active)	Wales/UK	South Wales Coalfield	Anthracite	Deep Mine	Oct-2013	-30.9 ± 1.4	3
Hunter Coalfield (active)	Australia	Whittingham Coal Measures	Bituminous coal	Surface and Deep Mines	Mar-2014	-66.4 ±1.3	12
Bulga Colliery (active)	Australia	Blakefield South Seam, from 130 to 510 m in depth (Bulga Underground Operation mining, 2015)	Bituminous coal	Deep Mine	Jan-2016	-60.8 ±0.3	10
Upper Silesian Basin (active)	Poland	502 (590-634m) and 510 seam (763-812m)	Bituminous coal	Deep Mine	Jun-2013	-50.9 ± 1.2	8

512 Table 1 δ¹³CCH₄ signatures of all the coal mines and coal basins surveyed with the Picarro mobile system. Errors in the δ¹³C signatures are calculated as 2 standard deviations. *Only

513 the coalfield is indicated, as the name of the seams exploited could not be retrieved.

Site	Coal Rank	δ ¹³ C (‰)	Author	δ^{13} C (‰) measured in this study
Powder River Basin, U.S.A	Sub-bituminous coal	-79.5 to -56.8	Bates et al. (2011	
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
Velenje Basin, Slovenia	Lignite	-71.8 to -43.4	Kanduč et al. (2015)	
Powder River Basin, U.S.A.	Sub-bituminous coal	-68.4 to -59.5	Flores et al. (2008)	
Illinois Basin, U.S.A.	Bituminous coal	-66.6 to -56.8	Strąpoć et al. (2007)	
Elk Valley Coalfield, Canada	Bituminous coal	-65.4 to -51.8	Aravena et al. (2003)	
Bowen basin, Australia	Bituminous coal	-62.2 to -48	Kinnon et al. (2010)	
Queensland Basin, Australia	From Sub-bituminous to high volatile bituminous	-57.3 to -54.2	Papendick et al. (2011)	
Tara region, Queensland Basin, Australia	From Sub-bituminous to high volatile	-56.7	Maher et al. (2014)	
San Juan basin, New Mexico and Colorado	High-volatile Bituminous coal	-43.6 to -40.5	Rice et al. (1989)	
Michigan Basin, U.S.A.	Bituminous coal	-56 to -47	Martini et al. (1998)	
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)	
Eastern Ordos Basin, China	Bituminous Coal	-42.3 to -39.8	Guo et al. (2012)	
South-eastern Ordos Basin, China	Low-volatile bituminous and semi- anthracite	-48.1 to -32.8	Yao et al. (2013)	
Qinshui, China	Anthracite	-41.4 to -34.0	Qin et al. (2006)	
Qinshui Basin, China	Semi-anthracite to Anthracite	-36.7 to -26.6	Su et al. (2005)	
Sichuan Basin, China	Bituminous to anthracite	-36.9 to -30.8	Dai et al. (2012) Xiong et al. (2004)	
Wales, UK	Anthracite			-33.3 ± 1.8 to -30.9 ± 1.4

514 Table 2 Literature isotope values obtained by methane samples CBM (Coalbed Methane) wells across the basin. Errors are not included in the sources.





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516 Figure 1 Methane mole fractions recorded in the Hunter Coalfield on 12th March 2014 (a) and during a more detailed

- 517 survey of the area highlighted by the black square on 18th March 2014 (b). Source: Esri, DigitalGlobe, GeoEye, icubed, Earthstar Geographics, CNES/Airbus DS, USDA, USGS, AEX, Getmapping, Aerogrid, IGN, IGP, swisstopo
- 518 519 and the GIS User Community.



- 520 Figure 2 Keeling Plots based on samples collected around English and Welsh coal mines (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
- 522 graph.
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Increase in Depth of Burial and Temperature



- 525 Figure 3 Coal maturation process. The δ^{13} CCH₄ signatures ranges are based on the literature values in Table 2 and on 526 the results of this study.
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