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### Paper:

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## RESEARCH REVIEW

## Towards a global assessment of pyrogenic carbon from vegetation fires

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## Abstract

The production of pyrogenic carbon (PyC; a continuum of organic carbon (C) ranging from partially charred biomass and charcoal to soot) is a widely acknowledged C sink, with the latest estimates indicating that ~50% of the PyC produced by vegetation fires potentially sequesters C over centuries. Nevertheless, the quantitative importance of PyC in the global C balance remains contentious, and therefore, PyC is rarely considered in global C cycle and climate studies. Here we examine the robustness of existing evidence and identify the main research gaps in the production, fluxes and fate of PyC from vegetation fires. Much of the previous work on PyC production has focused on selected components of total PyC generated in vegetation fires, likely leading to underestimates. We suggest that global PyC production could be in the range of 116–385 Tg C yr<sup>-1</sup>, that is ~0.2–0.6% of the annual terrestrial net primary production. According to our estimations, atmospheric emissions of soot/black C might be a smaller fraction of total PyC (<2%) than previously reported. Research on the fate of PyC in the environment has mainly focused on its degradation pathways, and its accumulation and resilience either *in situ* (surface soils) or in ultimate sinks (marine sediments). Off-site transport, transformation and PyC storage in intermediate pools are often overlooked, which could explain the fate of a substantial fraction of the PyC mobilized annually. We propose new research directions addressing gaps in the global PyC cycle to fully understand the importance of the products of burning in global C cycle dynamics.

**Keywords:** biochar, black carbon, carbon accounting, carbon emissions, carbon sequestration, charcoal, dissolved organic carbon, erosion, pyrogenic organic matter, wildfire

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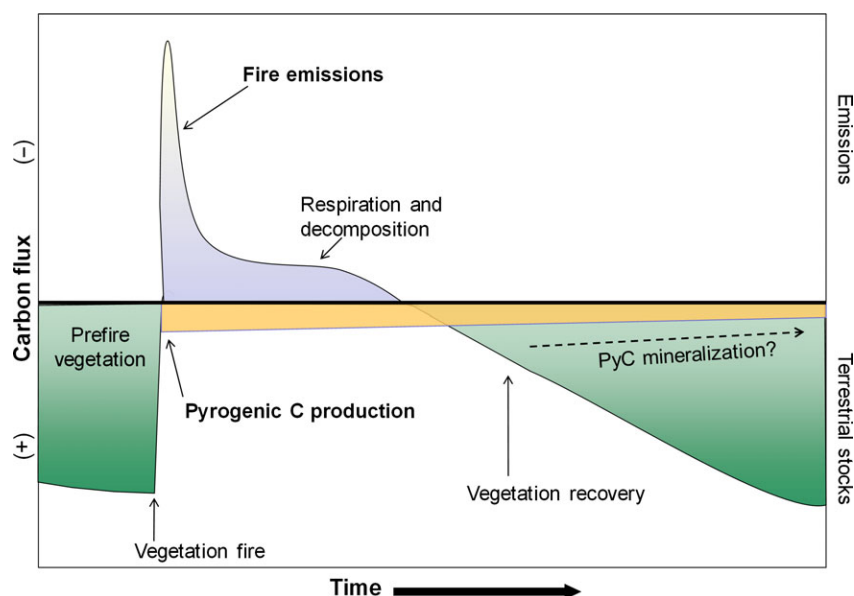
## Introduction

Vegetation fires affect 300–460 Mha globally per year (Randerson *et al.*, 2012; Giglio *et al.*, 2013), emitting 1.6–2.8 Gt carbon (C) to the atmosphere, the equivalent of 25–30% of the current annual C emissions from fossil fuel consumption (Van Der Werf *et al.*, 2010; Boden *et al.*, 2012). Over the longer term (i.e. decades), however, vegetation fires are widely considered as ‘net zero C emission events’ because C emissions from fires are balanced by C uptake by regenerating vegetation (excluding deforestation and peatland fires) (Bowman *et al.*, 2009; Van Der Werf *et al.*, 2010). This zero C emis-

sion scenario is potentially flawed, however, as it does not consider the role of pyrogenic C (PyC; Fig. 1).

Incomplete combustion during fires transforms part of the fuel C into PyC. The high diversity of fuel materials as well as the wide range of combustion conditions, especially in vegetation fires, do not allow PyC to be defined as a distinct chemical component, but instead as the organic C fraction of the whole range of pyrogenic organic materials from partially charred vegetal biomass and charcoal to soot (Goldberg, 1985; Schmidt & Noack, 2000). PyC is therefore not a homogenous organic C pool, but includes a broad continuum ranging from biolabile depolymerization products to highly resistant condensation products. Charring mainly induces condensation reactions, with the resulting

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**Fig. 1** Carbon fluxes driven by fire over time and through a whole fire cycle (i.e. complete recovery of vegetation). Emissions to the atmosphere are represented as negative fluxes. The timescale ranges from a few years to many decades depending on the ecosystem type and the severity of the fire (modified from Conard & Solomon, 2009).

polycyclic aromatic structures providing increased resistance against degradation. However, under low charring temperatures ( $\sim 250^\circ\text{C}$ ), depolymerization and dehydration of part of the biomass also occurs, which makes some of the PyC highly water soluble and biodegradable (Norwood *et al.*, 2013; Myers-Pigg *et al.*, 2015).

Overall, the pyrogenic process mostly confers the charred materials a longer mean residence time in the environment compared to their unburnt precursors (Schmidt *et al.*, 2011; DeLuca & Boisvenue, 2012; Singh *et al.*, 2014; Naisse *et al.*, 2015a). Therefore, a large fraction of the PyC continuum can be considered a C sink on a decadal/centennial timescale (Fig. 1) (Bird *et al.*, 2015). The enhanced resistance of PyC to degradation, for example, underpins the production of biochar (PyC intentionally produced for soil amendment) and its addition to soils as one of the most viable global approaches in offsetting C emissions to the atmosphere (Woolf *et al.*, 2010; Jeffery *et al.*, 2015). However, PyC produced naturally in vegetation fires is usually not considered in C budget and global warming investigations (Lehmann *et al.*, 2008; Le Quere *et al.*, 2009), and the net role of PyC in the global C cycle is thus not well elucidated. The main reason for this is the lack of robust knowledge on PyC production and degradation, fluxes and residence time in the environment (Masiello, 2004; DeLuca & Aplet, 2008; Schmidt *et al.*, 2011). As small changes in C cycle dynamics can have large effects in global climate change scenarios, there is an urgent need for improvement of the representation of

the terrestrial C cycle in climate and integrated assessment models (Moss *et al.*, 2010).

In this research review we discuss the identification and quantification of PyC from vegetation fires and review current knowledge and uncertainties on its production, degradation, mobilization and long-term fate in the environment. We provide updated estimates of the current global PyC production, fluxes and pools and highlight the main research gaps in the PyC cycle. This review concludes with suggestions of new research directions aimed at achieving a more complete and integrated understanding of the role of PyC from vegetation fires in the global C cycle.

### PyC identification and quantification

Identification and quantification of PyC in the different environmental matrixes (soils, sediments, air and waters) are essential for addressing the role of PyC in C budgets. However, these are difficult tasks as PyC is not a distinct chemical component but a continuum of C-rich solid organic materials. This represents a key challenge when studying PyC because the methodology used determines the overall amount and characteristics of the PyC quantified, and yields can also change depending on the environmental matrix the PyC is being isolated from (Schmidt *et al.*, 2001; Hammes *et al.*, 2007; Roth *et al.*, 2012).

A clear definition of the specific window of the PyC continuum that each study addresses is essential to avoid uncertainty when comparing, extrapolating or

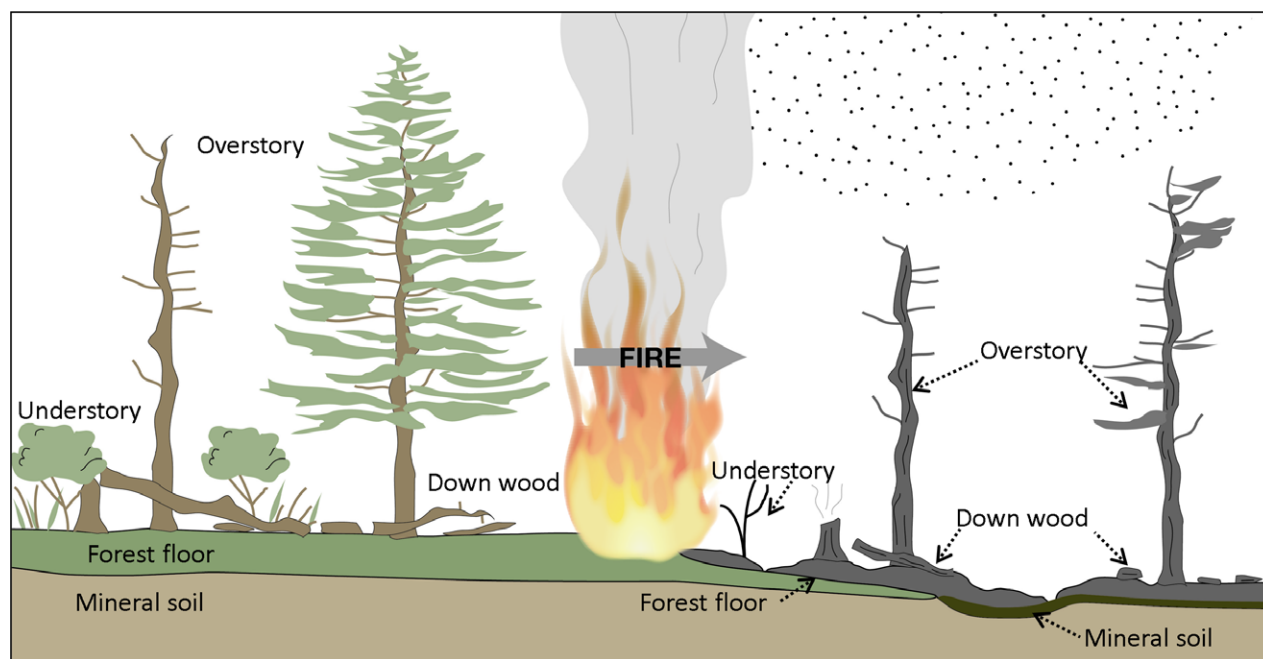
scaling up data from individual studies. Some components of the PyC continuum are measured through mutually exclusive techniques. For example, low-temperature pyrolysis can transform cellulose into anhydrosugars (e.g. levoglucosan) which have no aromatic rings, are detected by gas chromatography and can have a very short mean residence time (MRT) (Norwood *et al.*, 2013). Conversely, fire can also produce soot, which is characterized by a stable, condensed and aromatic chemical structure and is typically measured either by nuclear magnetic resonance or by thermal analyses (Hammes *et al.*, 2007). While these two forms of C (anhydrosugars and soot) are produced exclusively by pyrolysis, they have very different environmental pathways and MRTs. Neither can be treated as behaving in a way representative of the entire PyC pool. Currently, no technique representatively captures the entire PyC pool. Therefore, not only methodologies capturing a wide range of the PyC continuum are fundamental (e.g. visual identification, Santín *et al.*, 2015), but also those characterizing specific parts of the PyC continuum are necessary to understand how much of the total PyC is significant as a C sink in the long term (e.g. stable polycyclic aromatic C – SPAC, McBeath *et al.*, 2015).

### PyC production from vegetation fires

To elucidate the PyC cycle, we need first to know how much is formed during fire. To allow inclusion of PyC

production into C emission and budget models, a complete prefire fuel quantification is needed so that PyC production can be reported as a proportion of the fuel affected by fire (Fig. 2). However, comprehensive fuel data are rarely available, particularly for wildfires (Keane, 2012), and assessments often exclude relevant fuel components such as the canopy or woody debris (de Groot *et al.*, 2007; Possell *et al.*, 2015).

A detailed prefire fuel quantification is normally only available for small-scale experimental or prescribed fires (e.g. Alexis *et al.*, 2007), which are usually not representative of wildfire conditions. In the low-intensity burning conditions typical of prescribed fires (Certini, 2005), burning efficiency is generally low, only a small part of the fuel is exposed to thermal degradation, and overall little PyC is produced (Graça *et al.*, 1999; Schmidt & Noack, 2000). In contrast, high-intensity fires have a higher burning efficiency (Campbell *et al.*, 2007) and affect a greater proportion of the fuel available. The degree to which fire intensity translates into complete fuel combustion vs. PyC generation is governed by complex factors related to intrinsic fuel properties (e.g. density and composition), extrinsic fuel properties (e.g. arrangement, moisture and loads) and burning conditions (e.g. fire weather, oxygen availability and burning duration) (Brewer *et al.*, 2013). As a result, PyC produced in low-intensity experimental or prescribed fires may not be representative of PyC produced during the often more intense wildfires.



**Fig. 2** Quantification of pyrogenic C produced in a fire with respect to C affected by fire requires: (i) pre- and postfire unburnt fuel estimations and (ii) determination of pyrogenic C emitted to the atmosphere and the remaining in all fuel components (for this example of a forest fire: overstory, understory, down wood, forest floor (or litter) and mineral soil).

In addition to a complete prefire fuel assessment, the whole range of PyC materials generated needs to be considered and quantified (Fig. 2). However, PyC investigations are primarily divided into (i) on-site PyC (i.e. generated and remaining on-site immediately after the fire) and (ii) PyC emitted to the atmosphere within the smoke. To the best of our knowledge, no investigation has fully quantified simultaneously the production of on-site and emitted PyC during fire.

#### *On-site PyC*

Much of the PyC generated during a vegetation fire initially remains on-site mainly as PyC within (i) soil; (ii) the ash layer on the ground; and (iii) charred plant tissue (charcoal) on standing vegetation and downed wood (Scott, 2000) (Fig. 2).

(i) *PyC in soil*: When examining fire effects on soil PyC stocks, it is important to consider that most vegetation fires do not result in temperatures exceeding the minimum temperature required to initiate the charring process a few millimetres below the mineral soil surface ( $\sim 200^\circ\text{C}$ ; González-Pérez *et al.*, 2004). Therefore, much of the PyC found in fire-affected soils does not originate from *in situ* pyrolysed soil organic matter, but from PyC produced from the burning of the litter and above-ground vegetation (Bodí *et al.*, 2014; Boot *et al.*, 2015). This PyC can subsequently be incorporated into the soil profile through processes such as bioturbation and freeze–thaw or into newly forming horizons in areas where eroded sediments accumulate (Gavin, 2003; Wilkinson *et al.*, 2009). Notable exceptions are burning of tree roots, organic soils and peats, where smouldering combustion produces PyC *in situ* at depth (Kane *et al.*, 2010).

Studies investigating soil PyC often address exclusively either visually detectable charcoal or one of the chemically defined PyC components (i.e. quantified through thermal, chemical or spectroscopic techniques), reflecting a research focus, respectively, on either fire history (e.g. Ohlson *et al.*, 2011) or on mineral soil/biogeochemical issues (e.g. Czimczik *et al.*, 2005). Either approach misses part of the soil PyC spectrum. Studies focused on visual identification of charcoal pieces miss the PyC in the finest fraction of the soil, which can contain the largest PyC soil stock (Brodowski *et al.*, 2006). Those quantifying chemically defined PyC in soils usually examine the soil fraction  $<2\text{ mm}$  and therefore exclude macroscopic charcoal  $>2\text{ mm}$ , which can account for up to 60–90% of the visually detected charcoal (Ohlson & Tryterud, 2000; Nocentini *et al.*, 2010). This mutual neglect tends not only to underestimate the total amount of PyC, but also to overrepresent

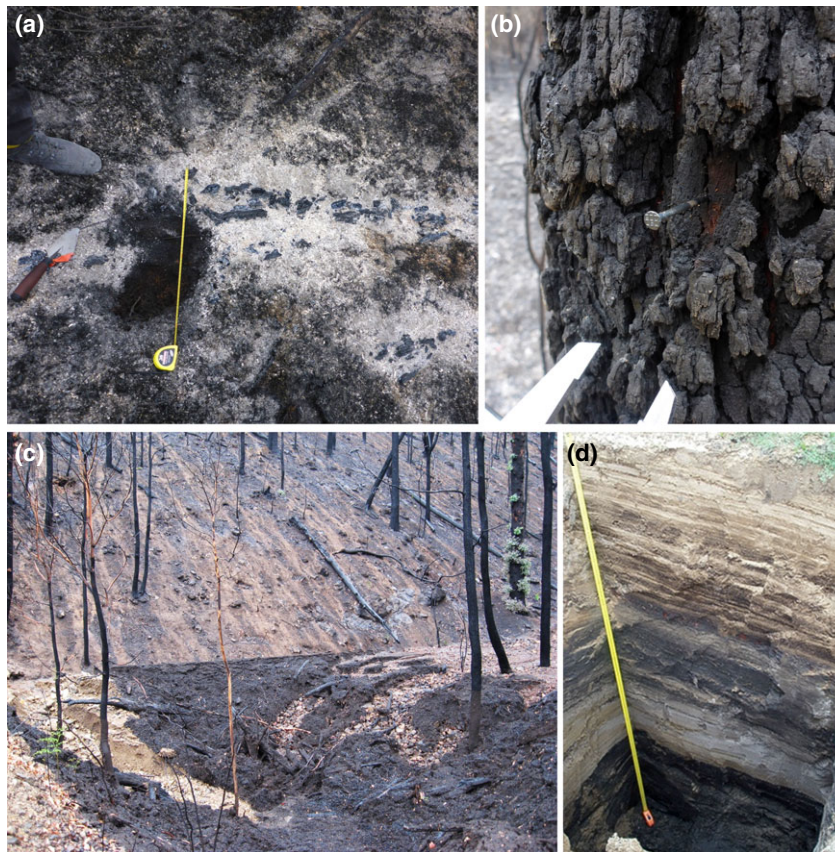
specific fractions and their characteristics. For instance, the biggest pieces of PyC are dominated by wood-derived charcoal, while the smaller fractions typically originate from needles, leaves, herbs and organic topsoil material. These fractions differ in their physical and chemical nature. Wood-derived PyC is generally more recalcitrant than the chemically more reactive nonwoody PyC fractions (Hilscher *et al.*, 2009; Nocentini *et al.*, 2010; De la Rosa & Knicker, 2011). They also differ in their mobility and MRTs, with large wood-derived charcoal particles prone to being incorporated into the soil and to persist there for millennia (Gavin, 2003; Ohlson *et al.*, 2009; de Lafontaine & Asselin, 2011; de Lafontaine *et al.*, 2011).

(ii) *PyC in the ash layer* is rarely considered in PyC inventories (Fig. 3a). A major reason for the neglect of ash is its often rapid redistribution within, and removal from, burnt sites by wind and water erosion, which often occurs before the commencement of postfire field investigations (Cerdeña & Doerr, 2008; Bodí *et al.*, 2014). Depending on formation conditions, ash can contain substantial amounts of PyC and should be included in PyC inventories (Forbes *et al.*, 2006; Bodí *et al.*, 2014). For example, Santín *et al.* (2012) estimated that  $6\text{--}8\text{ t PyC ha}^{-1}$  was transferred from burnt fuels to the ash layer during the catastrophic 2009 ‘Black Saturday’ wildfires in Australia.

(iii) *PyC on standing vegetation and downed wood* is another important, although frequently neglected, pool of PyC (Figs 2 and 3b). Tinker & Knight (2000) estimated  $6.4\text{ t ha}^{-1}$  of charcoal produced from coarse woody debris ( $>7.5\text{ cm}$  diameter) during a crown fire in a conifer forest in western USA. Donato *et al.* (2009) reported  $0.3\text{--}0.6\text{ t ha}^{-1}$  PyC generated from downed wood in a stand-replacing conifer forest fire in north-western USA. Santín *et al.* (2015) quantified a PyC production of  $1.9 \pm 0.2\text{ t ha}^{-1}$  in down wood and  $2.5 \pm 1.3\text{ t ha}^{-1}$  in bark of standing trees during a conifer forest fire in the boreal Canada. The importance of these aboveground PyC pools deserves further attention as most PyC compounds originating from woody materials are expected to have long MRTs (Ohlson *et al.*, 2009). Also large woody charcoal pieces may act as slow-release sources of PyC. The location of this woody PyC (e.g. standing timber vs. down wood) also needs consideration as it affects its persistence and mobilization.

In conclusion, for a comprehensive quantification of PyC produced, all the components discussed above need to be accounted for (Fig. 2). Santín *et al.* (2015) summarized the approaches used in 31 previous studies quantifying on-site PyC production and concluded that most inventories are incomplete and tend to underestimate total PyC production. Santín *et al.* (2015)





**Fig. 3** Examples of pyrogenic C (PyC) in the environment: (a) PyC derived mainly from burnt forest floor and down wood (boreal forest, NW Canada). Note the PyC-rich ash layer below the white ash; (b) bark-derived PyC on standing tree (dry eucalypt forest, SE Australia); (c) water erosion and redeposition of PyC-enriched sediments after a severe wildfire (wet eucalypt forest, SE Australia); (d) PyC-rich layers in reservoir sediments (excavated at low water level, pit depth 2 m, SE Australia).

quantified the complete range of PyC components found on-site immediately after a boreal forest fire and estimated that over a quarter of the C affected by fire was converted to PyC. This is well above the ~1–5% commonly considered (Preston & Schmidt, 2006) and highlights the importance of including the complete range of PyC in quantitative studies.

#### *PyC emitted to the atmosphere*

Part of the PyC produced during fire is emitted to the atmosphere within smoke. This 'atmospheric PyC' is situated at the smallest size end of the PyC spectrum (<1–2  $\mu\text{m}$ ) and is chemically the most recalcitrant (Bird & Ascough, 2012). Within the atmospheric sciences, it is usually referred to as black C (BC), elemental C or soot (for a detailed discussion of these terms see Buseck *et al.*, 2012). It is important to recognize that during fire, some macroscopic PyC (particles >120–150  $\mu\text{m}$ ) can also become airborne, but these are commonly not considered in PyC emissions as its mobilization is generally limited to the vicinity of the fire (Oris *et al.*, 2014).

At regional and global scales, estimations of emitted PyC based on bottom-up inventories are much lower than concentrations estimated from atmospheric observations (Kaiser *et al.*, 2012). Given that open biomass burning is not only one of the largest contributors to global PyC emissions, but also the one presenting the highest uncertainties (Bond *et al.*, 2004), a better understanding of the PyC emissions to the atmosphere during vegetation fires is essential. Bond *et al.* (2013) point to emissions factors (i.e. C emitted with respect to fuel combusted) representing the dominant uncertainty of the role of PyC aerosols in the global climate.

There is a major gap between 'atmospheric' and 'terrestrial' PyC research. On the one hand, research on fire emissions quantifies PyC emitted to the atmosphere, but overlooks the PyC remaining on-site (Ottmar, 2014); investigations on fire emissions generally assume that all burnt C is either volatilized as gases or contained in the emitted aerosols (Akagi *et al.*, 2011). On the other hand, research focusing on 'terrestrial' PyC (i.e. remaining on-site) often assumes that >80% of PyC produced remains on-site and <20% is emitted to the

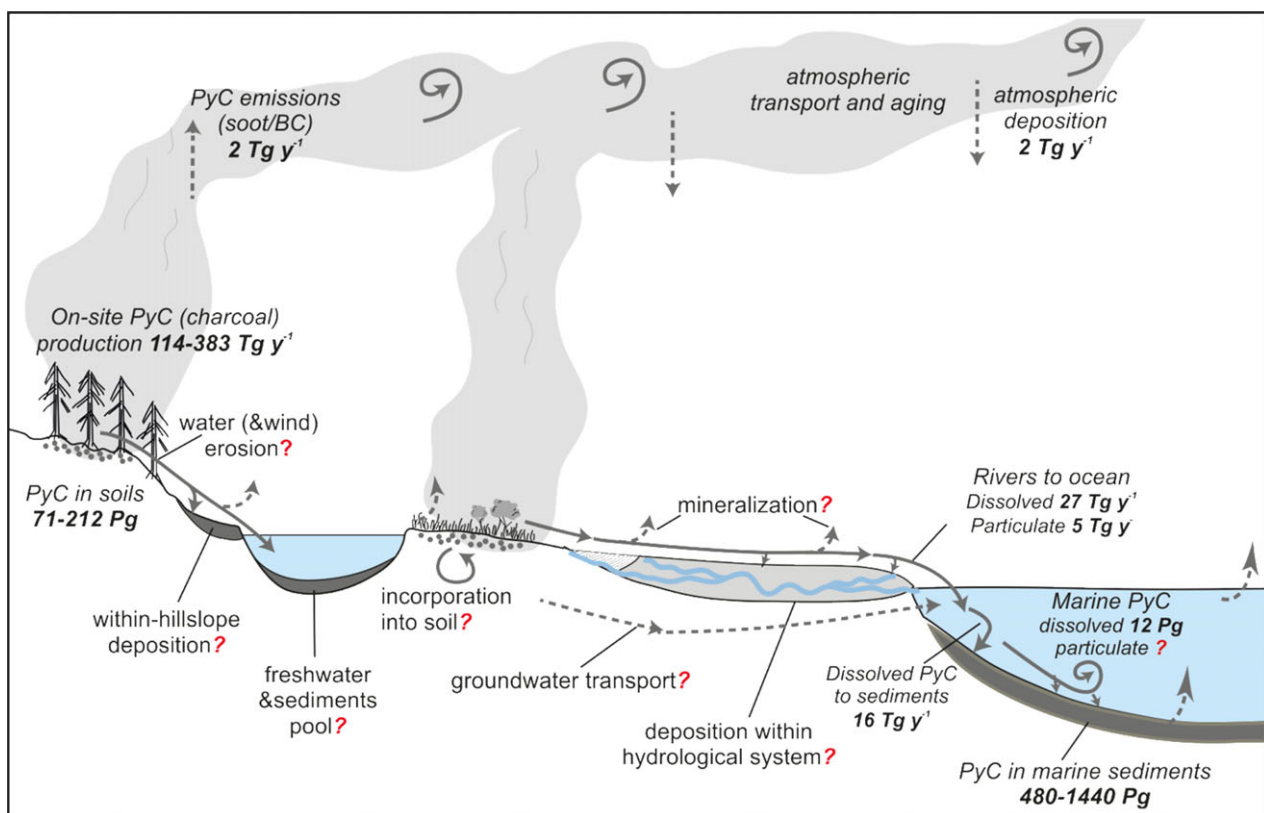
atmosphere as aerosol PyC. These numerical estimates have been used extensively (e.g. Forbes *et al.*, 2006; Alexis *et al.*, 2007; Zimmerman *et al.*, 2012), although their general applicability is questioned here. The ratio of 80/20% was obtained by Kuhlbusch & Crutzen (1995) by simple comparison of production rates of PyC that remained on-site following the laboratory and prescribed fires with emission factors for emitted 'aerosol PyC' obtained from other studies. For this calculation, emitted vs. remaining PyC was not determined for any specific fire nor was its variability with vegetation or fire characteristics examined. This ratio is likely to vary substantially with environment, fuel type and fire behaviour, and we suggest that site-specific validation studies are a critical need for future research. For example, Saiz *et al.* (2014a) quantified PyC produced during 16 small-scale experimental burns in tropical savannah, distinguishing between PyC remaining on the ground and 'distal' PyC (airborne 125–10  $\mu\text{m}$  particles and some soot material). They found that the distal component

was always <3% of the total amount PyC produced. Unfortunately, not all atmospheric PyC was accounted for, as particles <10  $\mu\text{m}$  were not quantified.

#### Global estimations of PyC production

Updated estimates of the global PyC production from vegetation fires are presented in Fig. 4, divided into on-site PyC (charcoal) and atmospheric PyC (i.e. BC emissions). Global BC emissions from vegetation fires have recently been estimated as 1.85 Tg BC  $\text{yr}^{-1}$  (average for the period 1997–2014 from the Global Fire Emissions Database GFED4s, 2015). These BC emissions include both BC and elemental C emissions (see Akagi *et al.*, 2011), so for simplicity, we assume a C concentration of 100% (i.e. 2 Tg BC  $\text{yr}^{-1}$  = 2 Tg C  $\text{yr}^{-1}$  in Fig. 4).

Regarding our on-site PyC (charcoal) production estimations, the proportion of fuel C affected by fire (CA) that is transformed to PyC (PyC/CA) has been previously assumed to be 1–5% (e.g. Forbes *et al.*, 2006;



**Fig. 4** Global cycle of pyrogenic C (PyC) from vegetation fires. PyC production (in Tg C  $\text{yr}^{-1}$ ) is divided in on-site (charcoal) and atmospheric (soot/BC) PyC. Fluxes between atmosphere, terrestrial and marine environments are given in Tg C  $\text{yr}^{-1}$ . Main PyC pools are given in Pg C. Main uncertainties and unknowns are represented by red question marks. Data derived from Schmidt & Noack, 2000; Hockaday *et al.*, 2007; Elmquist *et al.*, 2008; Dittmar & Paeng, 2009; Jaffé *et al.*, 2013; Coppola *et al.*, 2014; Scharlemann *et al.*, 2014; Bird *et al.*, 2015 and the GFED4 database. These estimates are based on data produced using different approaches which do not account for regional variability and may not distinguish between PyC from different sources. For more details see main text.

Preston & Schmidt, 2006). However, the most recent and comprehensive studies addressing different ecosystems reported substantially higher conversion rates: 16% PyC/CA for savannah fires (Saiz *et al.*, 2014a), 27% in a boreal forest fire (Santín *et al.*, 2015) and 16% in tropical slash and burn fires (Righi *et al.*, 2009). Therefore, it seems justified to apply an increased estimate here of ~5–15% of the total C affected by fire converted to PyC. Using this PyC/CA conversion rate, the annual amount of PyC can be derived from the amount of total C emitted globally (2.17 Pg C yr<sup>-1</sup>; average for the period 1997–2014 from GFED4s, 2015), according to the equation CA = 'C emitted' + PyC (Santín *et al.*, 2015). This translates into an annual global PyC production of 114–383 Tg C yr<sup>-1</sup> (Fig. 4).

The sum of on-site and atmospheric PyC production is 116–385 Tg C yr<sup>-1</sup>. This represents ~0.2–0.6% of the terrestrial annual net primary production (Huston & Wolverton, 2009), further stressing the global significance of PyC in the C cycle. Our estimations of PyC production exceed the high end of the previously reported ranges (50–270 Tg C yr<sup>-1</sup>, Kuhlbusch & Crutzen, 1995; 49–200 Tg C yr<sup>-1</sup>, Schmidt & Noack, 2000; 63–140 Tg C yr<sup>-1</sup>, Bird *et al.*, 2015). In the present calculations, atmospheric PyC accounts only for 0.5–1.6% of the total production, a lower proportion than what has been previously estimated (e.g. 3–12% Schmidt & Noack, 2000; 5 ± 3% Bird *et al.*, 2015).

It is essential to remember that not all produced PyC has the same MRT. Some PyC will be mineralized on the timescale of weeks (e.g. anhydrosugars, Norwood *et al.*, 2013), while other forms may persist for millennia (e.g. woody-charcoal, Ohlson *et al.*, 2009). Thus, as will be further discussed, an accurate incorporation of PyC into the C cycle would require consideration of this variability in MRTs.

### Degradation vs. mobilization of PyC

Based on a simple calculation, Goldberg (1985) estimated that if all PyC produced during vegetation fires remained, all the C on the Earth's surface would be transformed to PyC in <100 000 years. Obviously, not all PyC remains in the environment in the medium or long term and the question arises, where does it all go? To address this fundamental issue, two main mechanisms for PyC removal need to be considered together: degradation and mobilization.

#### PyC degradation

The assumption of PyC being inert has long been demonstrated to be wrong (Goldberg, 1985). What remains clear is that many pyrogenic transformations enhance

the chemical recalcitrance of the organic materials, which prolongs their MRTs in the environment (Schmidt *et al.*, 2011; DeLuca & Boisvenue, 2012; Knicker *et al.*, 2013). Estimated MRTs of pyrogenic materials (including biochar) are very variable, ranging from decades or centuries (e.g. Bird *et al.*, 1999; Hammes *et al.*, 2008; Steinbeiss *et al.*, 2009) to millennia (e.g. Thevenon *et al.*, 2010; de Lafontaine *et al.*, 2011). Critically, however, the MRTs of PyC products are generally one or two orders of magnitude longer than those of their unburnt precursors (Baldock & Smernik, 2002; Knoblauch *et al.*, 2011; Brunn & EL-Zehery, 2012; Santos *et al.*, 2012; Maestrini *et al.*, 2014a; Naisse *et al.*, 2015a). Furthermore, PyC, together with fossil C, is the only form of non-mineral-associated organic matter that shows long-term persistence in mineral soils (Marschner *et al.*, 2008). Thus, PyC is likely to be a potent C sink over the medium and long term (decades to millennia). An exception to this general statement is the water-soluble PyC fraction of low-temperature chars, with turnover rates in the order of weeks to months (Norwood *et al.*, 2013).

Experimental results for PyC mineralization in soils have been contradictory, with reported decomposition rates ranging from rapid (e.g. 0.07% day<sup>-1</sup>, Hilscher *et al.*, 2009) to slow (e.g. 0.0007% day<sup>-1</sup>, Kuzyakov *et al.*, 2014). However, short-term incubation experiments can lead to unrealistically low MRTs (Woolf & Lehmann, 2012; Kuzyakov *et al.*, 2014). In early stages, degradation of the labile and readily available compounds in PyC occurs, which is reflected in relatively fast degradation rates (Zimmerman, 2010). Mukome *et al.* (2014) illustrated this by showing that the labile aliphatic PyC fraction is degraded first, whereas the oxidation of the aromatic PyC portion occurs more slowly. Fast degradation of labile components also explains why PyC produced at low temperatures degrades faster than PyC from high temperatures as the labile fraction is relatively large in low-temperature PyC (Inoue & Inoue, 2009; Ascough *et al.*, 2011). The loss of labile PyC components with ageing leads to a decrease of PyC degradation rates over time (e.g. Hamer *et al.*, 2004; Bruun *et al.*, 2008; Kuzyakov *et al.*, 2009; Knoblauch *et al.*, 2011). Therefore, realistic long-term turnover dynamics of the different PyC forms have to be considered when estimating MRTs of PyC in the environment (Foereid *et al.*, 2011; Kasin & Ohlson, 2013). PyC degradation should not be estimated using a single-pool, single residence time model.

Recent studies investigating MRT of PyC in soils have used multipool models representing different biomolecular classes in soils (e.g. Singh *et al.*, 2012; Woolf & Lehmann, 2012; Knicker *et al.*, 2013; Bird *et al.*, 2015). The most recent model for PyC mineralization



proposed by Bird *et al.* (2015) differentiates three PyC pools: a labile (anhydrosugars and methoxylated phenols; half-life of weeks to months), an intermediate semilabile (polycyclic aromatic compounds <7 rings; half-life of years to centuries) and a stable pool or SPAC (polycyclic aromatic compounds >7 rings; half-life of centuries to millennia). The contribution of these pools varies with the formation conditions and original material. For PyC formed in natural fires, Bird *et al.* (2015) speculated that contributions for labile, semilabile and stable pools maybe are around 10%, 40% and 50%, respectively, which suggests that most of PyC formed is in relatively stable forms.

A major limitation of previous work on PyC decomposition rates is that much of it has focused on processes occurring in surface soils. However, surface soil horizons are only one type of environment where PyC accumulation has been identified (e.g. *Terra Preta* soils; Glaser & Birk, 2012). Most ancient charcoal deposits are found in environments with low decomposition rates such as peats, lake sediments, alluvial fans, flood plain deposits or deep marine sediments (Scott, 2000). Hence, knowledge on PyC degradation in environments where PyC accumulates such as deep soil horizons or depositional sites is required (Dungait *et al.*, 2012; Marin-Spiotta *et al.*, 2014). As a proxy, some studies examined PyC decomposition under differing environmental conditions such as oxygen availability (Nguyen & Lehmann, 2009; Knoblauch *et al.*, 2011), temperatures (Cheng *et al.*, 2006; Nguyen *et al.*, 2010) or alkalinity (Braadbaart *et al.*, 2009). In addition, more information is needed about interaction with the matrix in which PyC is held, given that physicochemical stabilization/protection (e.g. occlusion within aggregates, adsorption onto minerals, per-mineralization) is increasingly seen as a key factor in PyC preservation (de Lafontaine *et al.*, 2011; Cusack *et al.*, 2012; Bruun *et al.*, 2014). The only study to date examining the decomposition of PyC in subsoils points to soil physicochemical parameters being more critical for stabilization than microbial community characteristics (Naisse *et al.*, 2015a).

When examining the relationship between PyC accumulation in soils and C losses to the atmosphere, the effect of PyC on soil organic matter degradation must also be considered. This is especially relevant when PyC is added to the soil for C sequestration and soil amelioration purposes (i.e. biochar application). Studies testing the hypothesis that PyC can prime the decomposition of soil organic matter have had mixed results, with effects being negative (e.g. Cross & Sohi, 2011; Jones *et al.*, 2011; Zimmerman *et al.*, 2011; Whitman *et al.*, 2014; Naisse *et al.*, 2015a), positive (Wardle *et al.*, 2008; Zimmerman *et al.*, 2011; Singh *et al.*, 2014; Naisse *et al.*, 2015a) or absent (Hilscher *et al.*, 2009; Kuzyakov

*et al.*, 2009; Cross & Sohi, 2011; Brunn & EL-Zehery, 2012; Santos *et al.*, 2012). The direction of the priming effects depends on several factors such as soil type, original soil organic matter quantity and quality, climate/incubation conditions, and PyC amount and characteristics (Stewart *et al.*, 2013; Michelotti & Miesel, 2015). In a meta-analysis of 18 studies on PyC-induced priming, Maestrini *et al.* (2014b) suggested that overall the presence of a labile fraction in PyC may induce a positive priming effect in the short term, whereas in the long term, PyC may induce a negative priming by promoting physical protection mechanisms. The only available modelling estimates on long-term potential priming of PyC (biochar) additions on soil organic C concluded that, even for the worst-case scenario examined, the potential negative priming effect exceeds by far the potential positive priming effect (Woolf & Lehmann, 2012).

Biotic degradation is currently the better-understood pathway for PyC decomposition (Kuzyakov *et al.*, 2009; Santos *et al.*, 2012), although abiotic degradation is also important (Cheng *et al.*, 2006; Spokas *et al.*, 2014). Abiotic factors such particle disintegration during water erosion, cryoturbation or gelifluction are notable drivers of PyC degradation (Preston & Schmidt, 2006; Spokas *et al.*, 2014). For example, Naisse *et al.* (2015b) exposed PyC (biochar) to wetting/drying and freezing/thawing cycles and noted substantial losses (10–40% C) by leaching of dissolved and small particulate PyC (<20  $\mu\text{m}$ ).

Consumption of existing PyC by subsequent fires has also been highlighted as a possible major abiotic loss mechanism of PyC in soils (Ohlson & Tryterud, 2000; Czimczik *et al.*, 2005; Preston & Schmidt, 2006; Czimczik & Masiello, 2007; Kane *et al.*, 2010). However, none of these studies have produced direct evidence to support this suggestion. More recently, two studies measured PyC consumption by fire in contrasting environments: an experimental boreal forest fire (Santín *et al.*, 2013) and a prescribed fire in open savannah woodland (Saiz *et al.*, 2014b). Both found only minor losses of existing PyC (median mass losses <15% in Santín *et al.*, 2013; average mass losses <8% in Saiz *et al.*, 2014b), suggesting that subsequent fire is not a major cause of PyC loss.

It is essential to recognize that although PyC can be altered and degraded, only its transformation to CO<sub>2</sub> (and other gases) constitutes a net loss to the atmosphere. Through the alteration/degradation process, PyC can evolve into other PyC forms, which may still act as C sinks and need to be accounted for. For example, during degradation, some PyC can enter the dissolved organic matter pool (Hockaday *et al.*, 2006; Guggenberger *et al.*, 2008; Major *et al.*, 2010). It is not

clear how much of the PyC becomes soluble during ageing, but some exploratory results suggested that this soluble fraction tends to increase with ageing (Abiven *et al.*, 2011). This is consistent with the observation by Dittmar *et al.* (2012a) of a continuous flux of dissolved PyC from a burnt catchment decades after the fire. Ding *et al.* (2013) also suggested the continuous export of dissolved PyC over long timescales as a plausible explanation for the lack of correlation between dissolved PyC concentration and recent fire history (<20 years) in grassland streams. Some of this dissolved PyC is likely to become part of the recalcitrant pool of dissolved organic C in the deep ocean or sequestered in abyssal sediments where its MRT is in the order of thousands of years (Ziolkowski & Druffel, 2010; Coppola *et al.*, 2014). From a global C accounting perspective, it is important to note that this C, effectively sequestered from the atmosphere, is largely not accounted for.

In the case of atmospheric PyC, this is subjected to a range of alteration processes during ageing, including coating by coagulation and condensation with other aerosols, oxidation and incorporation into liquid water. All these processes have profound implications not only for PyC dynamics, but also for climate forcing and human health (Zhang *et al.*, 2008; Wang *et al.*, 2014a). It has been suggested that, through reactions with atmospheric oxidants, solubilization of atmospheric PyC could result in it entering the dissolved organic C pool (Masiello, 2004); however, the exact mechanisms still need to be identified.

#### *PyC mobilization*

The reservoir that a PyC particle initially enters and its further mobility are primarily determined by its size: for small PyC particles emitted during burning (atmospheric PyC, size <1–2  $\mu\text{m}$ ), the initial reservoir is predominantly the atmosphere, whereas for larger PyC particles, it is the burnt area and surroundings (although some large charred particles may become airborne during fire; Tinner *et al.*, 2006; Oris *et al.*, 2014). Following this basic division by size, atmospheric transport would be the main mobilization pathway for emitted PyC particles and transport by water for on-site PyC particles (Scott, 2010). In some cases aeolian transport may also contribute to mobilization of PyC particles that have initially been deposited on-site (Pereira *et al.*, 2015).

*Mobilization of atmospheric PyC.* Atmospheric PyC is transported globally through atmospheric circulation, which allows its deposition in remote environments such as deep-sea sediments (Lohmann *et al.*, 2009) or on ice sheets (McConnell *et al.*, 2007). Residence times

of PyC in the atmosphere are in the order of a few days, much shorter than the long-lived greenhouse gases (Feichter & Stier, 2012). Atmospheric deposition and fluvial fluxes are the main mechanisms for delivery of atmospheric PyC to marine sediments (Suman *et al.*, 1997), which are considered to be the final PyC sink (Masiello & Druffel, 1998). The relative importance and spatial patterns of these fluxes in the global context are still a subject of debate. Data by Elmquist *et al.* (2008) from the Arctic Ocean point to a predominance of inputs from terrestrial systems by rivers over direct atmospheric deposition, whereas Sánchez-García *et al.* (2012) accounted atmospheric deposition to be much larger than fluxes from rivers for the northern European shelf. Lohmann *et al.* (2009) estimated both fluxes being of similar quantitative importance for the South Atlantic Ocean.

Mobilization of atmospheric PyC in terrestrial systems and deposition in intermediate pools also warrant further investigation (Masiello & Druffel, 1998). For example Bisiaux *et al.* (2011) examined the inputs of atmospheric PyC nanoparticles in a lake after a large wildfire and found that most of this PyC reached the lake immediately after the fire by direct atmospheric deposition, rather than by fluvial transport and subsequent deposition.

*Mobilization of on-site PyC.* Soil erosion by water is usually enhanced after fire by loss of the vegetation cover and, in some cases, increase of soil water repellence and/or the destabilization of soil structure (Certini, 2005; Shakesby & Doerr, 2006). Given that PyC particles typically have a lower density than soil, and are located on or within the soil surface, a significant part of the PyC may thus become mobilized by postfire water erosion (Fig. 3c; Rumpel *et al.*, 2015; Wagner *et al.*, 2015). Rumpel *et al.* (2009) found that, even on a slope of only 1%, 7–55% of PyC produced in an experimental savannah fire was subject to erosion under simulated rainfall. Boot *et al.* (2015) did not detect substantial incorporation of PyC into the mineral soil four months after a conifer forest fire and concluded that most PyC generated aboveground was likely transported off-site through erosion events. In a study characterizing PyC pools across a boreal forest watershed, Ohlson *et al.* (2013) showed that the lake sediment contained more PyC per unit area than the forest soil surrounding the lake, which also supports the importance of lateral PyC mobilization.

It has also been demonstrated that ‘fresh’ PyC is preferentially transported *ex situ* by water erosion with respect to bulk soil organic matter, probably due to its low-density particulate nature and lack of immediate interaction with the mineral soil phase (Chaplot *et al.*,

2005; Rumpel *et al.*, 2006). However, even if lateral erosion is becoming widely recognized as one of the main mechanisms for PyC removal from surface soils (Major *et al.*, 2010; Foereid *et al.*, 2011), PyC flux by erosion has been scarcely quantified and relationships between soil erosion and PyC movement remain poorly understood (Rumpel *et al.*, 2015). The effects of PyC intrinsic characteristics (e.g. particle size, density, porosity, hydrophobicity; Kinney *et al.*, 2012; Brewer *et al.*, 2014) and environmental factors (e.g. topography, rainfall regime, soil type; Boot *et al.*, 2015; Rumpel *et al.*, 2015) in the transport of PyC are yet to be elucidated.

In addition to lateral movement of PyC by erosion and its potential off-site transport by wind and water, vertical transport through the soil profile can also occur (Rumpel *et al.*, 2015). This movement can be driven by water flow and is governed by intrinsic PyC properties and soil characteristics. For example, Wang *et al.* (2013) observed a greater vertical mobility for PyC particles with smaller sizes and lower surface charges. Haefele *et al.* (2011) found that 50% of the biochar moved below 0.30 m in the soil profile within 4 years after its application to a sandy soil, whereas vertical movement was inappreciable in another soil with poor percolation rates.

Vertical movement can also occur by bioturbation or physical processes such as gelifluction and cryoturbation (Schmidt & Noack, 2000; Preston & Schmidt, 2006; Elmer *et al.*, 2015). Although the main direction is downward, upward vertical movement of PyC can occur by, for example, bioturbation or uprooting of trees (Carcaillet, 2001). The oxidation of PyC with ageing may enhance vertical transport of PyC by increasing its polarity, which may promote its movement through the profile with water (Knicker, 2011). Notwithstanding this, PyC oxidation may also enhance its interaction with the soil mineral phase, which in turn could increase its stabilization within the soil (Brodowski *et al.*, 2006). Singh *et al.* (2014) found that these PyC–mineral interactions were formed in <1 year in a temperate forest Cambisol.

Vertical movement and subsequent accumulation of PyC in deeper soil horizons can contribute to its preservation (Dungait *et al.*, 2012; Lorenz & Lal, 2014), but also, in the case of very small particles or dissolved PyC, could facilitate further transportation by groundwater (Hockaday *et al.*, 2007; Dittmar *et al.*, 2012b). In addition to this, PyC incorporated into the soil matrix disintegrates and oxidizes into water-soluble low-molecular mass compounds (Abiven *et al.*, 2011; Spokas *et al.*, 2014). Water fluxes carry these dissolved PyC compounds horizontally across landscapes into rivers. This is a slow but continuous process that affects land–ocean fluxes globally (Dittmar *et al.*, 2012a; Jaffé

*et al.*, 2013). Wagner *et al.* (2015) reported simultaneous measurements of dissolved and particulate PyC fluvial export one year after a wildfire. They found that their dynamics were decoupled: dissolved PyC fluxes were not significantly affected by recent fire activity, whereas particulate PyC export was substantially larger in recent fire-affected areas when surface run-off occurred. This highlights the need to understand the specific mobilization mechanisms for the different forms of PyC.

It is worth noting that transformation of PyC may take place during transport. For example, PyC particles subjected to water mobilization may suffer abrasion and fragmentation (Scott, 2010; Crawford & Belcher, 2014). Moreover, as is the case for other forms of C within soil organo-mineral complexes, PyC can become exposed during soil erosion and transport through the breaking of soil aggregates and could therefore be more susceptible to degradation (Berhe *et al.*, 2007). Finally, the degradation of dissolved PyC can also be quantitatively and qualitatively important during its transport in surface waters. Stubbins *et al.* (2012) pointed to photo-degradation as being responsible for the shift from highly condensed aromatics in terrestrial waters to less condensed PyC structures in the open ocean. Myers-Pigg *et al.* (2015) estimated that half of the low-temperature dissolved PyC is lost in Arctic rivers during the transport from fire source to the ocean.

#### *Global estimations of PyC fluxes*

A representation of the main PyC fluxes is shown in Tg C yr<sup>-1</sup> in Fig. 4. For this annual timescale, atmospheric deposition is equal to atmospheric emissions, given that the residence time of PyC in the atmosphere is in the range of days (Feichter & Stier, 2012). It has been estimated that around half of the atmospheric deposition in the oceans takes place over the continental margins and the other half in the open ocean (Suman *et al.*, 1997); however, to the authors' knowledge, no robust data are available distinguishing between deposition on land vs. ocean. Therefore, the deposition of atmospheric PyC in Fig. 4 is given as a single value. Regarding transport from land to ocean, we focus on riverine fluxes although some minor remobilization and short-distance deposition by wind may also take place (Suman, 1986). Together, riverine fluxes of particulate and dissolved PyC to oceans may account for about 8–27% of the total annual production of PyC (Fig. 4). Riverine particulate PyC inputs to marine sediments were first estimated as 12.2 Tg PyC yr<sup>-1</sup> by Suman *et al.* (1997), with most of the PyC (94–96%) being deposited on the continental shelf. Here, we use the revised values presented by Elmquist *et al.* (2008),

which increase global riverine PyC flux to 26 Tg C yr<sup>-1</sup>, but with only 20% derived from vegetation burning (i.e. 5.2 Tg C yr<sup>-1</sup>; Fig. 4), and the rest derived from <sup>14</sup>C-extinct sources. This is probably a low estimate of riverine particulate PyC inputs because the method used by Elmquist *et al.* (2008) does not detect the less recalcitrant component of particulate PyC (Hammes *et al.*, 2007).

For dissolved PyC, Jaffé *et al.* (2013) identified a global flux of 26.5 Tg C yr<sup>-1</sup>, corresponding to ~10% of the global riverine flux of dissolved organic C. Importantly, Jaffé *et al.* (2013) quantified only the most recalcitrant forms of PyC, and therefore, values would be higher if labile dissolved PyC forms were also considered (Myers-Pigg *et al.*, 2015). Regarding mechanisms for transfer of PyC from waters into sediments, Coppola *et al.* (2014) suggested that sorption of dissolved PyC to sinking particulate organic C and deposition into abyssal sediments could account for ~16 Tg C yr<sup>-1</sup> (Fig. 4). Other transfer mechanisms need yet to be quantified. Fluxes from terrestrial ground waters to oceans also remain unquantified (Fig. 4), representing another gap in our understanding of the global PyC cycle.

### Long-term fate of PyC

Burial in marine sediments is usually considered the ultimate fate of PyC (Masiello, 2004). Storage conditions in this anoxic environment are ideal for PyC preservation, with estimated MRTs of several thousands of years (Masiello & Druffel, 1998). However, at the global scale, PyC concentrations measured in marine sediments do not account for all the PyC generated, even considering partial degradation of PyC in the depositional marine environments (Masiello, 2004). This points to the potential importance of 'intermediate' PyC reservoirs, which are poorly understood. Regnier *et al.* (2013) estimated the current lateral anthropogenic-induced fluxes of C from land to ocean and reported that, globally, only <20% of this C is exported to the open ocean and ~30% is emitted, whereas another ~50% is instead accumulated in 'intermediate reservoirs' along the continuum of freshwater, estuaries and coastal environments.

Deep soil is one of the intermediate reservoirs gaining attention in the C sequestration context (Lorenz & Lal, 2014). However, it is still not clear how quantitatively relevant the 'deep' PyC in soils is, as it remains unaccounted for in studies that do not consider the whole soil profile (Rumpel & Kögel-Knabner, 2011). Other poorly understood intermediate PyC reservoirs are depositional sites within terrestrial environments such as colluvial and alluvial deposits, lake and reser-

voir sediments, peats and other types of wetlands, and river bank and floodplain deposits (Gerlach *et al.*, 2012; Springer *et al.*, 2012; Wang *et al.*, 2014b; Matthews & Seppälä, 2015).

In terrestrial depositional environments, MRTs of PyC are expected to be relatively long as environmental conditions promote low decomposition rates through, for example, oxygen deficiency, physical protection and/or substrate-driven biological rate limitation (Fig. 3d; Knicker, 2011; Dungait *et al.*, 2012). However, these terrestrial depositional environments are not as stable as marine depositional environments. They are subjected to disturbances over a range of temporal and spatial scales, which can lead to the remobilization of PyC. For example, Ryan *et al.* (2011) reported PyC enriched fluvial discharges several years after a wildfire, caused by soil remobilization after intense rainfall. Hatten *et al.* (2012) found that flood events can lead to input of particulate PyC to rivers by mobilization of PyC stored in near-stream deposits. However, even if remobilization from intermediate terrestrial environments takes place, it is necessary to bear in mind that this is a recurring and natural geomorphological process acting at the landscape scale, with most of the material being redeposited within the landscape (Chaplot *et al.*, 2005; Shakesby & Doerr, 2006; Rumpel *et al.*, 2009). Therefore, fluxes between sites do not necessarily imply a net loss or export of PyC from terrestrial systems.

In addition to these terrestrial PyC reservoirs, transitional environments at ocean margins such as estuaries and other coastal wetlands may also hold substantial PyC pools, considering that most of the PyC deposition from rivers to marine sediment occurs near shore (Golding *et al.*, 2004; Ding *et al.*, 2014; De la Rosa *et al.*, 2015). In these transitional environments, particulate PyC can shift from being partially saturated (and thus capable of floating and long-distance transport) to fully saturated and deposited. Also, fluctuations in water table position frequently cause changes in redox conditions in wetlands and coastal zones, which can promote coprecipitation of dissolved PyC with iron hydroxides and other minerals (Riedel *et al.*, 2013). Once enclosed in a mineral matrix and buried in sediments, PyC may be stabilized over long periods of time (Riedel *et al.*, 2013).

### Global estimations of PyC pools

Robust quantifications of the global PyC pools are currently unavailable, although some estimates can be provided (Fig. 4). Hockaday *et al.* (2007) estimated that if the PyC contents of soil, freshwater and coastal waters and sediments are assumed to be in the order of 5–15%



of the total organic C, it would imply a global PyC reservoir of 300–500 Pg C. If the same approach is taken for soils with updated figures (global soil organic C pool 1416 Pg; Scharlemann *et al.*, 2014), it translates into a global PyC pool in soils (0–100 cm depth) of 71–212 Pg C (Fig. 4). Past estimates of PyC stored in marine sediments were 2400–6000 Pg (Schmidt & Noack, 2000). However, those numbers did also include PyC derived from lithogenic graphite (Dickens *et al.*, 2004), and updated estimates are now in the range of 400–1200 Pg in coastal and 80–240 Pg in open ocean sediments (Bird *et al.*, 2015) (Fig. 4). Future studies examining mechanisms of PyC stabilization in marine sediments are necessary to obtain estimates for the size of this pool.

Recent estimates for dissolved PyC in the ocean are 12 Pg C (Dittmar & Paeng, 2009) and 26–145 Pg C (Ziolkowski & Druffel, 2010). In Fig. 4, we use the more conservative estimate by Dittmar & Paeng (2009) because their study included hundreds of samples of major oceanic water masses. We stress again that estimates are still lacking for key PyC pools such as terrestrial sediments (e.g. lakes, reservoirs, floodplains), freshwaters and particulate PyC in ocean waters (Fig. 4); quantification and characterization of PyC in these pools is an important area for future research.

## Conclusions

### *New directions and challenges: an integrated view of PyC in the environment*

Fire is a globally important perturbation in the Earth system, and the extent and intensity of vegetation fires are expected to increase in some regions under predicted future climatic scenarios (Flannigan *et al.*, 2013; Moritz *et al.*, 2014). Thus, irrespective of current efforts for decreasing global anthropogenic PyC emissions, natural PyC production from vegetation fires will remain a major and potentially increasingly important player in the global C cycle. To quantitatively assess the role of PyC in the global C budget and climate prediction models, an integrated view including multiple pools and fluxes of PyC is required. In addition, a full understanding of PyC generation from vegetation fires can provide us with new opportunities for mitigating climate change through, for example, optimizing management of burns for maximum PyC production (Ottmar, 2014). Furthermore, the lessons learned from natural PyC can be used to elucidate the longer-term implications of biochar as a tool for C sequestration and climate change mitigation (Woolf *et al.*, 2010; Lorenz & Lal, 2014). We conclude here by proposing further

directions in PyC investigations that may help to achieve these ambitious objectives:

*Complete PyC production inventories and conversion factors.* Simultaneously acquired quantitative data are needed for the whole spectrum of PyC produced, both PyC remaining on-site and emitted to the atmosphere. These data are required with respect to fuel consumed for a range of fuel types and fire behaviours. When emission factors regarding gases and aerosols are estimated for different fuel types and burning conditions (e.g. Akagi *et al.*, 2011; Urbanski, 2014), conversion factors for PyC production could be determined simultaneously. This would allow direct incorporation of PyC production into C emissions models.

*Characterization of the whole range of PyC products and their MRTs.* The assessment of the types and relative proportions of all PyC generated would allow not only robust estimation of total PyC produced, but also determination of their characteristics and MRTs. MRT of PyC is not only determined by chemical recalcitrance (e.g. SPAC) but also by physical properties and environmental factors; all of these parameters need to be considered for realistic estimations of MRTs for different PyC types and for accurate appraisals of their roles as C sinks. In addition to this, the importance of pyrogenic organic matter in the cycles of other elements such as nitrogen, phosphorous or sulphur deserves further attention (Knicker, 2010).

*Full understanding of PyC intermediate pools and fluxes.* The relative importance of the whole range of intermediate PyC pools (and their fluxes) still need to be quantified to understand how much PyC is actually lost (i.e. mineralized) and how much is just moving between reservoirs. A key step forward in this regard has been the recognition of riverine fluxes of dissolved PyC to oceans as one of the major mechanisms for mobilization of PyC from soils (Jaffé *et al.*, 2013). Global-scale quantification of other major PyC fluxes should follow this example. In this context, simultaneous determinations of different PyC types would help elucidating whether their dynamics are coupled (e.g. Wagner *et al.*, 2015).

*Terrestrial PyC erosion–deposition as a C sink.* Within terrestrial environments, the potential for soil C erosion and subsequent deposition as a C sink is widely recognized (Berhe *et al.*, 2007; Van Oost *et al.*, 2007). Considering the characteristics of PyC (i.e. the recalcitrant nature of some fractions combined with its high susceptibility to water erosion), PyC erosion–deposition could be one of the key mechanisms of PyC preservation. An understanding of not only how PyC is transported

away from production sites, but also where it is deposited, is essential to quantify PyC fluxes and its ultimate role as long-term C sink. Substantial efforts have been made to measure and model postfire soil erosion and redistribution by the soil and geomorphology communities (Moody *et al.*, 2013), but PyC has to date not been examined as a component within these fluxes. Adding PyC to monitoring studies, or applying established methods in new investigations focusing on PyC, would deliver a fundamental understanding of pathways and quantities involved.

*Integration of PyC in models through interdisciplinary collaboration.* The PyC production estimates presented in this review suggest that the inclusion of PyC in the global C budget estimations could identify up to 25% of the current missing or residual terrestrial C sink ( $\sim 1.5 \text{ Pg C yr}^{-1}$ , Ciais *et al.*, 2013), with the majority of this expected to survive over decades or centuries (Bird *et al.*, 2015). However, few attempts have been made to incorporate PyC into C budgets and models. To date, this work has mainly focused on soil C models for savannah and agricultural soils (Skjemstad *et al.*, 2004; Lehmann *et al.*, 2008). In addition to soil C models, a wide range of advanced erosion-, fire-related emissions and sediment transport models exist that provide suitable platforms for including PyC (e.g. CASA-GFED, FO-FEM, CONSUME, CanFIRE, ERMiT, LISEM). A closer collaboration between the often distinct research communities specializing in fire behaviour and combustion, fire emissions, fire history, biogeochemical cycling, soil erosion and sediment fluxes could provide the knowledge and data required for incorporating PyC in such models. This integration would bring us closer to a robust global assessment of PyC from vegetation fires.

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