We would like to thank both reviewers for their time and for the detailed comments that they have provided. As a result of these comments we have made a number of changes to the paper. The major alterations to the paper have been: a re-written paragraph in the discussion highlighting the methodological differences between our study and others, why it is difficult to compare them and a suggestion of how to tackle this problem; the addition of a sampling plot design figure; and the addition of a schematic of the combustion analysis setup in the supplementary material. We have also added sentences or short passages where the reviewers have suggested they are required and we believe these changes improve the accuracy and readability of the paper. We have sought to address all the individual comments of the reviewers and have made corrections and clarifications where necessary. We have reproduced the reviewers' comments in italics below and our reply to each comment can be found underneath in blue font. A marked-up version of the revised manuscript is attached to the end of this document

Reviewer One

General comments

This paper presents a study that measured emission factors (EF), fuel load, and carbon content for different fuel types from a temperate forest (Eucalyptus) in south-eastern Australia. The measurements are used to estimate direct carbon emissions due to the application of prescribed fire. The carbon emission estimates derived from the studies measurements are compared to estimates based on less specific input and other methods.

Emission ratios (Table S3) and emission factors (Table 4) for the pooled species (CH4, NMHC and PM) are dramatically at odds all of the biomass burning EF literature of the last 20-30 years!
First, the pooled species are described as the sum of CH4, PM and non-methane hydrocarbons (NMHC), however the in the analysis used the pooled species represent all carbon containing species other than CO2 and CO, organic compounds (volatile and semi-volatile organic compounds) and PM. In biomass smoke a significant fraction of emitted carbon is present in oxygenated organic compounds (e.g. methanol, formaldehyde, furan, …,) that are not hydrocarbons (see Akagi et al., 2013) and the authors should have described the gas portion of the pooled species as VOC or simply organic compounds.

Improper terminology aside, the pooled EF (g-C/g-C) reported in this study are for the most part far higher than that inferred from virtually all previous studies that I am familiar with (e.g. ….).
The median of the EFpooled reported in Table 4 is 0.23 g-C/g-C with maximum of 0.97 gC/g-C. The pooled emissions account for 24% of carbon emitted on average with a maximum of 43% (leaf litter – Oliver). For comparison one may use the laboratory measurements reported in Yokelson et al. (2013). This study combined multiple instruments and methods (open path FTIR spectrometer, proton-transfer-reaction mass spectrometry (MS), proton-transfer ion-trap MS, negative-ion proton-transfer chemical ionization MS, and gas chromatograpghy with MS to measure emissions of over 300 compounds for large scale laboratory burns of forest and chaparral fuels. The supplemental material for Yokelson et al. (2013) supplemental includes EF for CO2, CO, CH4, over 300 organic gases, and PM2.5 for 25 lab burns of forest fuels. Following conversion of these EF from units of g/kg to units g-C/g-C it may be found that the sum of the EF for PM2.5, CH4, and all organic gases (>300 species) has a median of 0.039 g-C/g-C with a range of 0.008 to 0.139 g-C/g-C. The percent of emitted carbon contained in PM2.5, CH4, and all organic gases averaged 3.8% with a range of 0.7% to 13.6%. The EF data from Table 4 of the manuscript and from the supplemental material of Yokelson et al. (2013) are summarized in Table 1 below.

In this manuscript the pooled EF and the fraction of emitted carbon in pooled species are far higher than observed in the comprehensive study of Yokelson et al. (2013). The Yokelson et al. (2013) is consistent with previous and subsequent EF reviews (Andreae & Merlet, 2001; Akagi et al., 2011; Urbanski 2014) and field and laboratory studies (e.g. Burling et al. 2010; Burling et al., 2011; Akagi et al. 2013). For a wide range of biomass burning, the fraction of carbon emitted in species other than CO2 and CO is typically < 5% and rarely greater than 10%. In this manuscript the fraction of emitted carbon attributed to species other than CO2 and CO is on average 24% (with maximum of 43%) and far exceeds what is found the previously published literature. The fact that the MCE of the burns reported in this study are relatively high (average =0.96) and therefore indicative of high efficiency burns with low PM and VOC emissions makes the pooled EF even less believable. If the authors were to conduct a similar comparison against EF data from other studies / reviews they would arrive at similar results. The authors simply made a comparison versus Hurst et al (1994b) and concluded without any justification that the discrepancies were due to PM. The authors clearly failed in their duty to seriously compare their findings to previously published work. I can only conclude that significant errors were made in the calculation of the pooled EF and this study is therefore seriously flawed and I recommend that is rejected for publication.

Table 1.

		EF pooled[1] (g-C/g-C)		
	Average MCE	Median	Minimum	Maximum
Table 4	0.962	0.24	0.00	0.97
Yokelson et al. (2013)[3]	0.935	0.039	0.008	0.139

[1]Labeled as sum of CH4, NMHC, PM in manuscript and is the sum of PMm2.5, CH4, and > 300 organic gases for Yokelson et al. (2013)
[2]MCE = ΔCO2/(ΔCO + ΔCO2) and was calculated from Supplemental Table 3
[3]Derived from EF reported in the Supplemental Material for 25 forest fuel burns

We would like to thank the reviewer for the in-depth comparison of our pooled EF data to that of the literature. However, the primary aim of this manuscript was not to specifically discuss the form that the carbon was lost as (i.e CO_2, CO, VOCs or PM). The aim was to estimate the total carbon lost from south-east Australian forests because of prescribed fire, and how variable they can be when the amount of information you have changes. Specific emission factors for the prescribed burning sites were not available so we had to generate our own. These EFs were then used to create bounds for the random setting of EFs in two of the scenarios in the Monte-Carlo simulations, which are then summed to estimate the total carbon loss (Eqns. 4 and 5). Therefore, what is more important in this study is the total C loss through the sum of the different EFs rather than any one particular EF and we already demonstrate those sums as $\Sigma C_{emit}/C_{fuel}$ in Table 3.

With regard to the pooled data, we acknowledge that, yes, there does appear to be some major differences among the pooled values with those from the literature but the comparison the reviewer has made is a not direct one and not necessarily valid. The EF values for our study relate to different fuel components (twigs, litter, grass, etc.) while many of the published studies, and the reviews that have compiled them, have calculated their EFs based on laboratory studies of complete reconstructed fuel beds (e.g. Burling et al. 2010), and field based studies either on the ground (e.g.

Burling et al., 2011) or from aircraft (e.g. Akagi et al., 2013). Therefore, it is entirely plausible that the values recorded for some components of the total fuel may record higher than the aggregated values measured with complete fuel beds, either in the lab or field. Indeed, our values do overlap with the aggregated values but have a different distribution to the Yokelson et al. (2013) study, which is based upon fuels from range of US forest types measured in both the field and laboratory. Hence, we have included in the discussion how the methodological differences of our work make it difficult to compare with other Australian forest studies and that there is a need for measurements from south-east Australia to be made in a similar manner to Yokelson et al. (2013) to reconcile laboratory and field measurements (see page 15, L11-28 of the revised manuscript).

With regard to the comparison with Hurst et al. (1996), it was done because, at the time of submission, this was the closest study to ours that was on Australian forests. Indeed, a number of the studies the reviewer cites compare results among the same ecosystems within the United States. The reviews by Akagi et al. (2011) and Andreae and Merlet (2001) show that there are considerable differences between ecosystems and south-east Australian Eucalypt forests, by extension, are likely to be varied to. Of course, we acknowledge that the environment plays a role in the EF measured, which is why a laboratory bench top study was chosen to control for these environmental effects, as well as to produce some unique, site specific EFs. We have included in the discussion a statement that there is a need for more comprehensive emissions measurements specifically for south-east Australian forests. As previously mentioned, if these measurements are conducted in a manner similar to those for the south-eastern and south-western US (e.g. Yokelson et al., 2013), field and laboratory measurements may be reconciled (see page 15, L11-28 of the revised manuscript).

As a minor comment, we would also like to point out a typographical error in the reviewer's table that could potentially be misleading for readers of this discussion because it is reporting incorrect values from our Table 4. The maximum value from Table 4 would be 0.46, producing a median value of 0.22.

The revised paragraph, which also addresses the above comments and the comment regarding P13826, L11 -14 (below), reads as follows (see page 15, L11-28 of the revised manuscript):

"Across the four sites, the mean proportion of fuel carbon lost to the atmosphere relative to the total amount of carbon ($\Sigma C_{emit}/C_{fuel}$) was 86%. This is less than the 97% suggested by Hurst et al. (1996) for the one planned burn they measured in a south-east Australian forest. However, a direct comparison of this study with the Hurst et al. (1996) study cannot be made due to the significantly different methodological approaches taken that they may bias either study. These methodological differences include factors such as: the measurement of aggregated emissions from naturally structured fuels taken using an aircraft, compared to individual fuel components measured at a very small scale in the laboratory; and neither study measures the same range of compounds. Indeed, these methodological differences also prevent direct comparison of emission factors, not just with Hurst et al. (1996) but also the recent work of Paton-Walsh et al. (2014) who made ground-based emission measurements from planned fires in temperate south-east Australian fires. There are large variabilities in emission factors for certain compounds among different ecosystems (see reviews by Andreae and Merlet (2001) and Akagi et al. (2011)). This demonstrates the need for more comprehensive emissions measurements for specific ecosystems and regions, including south-east Australian forests. If these measurements are conducted in a manner similar to those for the south-eastern and south-western US (e.g. Yokelson et al., 2013), field and laboratory measurements may be reconciled."

Additional Comments

Combustion analysis method

I find it very uncertain that the combustion analysis employed is a reasonable proxy for the following reasons:

For many of the fuel components (twigs, ground layer, understory, overstory) filling a 10 cm × 10 cm × 3 cm sample holder seems to be a great distortion of the structure and arrangement of the natural fuelbeds which should have a significant impact on the manner in which the fuel burn and the subsequent emissions.

We acknowledge that there will be a distortion of the structure and arrangement of the fuel and this could be a potential source of error in terms of the subsequent emission factors. As mentioned previously, we have added a comment about this to the discussion. The small nature of the laboratory bench top testing equipment precludes us burning multiple components in their original configuration. However, for the purposes of this comparative study, every sample would have been treated and combusted in exactly manner and would carry the same intrinsic error.

The samples were combusted at a fixed irradiance of 25kW/m2 it is unclear how this approach replicates a natural free burning fire. It seems as though this approach could shift the combustion process towards flaming relative to natural fires.

The capacity to alter the flux used during combustion of samples using the MLC is limited. Regardless of this, work by Cruz et al. (2011) and Silvani et al. (2009) indicate that irradiances of 25 kW m^{-2} are achievable during a natural fire at the fire front and can remain that high for some time once the front has passed. This period of time is comparable to the length of time each burn was conducted (300 to 600 seconds).

Cruz, M. G., Butler, B. W., Viegas, D. X., and Palheiro, P.: Characterization of flame radiosity in shrubland fires, Combust. Flame, 158, 1970-1976, 10.1016/j.combustflame.2011.03.002, 2011.

Silvani, X., and Morandini, F.: Fire spread experiments in the field: Temperature and heat fluxes measurements, Fire Saf. J., 44, 279-285, 10.1016/j.firesaf.2008.06.004, 2009.

2.3 Combustion analysis

A diagram of the combustion analysis set-up is needed.

A diagram has been added to the supplementary material (Supplementary Figure 1) and a reference to this figure has been added to page 7, line 26-27.

It is stated that the mass of the samples were before burning and the mass of the residue after burning were recorded (P13817, L11-12). Was the carbon content of the fuel and residue also measured? It is unclear, but the laboratory combustion analysis should measure the mass and carbon content of fuel prior to burning and the mass and carbon content of the post fire residue to derive $\Sigma C_{emit}/C_{fuel}$ *for the carbon emission factor calculations (Eq. (1)).*

Yes, it was. This was stated on P13819, L4-6. (also see page 9, L7-9 of the revised manuscript)

Specific Comments

P13811, L26-28: The authors should discuss more broadly factors that affect fuel accumulations such as disturbance history (previous land use, fire, insects, etc.), topography, and soils.

We have added these factors into this paragraph. That part of the paragraph now reads: "Even so, fuel accumulation varies widely in space and time as a result of the interaction of many factors such as topography, soils, disturbance history (e.g. previous land use, insects, fire) and climate (e.g. due to variations in rainfall patterns; Bradstock, 2010); hence, remote sensing techniques will require intensive calibration." (see page 2, L30 to page 3, line 1 of the revised manuscript)

P13814, L14-15: The three fuel sampling plots at each were selected to have "similar slope and aspect". I suspect slope and aspect may have an important influence on the fuel loading as well as burning efficiency and possible fire severity. It seems that randomly locating the plots within the burn units to capture the variability of slope and aspect would have provided a better representation of the natural variability of emissions from prescribed fires in these forest types. Please comment.

This was a mistake on our part. It should have read that the three selected plots had similar slope and aspect due to the narrow elevation change of the general study area (64 m). We have changed the sentence to read:
"Within each study site, three permanent circular plots were established at least 500 m apart in similar vegetation type prior to planned burning. Due to the small elevation change of the of the general study area, all study sites had similar slope and aspect." (see page 5, L3-5 of the revised manuscript)

P13814 – 13815: Sampling Protocol.
The ground layer and forest floor (decomposing litter, twigs, leaf litter) was sampled using destructive / disruptive methods and separate post-fire quadrants would be required to estimate fuel consumption. Please describe where the post-fire quadrants were located relative to the pre-fire quadrants. I recommend including a diagram showing the sampling design.

We have included a diagram showing the sample design. This is now Figure 1 (page 27 of the revised manuscript and referred to at page 5, L7). The position of the post-fire quadrats is now addressed with the following sentence added to section 2.2.1: "The mass of ground layer vegetation, twigs and litter (see below) remaining after prescribed burning was measured in the same way using quadrats positioned 2-3 m from the position of the original quadrat to avoid the influence of biomass removal prior to prescribed burning." (See page 5, L32 to page 6 L1-3 of the revised manuscript.)

Please note if the "decomposing litter" included unidentifiable decomposing organic matter in the upper layer of soil that could be consumed by fire? I'm thinking of the 'duff' layer or organic soil layer typically found in Northern Hemisphere temperate and boreal forests. Is such a layer present and important in the forest examined in this study or Australian temperate forest in general? Please comment.

No observable duff layer was present in these forests, therefore explaining its absence from this study.

P13818: Define DeltaCO2, DeltaCO, etc. including units. Presumably these are molar mixing ratios as in Hurst et al. (1994b) but this must still be defined.

Yes, they are mixing ratios and a definition has been added. (See page 8, L12-14 of revised manuscript.)

P13818 L10-13: Define DeltaX and give units.

The definition has been added. (See page 8, L17 of revised manuscript.)

P13818: NMHC should be VOC (volatile organic compounds) as a significant fraction of emitted carbon is present in oxygenated organic compounds (e.g. methanol, formaldehyde, furan, …,) that are not hydrocarbons (see Akagi et al., 2013).

NMHC has been changed to VOC throughout the manuscript and the supplementary material.

P13826, L11 -14: "Across the four sites, the mean proportion of fuel carbon lost to the atmosphere relative to the total amount of carbon (ΣCemit/Cfuel) was 86 %. This is significantly less than the 97% suggested by Hurst et al. (1996). Hurst et al. (1996) based their analysis on the assumption that the carbon content of ash was constant at 6 %."
It seems that this is an invalid comparison as the Hurst et al. (1996) numbers alluded include forest fires of all types – clearing, prescribed, and wild while the current study examines only prescribed fires.

With hindsight, the comparison with Hurst et al. (1996) and the very recently published Paton-Walsh et al. (2014) values cannot be directly compared because of the different methodologies. The Hurst et al. (1996) study was an aircraft study and the Paton-Walsh et al. (2014) study was a series of ground based measurements. Therefore, both studies have measurements made on aggregated fuels rather than individual components. We have changed this paragraph to reflect this caveat and to also highlight the need for work similar to that of Yokelson et al. (2013) for south-east Australian forests. These revisions also address the reviewer's general comments and the revised paragraph can be read towards the end of that section (above). (Also see page 15, L11-28 of the revised manuscript.)

Technical Corrections
P13812, L2: Volkova and Weston (2013) reference missing from bibliography
Missing reference added. (Volkova, L., and Weston, C.: Redistribution and emission of forest carbon by planned burning in *Eucalyptus obliqua* (L. Herit.) forest of south-eastern Australia, Forest Ecology and Management, 304, 383-390, 10.1016/j.foreco.2013.05.019, 2013.) (See page 22, L11-13 of the revised manuscript.)

P13812, L19: change "whether" to "demonstration that"
Change made. (See page 3, L19 of the revised manuscript.)

P13812, L23: change "shorter" to "longer"
Change made. (See page 3, L22 of the revised manuscript.)

P13812, L27 insert "that" between "burning" and "reduces"
This sentence has been re-written. (See page 3, L25-27 of the revised manuscript.)

P13815, L26-28: The sentence beginning with "sample" does not makes sense the text "and a subsample of pre-fire fraction ground" seems out of place. Insert "were" between "fraction" and "ground"?
Change made. (See page 6, L14 of the revised manuscript).

P13862, L22: Eq. (4) predicts emissions not emission factors. Is this a typo? Should it cite Eq. (1)?
This is a typographical mistake and the re-writing of this paragraph now omits referring to this equation (See page 15, L11-28 of the revised manuscript).

Reviewer 2

General comments

This manuscript presents a study of carbon (C) emissions from prescribed fire from a temperate forest in south-eastern Australia. The authors measured emission factors, fuel load, and carbon content for different fuel types from four field sites to estimate direct C emissions after a prescribed fire. They used Monte-Carlo simulations to generate probability density functions of the parameters to account for low quality of data when calculating C emissions. Their results show that the uncertainty in their estimates of C emission declines with declining information quantity and coarse woody debris' inclusion in estimates increases the median C emissions and overall uncertainty in C emissions.

Specific Comments:

Abstract

L1: remove "of" before emissions

Removed. (see page 1, line 11 of revised manuscript.)

L10: effect of what on what? Please rephrase.

This has been rephrased to: "In order to assess the effect of declining information quantity and the inclusion of coarse woody debris when estimating emissions, Monte-Carlo simulations were used to create seven scenarios where input parameters values were replaced by probability density functions." (See page 1, L19-20 of the revised manuscript.)

Introduction

P13812 L21-24: Check to make sure this is correct, "Return frequencies of wildfires in extra-tropical (temperate) forests in Australia are typically shorter than that of tropical grassland and savanna and are often decadal compared to annual and biannual (Russell-Smith et al., 2007)." I could not find this claim in Russell-Smith et al 2007.

We have used an incorrect reference here and have changed the citation to Bradstock (2010) and Adams (2013). Both citations are already in the reference list. Also, the return frequency has been changed to 'longer' (see Reviewer one's comment). (See page 3, L21-23 of the revised manuscript.)

P13812 L24-25: Strange sentence structure, "In addition, and on an annual basis,"please rephrase, doesn't make sense.

We changed the sentence to: "In addition, the total area of temperate forest burnt on an annual basis is considerably smaller (Russell-Smith et al., 2007), notwithstanding large single fire events (Adams, 2013)." (See page 3, L23-25 of the revised manuscript.)

P13812 L26-28: "On the other hand, planned or prescribed burning reduces fuel loads in temperate forests is used at moderate return frequencies (e.g. 7–10 years) to mitigate risks to life and property from wildfires (Penman et al., 2007; McCaw, 2013)." Rephrase and look at multiple verbs in this sentence.

We have rephrased the sentence so it now reads: “Planned or prescribed burning in temperate forests to mitigate risks to life and property from wildfires is used at moderate return frequencies (e.g. 7 – 10 years) (Penman et al., 2007; McCaw, 2013).” (See page 3, L25-27 of the revised manuscript.)

What is TC, Tons of C?
It is metric tonnes of carbon but we have changed the units to megagrams (Mg) where one Mg equals one metric tonne throughout the manuscript.

P13812, L2: Volkova and Weston (2013) reference is missing from the references.
The reference has been added. (See page 22, L11-13 of the revised manuscript.)

Materials and Methods
Study sites
P13818, L5: Cfuel is defined as “the total mass of carbon contained in fuel that is burnt”, this does not corresponds to the caption of table 3, “Cfuel is the initial carbon content of fuel”.

The definitions have been changed to match (initial C content of the fuel). (See page 8, L11 of the revised manuscript.)

P13818 L1-8: Define DeltaCO2, DeltaCO, DeltaCH4, Delta PM and DeltaSigmaNHMC and give units.
As per Reviewer One’s comments the definition has been added. (See page 8, L12-14 of revised manuscript.)

P13818 L10-13: Define DeltaX and give units.
As per Reviewer One’s comments the definition has been added. (See page 8, L17 of revised manuscript.)

P13862, L22: Eq. (4) predicts emissions not emission factors. Is this a typo? Should it cite Eq. (1)?
Due to re-writing of this paragraph, the reference to the equation has been removed. (See page 15, L11-28 of revised manuscript.)

Results
Table 1 and 2 both have Carbon Content listed but different values and unit. Table 1 shows carbon content varying from 0-1 and it is a unitless number. Table 2 has unit in % dry weight and value is mostly >1. Are you referring to different things?

The definition in Table 1 has been changed to match that of Table 2. (See pages 23 and 24 of revised manuscript.)

Table 3-What is the rationale for assuming the same ΣCemit/Cfuel for twigs and CWD?
CWD was defined as woody material greater than 25 mm in diameter. ‘Twigs’ was the term given to woody material (twigs, wood and bark) that was between 10-25 mm in diameter. Both are effectively the same material, just different sizes.

Figure 1 – Change the y axis label to “Proportion of total biomass (%)” because you are showing both pre and post burn measurements. Correct the legend as well.
Changes have been made to now read ‘proportion of total biomass (%)’. (This is now Figure 2 - see page 28 of revised manuscript.)

Figure 2-Describe 7 scenarios, all figures and tables should be stand alone.

The scenarios were original described in the legend but during the technical review the Editor asked us to reduce the length of caption and describe them in a Table.

Emissions from prescribed fire in temperate forest in south-east Australia: implications for carbon accounting

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Abstract

We estimated emissions of carbon, as CO_2-equivalents, from planned fire in four sites in a south-eastern Australian forest. Emission estimates were calculated using measurements of fuel load and carbon content of different fuel types, before and after burning, and determination of fuel-specific emission factors. Median estimates of emissions for the four sites ranged from 20 to 139 Mg $CO_{2\text{-e}}$ ha^{-1}. Variability in estimates was a consequence of different burning efficiencies of each fuel type from the four sites. Higher emissions resulted from more fine fuel (twigs, decomposing matter, near-surface live and leaf litter) or coarse woody debris (CWD; >25 mm diameter) being consumed. In order to assess the effect of declining information quantity and the inclusion of coarse woody debris when estimating emissions, Monte-Carlo simulations were used to create seven scenarios where input parameters values were replaced by probability density functions. Calculation methods were: (1) all measured data were constrained between measured maximum and minimum values for each variable; (2) as for (1) except the proportion of carbon within a fuel type was constrained between 0 and 1; (3) as for (2) but losses of mass caused by fire were replaced with burning efficiency factors constrained between 0 and 1; and (4) emissions were calculated using default values in the Australian National Greenhouse Accounts (NGA), National Inventory Report 2011, as appropriate for our sites. Effects of including CWD in calculations were assessed for calculation Method 1, 2 and 3 but not for Method 4 as the NGA does not consider this fuel type. Simulations demonstrate that the probability of estimating true median emissions declines strongly as the amount of information available declines. Including

CWD in scenarios increased uncertainty in calculations because CWD is the most variable contributor to fuel load. Inclusion of CWD in scenarios generally increased the amount of carbon lost. We discuss implications of these simulations and how emissions from prescribed burns in temperate Australian forests could be improved.

1 Introduction

Fire affects the carbon balance of terrestrial biomes by immediately releasing carbon dioxide (CO_2), carbon monoxide (CO), methane (CH_4), volatile organic compounds (VOCs) and particulate matter (PM) into the atmosphere through the consumption of fuel (e.g. Urbanski et al., 2009) and by modifying carbon stocks in post-fire vegetation. Immediate modification of carbon stocks results from combustion of fuels while post-fire changes are due to alteration in activity of microorganisms responsible for decomposition of organic matter and uptake of CO_2 via photosynthesis by vegetation regrowth. Over the period 1997-2009, global fire emissions were estimated to contribute, on average, 2 Pg C yr^{-1} to the atmosphere, with 15% of those emissions coming from extra-tropical fires (van der Werf et al., 2010). Australia contributes about 6.7% of the global fire emissions, the fourth largest contributor behind Africa (51.6%), South America (14.5%), and Equatorial Asia (9.5%) (van der Werf et al., 2010). A recent study estimated that fires in Australia contribute 127 Tg C yr^{-1} to the atmosphere, about 6% of the net primary productivity with the greatest contribution coming from fires in tropical and savanna bioclimatic regions (Haverd et al., 2013). In contrast, contributions from cool and warm temperate bioclimatic regions to total annual fire emissions were limited except during severe bushfire seasons (Haverd et al., 2013).

Emissions from fires are still widely estimated as products of fuel load, burning efficiency, area burnt and emission factors for gases and particles of interest (Seiler and Crutzen, 1980; Langmann et al., 2009). Uncertainties in any of these variables can lead to a wide range of estimates for different gases. In large part these uncertainties are a function of burning efficiency and vegetation characteristics (e.g. Stropiana et al., 2010), and spatial and temporal scales of measurement (e.g. Urbanski et al., 2011). Techniques such as LIDAR are being used to improve estimates of fuel load (e.g. Loudermilk et al., 2009). Even so, fuel accumulation varies widely in space and time as a result of the interaction of many factors such as topography, soils, disturbance history (e.g. previous land use, insects, fire) and climate (e.g. due to variations in rainfall patterns; Bradstock, 2010); hence, remote sensing techniques will

require intensive calibration. In Australia, estimates of emissions from forest fires are based on fine fuels (e.g. grass, leaves, bark and twigs) and tend to ignore fuel types such as coarse woody debris (CWD) or understorey fuels (Volkova and Weston, 2013). A more comprehensive set of fuel load measurements is required to develop reliable fuel load models.

A major source of uncertainty in estimates has been emission factors as they invariably contain large uncertainties (Andreae and Merlet, 2001; Akagi et al., 2011; Urbanski et al., 2011). Published emission factors for forests in south-east Australia are few. One study developed emission factors for a small set of gases directly using aircraft-based sampling (Hurst et al., 1996), while another used Fourier transform infrared spectroscopy at ground level (Paton-Walsh et al., 2014). Ground-based spectrometry or satellite-derived enhancement ratios have also been used to derive emission factors (Paton-Walsh et al., 2004; Paton-Walsh et al., 2005; Young and Paton-Walsh, 2011; Glatthor et al., 2013). These non-direct methods often use as a reference, an emission factor for CO. However, that factor too is often assumed rather than measured. Compared to emission measurements made for savanna and grassland in Australia (e.g. Hurst et al., 1994a; Hurst et al., 1994b; Paton-Walsh et al., 2010), emission factors from Australian temperate forests are usually aggregated for all fuel types and do not account for factors such as fire severity and patchiness (cf. Russell-Smith et al., 2009). There have been no studies of seasonal variation in emission factors in Australian forests or demonstration that such variation is minimal, as found for savanna in Australia for certain trace gases (Meyer et al., 2012).

Return frequencies of wildfires in extra-tropical (temperate) forests in Australia are typically longer than that of tropical grassland and savanna and are often decadal compared to annual and biannual (Bradstock, 2010; Adams, 2013). In addition, the total area of temperate forest burnt on an annual basis is considerably smaller (Russell-Smith et al., 2007), notwithstanding large single fire events (Adams, 2013). Planned or prescribed burning in temperate forests to mitigate risks to life and property from wildfires is used at moderate return frequencies (e.g. 7 – 10 years) (Penman et al., 2007; McCaw, 2013). Bennett et al. (2013) recently demonstrated that in a mixed species eucalypt forest, repeated prescribed burning at shorter intervals (e.g. 3 - 5 years) reduces tree-based carbon stocks. The generality of such findings requires further research, as does the fate of the carbon released during combustion. Among the few indirect analyses of emissions from temperate forests (based on changes in litter and biomass C), Volkova and Weston (2013) estimated that 6.7 Mg C ha^{-1} was emitted to the atmosphere from

prescribed burning in *Eucalyptus obliqua* forests in south-east Australia. However, there remains a general paucity of direct empirical data on emissions, and this impedes efforts to calibrate indirect estimates.

Here we present emission factors for different fuel types from a temperate *Eucalyptus* forest in south-east Australia and use these in conjunction with measurements of fuel load and carbon content to estimate emissions from this forest type. We compare our estimates to those made using more restricted datasets and based upon the methodology described in the Australian National Greenhouse Gas Inventory Report 2011 (DIICCSRTEE, 2013) and discuss the merits of the different approaches.

2 Materials and methods

2.1 Study sites

The general study area was located in East Gippsland, Victoria, Australia (37°42′ 0″ S, 148° 27′ 0″). The elevation of study sites range from 56 to 124 m above sea level and the study area has an average annual precipitation of 850 mm. Sites were selected using the Victorian Department of Environment and Primary Industries (DEPI; Victoria, Australia) fire operations plans for the area. Three sites west of Orbost were burnt in planned fires in 2011 and one site east of Orbost was burnt in a planned fire in 2012. The selected sites were named according to the nearest crossroad or location: Oliver, Pettmans, South Boundary and Upper Tambo. All sites are classified as Lowland Forest (Ecological Vegetation Class 16; Victoria Department of Sustainability and Environment, 2004). Sites varied in overstorey tree species composition although all were dominated by Yellow Stringybark (*Eucalyptus muelleriana* A.W.Howitt), White Stringybark (*E. globoidea* Blakely) or Yerchuck (*E. consideniana* Maiden). The understorey vegetation in the western sites (Pettmans, South Boundary and Upper Tambo) is dominated by Sunshine Wattle (*Acacia terminalis* (Salisb.) J.F.Macbr.), Black Wattle (*A. mearnsii* De Wild.) and Burgen (*Kunzea ericoides* (A.Rich.) Joy Thomps.) with Bracken (*Pteriduim esculentum* (G.Forst.) Cockayne) as the most common groundcover species. The eastern site (Oliver) was selected primarily because the understorey composition differed from the western sites. Here the understorey is dominated by Forest Geebung (*Persoonia silvatica* L.A.S.Johnson) and Sunshine Wattle (*A. terminalis*) with Wire Grass (*Tetrarrhena juncea* R.Br.) as groundcover. Soils at all sites were formed on Pliocene (2-5 Ma) sands and gravels (Hendrickx et al., 1996; Van den Berg et al., 1996).

2.2 Sampling protocol

2.2.1 Overstorey and understorey biomass

Within each study site, three permanent circular plots were established at least 500 m apart in similar vegetation type. Due to the small elevation change of the general study area, all study sites had similar slope and aspect. Plots were located close to the road (20-50 m) to ensure they were burnt during the planned fire and were circular in shape (22.5 m radius; 1590.4 m^2). A schematic of the plot and sampling design is shown in Figure 1. All pre-fire data were collected 1-3 months prior to the planned burning and post-fire data were collected within 1 month of burning. Diameter at breast height over bark (DBHOB; 1.3 m) and number of individuals of trees in two size classes (≥2 cm to <20 cm; ≥20 cm) were measured for all overstorey (whole plot) and understorey tree species found in four circular subplots (radius = 5 m) located 5 m along the north-south and east-west axes of each of the larger plots, as measured from the centre point. At least six trees per plot were measured for tree height to provide a representative stand height.

To determine aboveground biomass and carbon stocks represented by overstorey and understorey trees (equivalent to overstorey and intermediate tree canopy fuel layers, respectively in Gould et al., 2011), Understorey allometric equations were developed for Yellow Stringybark (*E. muelleriana*; n = 10 individuals harvested) and Silver Wattle (*Acacia mearnsii*; n = 11 individuals harvested) using destructive harvesting. When species-specific allometric equations were not available or could not be developed by destructive sampling (i.e. overstorey), equations from Bi et al. (2004) for the species with the most similar size and growth form were used instead. Tree diameter and density were measured before planned burning. Data for overstorey species of *Eucalyptus* were pooled to represent a single biomass component (hereafter referred to as 'Overstorey') and data for all other tree species were pooled to form a second biomass component (hereafter referred to as 'Understorey').

Ground layer vegetation (ground cover of grasses and Bracken; equivalent to the near-surface live fuel layer in Gould et al., 2011) together with any scattered small shrubs (equivalent to the elevated fuel layer in Gould et al., 2011), was collected by pruning at ground level four 1 m^2 quadrats, each located 17.5 m along the north-south and east-west axes of each plot, as measured from the centre point. Samples were dried to constant weight at 70°C and subsamples were ground and analysed for total carbon content (% dry weight) by combustion analysis (Elementar Vario Max CNS, Analysensysteme GmbH, Hanau, Germany). The mass

of ground layer vegetation, twigs and litter (see below) remaining after prescribed burning was measured in the same way using quadrats positioned 2-3 m from the position of the original quadrat to avoid the influence of biomass removal prior to prescribed burning.

2.2.2 Litter and coarse woody debris

Litter on the forest floor (<25 mm diameter; equivalent to the surface fuel layer in Gould et al., 2011) was collected from the same quadrats used for sampling near-surface live biomass. Samples were carefully collected from the soil surface to avoid contamination from the underlying mineral soil. Samples were dried to constant weight at 70°C, weighed and sorted into size fractions. Fractions included plant material that was <10 mm diameter (hereafter referred to as 'Decomposing litter'); twigs, wood and bark that was 10-25 mm diameter (hereafter referred to as 'Twigs'), and partial or whole leaves between 10-25 mm diameter (hereafter referred to as 'Leaf litter'). Samples were collected pre- and post-fire, dried at 70°C to constant weight and subsamples of the pre-fire fraction were ground and analysed for total carbon content (% dry weight) by combustion analysis (Elementar Vario Max CNS, Analysensysteme GmbH, Hanau, Germany).

The volume of CWD was determined using the line intersect method (Van Wagner, 1968), where the north-south and east-west axes of each plot were used as transects (45 m each). The diameter and length and state of decomposition (sound or rotten) of all pieces of CWD (>25 mm diameter) intersecting each transect was measured. Subsamples of sound and rotten CWD were used to determine specific gravity (Ilic et al., 2000) and dried pre-fire subsamples ground and analysed for total carbon content (% dry weight) by combustion analysis (Elementar Vario Max CNS, Analysensysteme GmbH, Hanau, Germany). The volume of CWD was determined before and after planned burning.

2.3 Combustion analysis

A ventilation-controlled Mass Loss Calorimeter (MLC; Fire Testing and Technology, East Grinstead, UK) with a porous holder was used for the combustion analysis. The MLC consisted of a conical heater and a load cell to measure the change in mass of a sample over time. The cone heater and load cell were contained within a stainless steel enclosure, which was supplied with compressed air at a known flow rate of 140 L min^{-1}. A 90 cm tall, 12 cm

diameter stainless steel chimney on top of the enclosure contained a gas sampling ring probe mounted 60 cm above the enclosure. Air was drawn through the gas sampling ring at 2 L min^{-1} into a stainless steel housing (Model H130; Head line Filters, Aylesford, UK) containing a silica-bonded borosilicate glass microfibre filter (Head Line Filters, Aylesford, UK) and heated to 200°C to remove PM from the airstream. Air movement continued from the heated filter via a heated line (200°C) into a sampling manifold. Air in the sampling manifold was diluted with ambient air, filtered through a 1 micron PTFE filter (Pall Australia Pty. Ltd., Cheltenham, Australia) and pumped into the manifold to ensure that gas concentrations in the manifold were within the linear range of the various analysers used. Flow rates from the sample and dilution line were controlled by mass flow controllers (Aalborg, Orangeburg, US). The air temperatures in the manifold and stainless steel chimney were measured at 1 Hz using type K thermocouples connected to a digital acquisition board (Model NI USB-9211A; National Instruments, Sydney, Australia).

Mixing ratios of CO_2 and CO were measured at 1 Hz using non-dispersive infra-red gas analysers (Models 410i and 48i; Thermo Fisher Scientific Australia Pty. Ltd., Melbourne, Australia) and were calibrated using high purity CO_2 or CO diluted in zero air (BOC Ltd., North Ryde, Australia).

In the MLC, a sample holder (10 × 10 × 3 cm) with a porosity of 27% was used to allow diffusion of air through the samples. For all material, samples were trimmed to fit the holder to uniformly fill the sample holder so that the sample thickness was maintained at approximately 3 cm. The mass of the samples were recorded before burning and the mass of the residue after burning. The bulk density of the sample (kg m^{-3}) was calculated as the initial sample mass divided by the volume of the sample holder. The moisture content (MC) of combusted samples (dry weight basis), determined by drying at 70°C until constant weight, ranged between 2-14%. Samples were combusted in triplicate at an irradiance of 25 kW m^{-2} and a 10 kV spark ignitor was used to provide piloted ignition. A schematic of the equipment used for the combustion analysis is provided in Supplementary Figure 1.

2.4 Emission factors

Emission factors for the gas species CO_2 ($EFCO_2$) and CO (EFCO) from each fuel (biomass) type were calculated in g kg^{-1} dry fuel burnt. The mass of CO_2 or CO released was calculated

by summing products of excess CO_2 or CO concentrations and flow rate measured at each time step for the duration of the burn.

Using the carbon-mass balance method approach described by Radke et al. (1988) and outlined in Hurst et al. (1994b), emission factors for each fuel type were also expressed relative to elemental carbon content of dry fuels (g C g C^{-1}). The $EFCO_2$ was calculated from the fraction of total fuel carbon released to the atmosphere during combustion and CO_2-normalised emission ratios of CO, CH_4, non-methane hydrocarbons (VOC) and PM. $EFCO_2$ was calculated as:

$$EFCO_2 = \frac{\Delta CO_2}{C_{fuel}} = \frac{\frac{\Sigma C_{emit}}{C_{fuel}}}{1 + \frac{\Delta CO}{\Delta CO_2} + \frac{\Delta CH_4}{\Delta CO_2} + \frac{\Delta \Sigma NHMC}{\Delta CO_2} + \frac{\Delta PM}{\Delta CO_2}} \quad (1)$$

where ΣC_{emit} is the mass of carbon released to the atmosphere during burning and C_{fuel} is the initial carbon content of the fuel. Therefore, ΣC_{emit}/ C_{fuel} represents the fraction of fuel carbon that is burned and released to the atmosphere during combustion. Δ represents the excess molar mixing ratio of a species (CO_2, CO, CH4, ΣVOC and PM) over the background (the difference between its mixing ratios in smoke and clean air) (Hurst et al., 1994b). Emission factors (g C g C^{-1}) for carbon-based species other than CO_2 were calculated as:

$$EF_x = \frac{\Delta X}{\Delta CO_2} \times n \times EFCO_2 \quad (2)$$

where ΔX is the excess mixing ratio of species X (CO, CH4, ΣVOC or PM) and n is the number of carbon atoms per molecule of species X. By definition, the sum of the emission factors for the carbon gases and PM, when measured on a g C g C^{-1} basis, will equal ΣC_{emit}/ C_{fuel}.

Emission factors measured relative to elemental carbon content can be converted to emission factors (g kg^{-1} dry fuel) using Equation (3):

$$EF_x\left[gX\ kg^{-1}\ fuel\right] = \frac{EF_x\left[gC\ gC^{-1}\right] \cdot C_{fuel}}{\left(12 / Mw_x\right)} \times 1000 \quad (3)$$

where Mw_x is the molecular weight (g mol^{-1}) of chemical species X and 12 is the molecular weight of carbon.

In this study, CH_4, VOC and PM concentrations were not measured and hence the CO_2-normalised emission ratios of these compounds are not available for the direct calculation of EFCO2 according to Equation (1). Using $EFCO_2$ (g CO_2 kg^{-1}), $EFCO_2$ (g C g C^{-1}) was solved for each fuel type by re-arranging Equation (3). This allowed for calculation of EFCO (g C g C^{-1}) using Equation (2) and known [CO]/[CO_2] ratios. As the sum of emission factors for carbon gases and PM, when measured on a g C g C^{-1} basis, will equal $\Sigma C_{emit}/ C_{fuel}$, CH_4, VOC and PM were treated as pooled species (Σ(CH_4, VOC, PM)). $\Sigma C_{emit}/ C_{fuel}$ ratios were measured for each fuel fraction by subtracting the mass of carbon remaining in the ash after combustion from the amount of carbon measured before combustion. The excess Σ(CH_4, VOC, PM) to excess CO_2 ratio was then solved through optimisation (MS Excel v.14; Microsoft Corporation, Redmond, US) in order to make the sum of $EFCO_2$, EFCO and $EF\Sigma$(CH_4, VOC, PM) equal to the measured $\Sigma C_{emit}/ C_{fuel}$. This method assumes that the value of *n* used in Equation (2) in order to calculate $EF\Sigma$(CH_4, VOC, PM) is equal to one.

2.5 Emission calculations

Emissions, in terms of CO_2-equivalents (E_j; Mg $CO_{2\text{-e}}$ ha^{-1}), from each plot at each site (*j*) were calculated as the sum of the emissions from each fuel (biomass) class (*k*) for each carbon species (*x*):

$$E_j = \sum_{xk} EF_{xjk} \left(C_{fuel\,jk} \times \left(m_{pre_{jk}} - m_{post_{jk}} \right) \right) \times 3.66 \qquad (4)$$

where m_{pre} and m_{post} are the fuel loads (Mg ha^{-1}) before and after burning and 3.66 is a conversion factor from C to CO_2. C_{fuel} for CWD was assumed to equal that measured from twigs (<25 mm diameter).

Emissions can also be calculated using Equation (4) but by substituting m_{pre} - m_{post} with the product of the pre-fire fuel load and a burning efficiency factor (BEF).

$$E_j = \sum_{xk} EF_{xjk} \left(C_{fuel\,jk} \times m_{pre_{jk}} \times BEF_{jk} \right) \times 3.66 \qquad (5)$$

The BEF is defined as the mass of fuel that is exposed to fire that is pyrolysed (Russell-Smith et al., 2009). It is determined from the mass of fuel (m_{pre}) before combustion and the mass of the unburnt fuel residue and ash remaining after combustion (m_{post}):

$$BEF = 1 - \frac{m_{post}}{m_{pre}} \quad (6)$$

Equation (5) was used to calculate emission estimates for the sites as described in the Australian National Greenhouse Gas Inventory Report 2011 (AUSNIR; DIICCSRTEE, 2013) for a prescribed burn. Default values for the parameters in Equation (5) are described in AUSNIR as: emission factors are taken from Hurst et al. (1996) ($\Sigma C_{emit}/C_{fuel}$ = 0.9684), C_{fuel} is 0.5, BEF is 0.42 and the fuel load is 17.9 Mg ha^{-1}.

2.6 Uncertainty analysis of emission calculations

We completed seven different Monte-Carlo simulations for each site, in which input parameters were replaced by normally distributed probability density functions (PDFs). Table 1 outlines for the seven different scenarios the equation used to do the calculations (Equation 4 or Equation 5), the range of the values used for each input parameter (for each fuel fraction and site) and whether coarse woody debris was included in the calculations. Scenario 7 used the default fuel load applicable to these sites from the Australian National Greenhouse Accounts, National Inventory Report 2011 (DIICCSRTE, 2013). A priori analysis of the initial number of iterations for each Monte-Carlo simulation needed to produce an analysis where the true mean of the distribution lies within 1% of the estimate were made before each simulation. The maximum estimated number of simulations for any one set of sites and scenario was 71,233. The true error of the estimated mean for each site and scenario was always less than 1%. Results of the simulations are expressed as 95% uncertainty ranges defined by the 2.5 and 97.5 percentiles. The simulations were performed using Microsoft Excel 2010 (Microsoft Corporation, Redmond, WA, USA).

2.7 Statistics

Linear mixed models (IBM SPSS Statistics, v. 21.0; IBM, Armonk, US) were used to analyse effects of fire on fuel (biomass) type; with site, plot and fuel type as subject variables and time as the repeated variable. Time, site and time × site interactions were used as fixed effects. Fuel loads for the different types of fuel (i.e. twigs, decomposing matter, near-surface live, leaf litter, CWD, understorey and overstorey), before and after burning, carbon content, $\Sigma C_{emit}/C_{fuel}$, and emission factors were analysed with linear mixed models where site, plot and

fuel type were subject variables. Site, fuel type and site × fuel type interactions were used as fixed effects. The Bonferroni test was used for pairwise comparisons of the site and fuel type factors. Carbon content, $\Sigma C_{emit}/C_{fuel}$, and the emission factors were arc-sine transformed to meet assumptions of normality and homogeneity of variance.

3 Results

3.1 Fuel load and carbon content

Total fuel load before planned burning ranged from 61.7 ± 15.3 Mg ha^{-1} (mean ± standard deviation) at South Boundary to 111.3 ± 26.2 Mg ha^{-1} at Upper Tambo but were not significantly different among sites (Linear mixed model; P = 0.303). There was 10-fold more CWD than all other fuel types at all sites (P <0.001; Table 2). Masses of all remaining fuel types at each site were similar (less than 8 Mg ha^{-1}; P = 1.000) and there were no significant site × fuel type interactions (P = 0.692). After burning, total fuel loads at all sites were significantly reduced (P <0.001) and ranged from 20.1 ± 7.2 Mg ha^{-1} at Upper Tambo to 97.2 ± 24.7 Mg ha^{-1} at Oliver (Table 2). Reductions in fuel load due to burning were not consistent, resulting in significant time × site (P = 0.025) and time × fuel type interactions (P = 0.003; Table 2; Fig. 1). Time × site interactions resulted mainly from an 80% reduction in total fuel load at Upper Tambo, but only a 10% reduction at Oliver (Fig. 1). Fuel loads at Pettmans were reduced by an average of 28% and at South Boundary by 40% (Fig. 1). A significant time × fuel type interaction was expected given small reductions in CWD mass after burning compared to other fuel types (P = 0.002; Table 2; Fig. 1). Even so, there were significant differences in amounts of CWD burnt among sites. At Oliver, Pettmans and South Boundary, amounts of CWD biomass consumed were significantly less than at Upper Tambo (P = 0.017; Table 2; Fig. 1).

Twig mass (up to 8 Mg ha^{-1} pre-burn) was significantly reduced by burning (P <0.001) with an average loss of close to 5 Mg ha^{-1}. There were no time × site interactions (P = 0.656) but the mass of twigs measured at Oliver was significantly greater than at Upper Tambo both before and after burning (P = 0.05; Table 2; Fig. 1). Burning significantly reduced the mass of decomposing matter at all sites (up to 7 Mg ha^{-1} pre-burn) by almost 5 Mg ha^{-1} (P <0.001). Reductions in mass were greater at Pettmans, South Boundary and Upper Tambo than at Oliver. Again, there was a significant time × site interaction (P = 0.007).

Fuel loads represented by the ground layer vegetation (up to 0.6 Mg ha^{-1} pre-burn for Pettmans, South Boundary and Upper Tambo) were significantly less after burning (P = 0.002; Table 2; Fig. 1). There were significant site × time interactions (P = 0.004) as a consequence of substantially greater amounts of such vegetation at Oliver before burning (3 Mg ha^{-1}) than any of the other sites. None or very little of this fuel type remained after burning. Fire strongly reduced the mass of leaf litter and there were no major differences among sites before and after burning (2-9 Mg ha^{-1}; P = 0.398; Table 2).

Understorey biomass was not significantly different after burning compared to before burning at all sites (P = 0.392), but was significantly different among sites (P = 0.001). Understorey biomass at Oliver was significantly greater (nearly 2 Mg ha^{-1} pre-burn) than at any of the other sites before and after burning (P = 0.001 to 0.013). Overstorey biomass was significantly different among sites before (ranging from 6-15 Mg ha^{-1}; P <0.001) and after burning (ranging from 2-12 Mg ha^{-1}; P = 0.009). There was no interaction between site and time (P = 0.167). Understorey fuel loads at all sites decreased after burning by a little more than 1 Mg ha^{-1}.

Mean carbon contents of decomposing matter (30 ± 2%) were significantly less than of other fuel types at all sites (Linear mixed model; P <0.001; Table 2). Carbon contents of all other fuel types were in a narrow range (45-56%) resulting in significant site × fuel type interactions (P = 0.009; Table 2).

3.2 Emission factors

Amounts of carbon lost to the atmosphere relative to amounts held in aboveground biomass (the so called 'fuel carbon') were similar among the four sites (Linear mixed model; P = 0.456; $\Sigma C_{emit}/C_{fuel}$; Table 3). For the four sites, the mean proportion of fuel carbon lost to the atmosphere was 86% with a 95% confidence interval range of 77-95%. There were significant differences among different fuel types (P <0.001). $\Sigma(C_{emit}/C_{fuel})$ was significantly less in decomposing matter compared to other fuels (P <0.001; Table 3). Twigs, CWD and understorey biomass had statistically similar $\Sigma(C_{emit}/C_{fuel})$ (P >0.05). These $\Sigma(C_{emit}/C_{fuel})$ were all less than those for ground layer, overstorey and leaf litter (P <0.04). The latter three fuel types had statistically similar $\Sigma(C_{emit}/C_{fuel})$ (P >0.05).

For the four sites, the mean proportion of carbon lost to the atmosphere in the form of CO_2 was 71% with a range of 65-80% (Table 4). In contrast, proportions of carbon lost to the atmosphere as CO were much smaller (2-4%). Emission factors for CO_2 were similar among

the four sites (P = 0.456) albeit with significant differences among different fuel types (P <0.001). Emission factors for CO_2 ranged from 0.43-1.00 g C g C^{-1}among the different fuel types. Twigs and leaf litter produced significantly smaller emission factors than decomposing matter and overstorey biomass (P <0.05). Emission factors for ground layer and understorey biomass were similar to those for twigs and leaf litter. Emission factors for CO were dependent on site × fuel type interactions (P = 0.026; Table 3). At South Boundary and Upper Tambo, emission factors for CO were greater for decomposing matter and ground layer fuels relative to the other types (P <0.05; Table 3). In contrast, at Oliver and Pettmans, decomposing material had greater emission factors for CO than other fuel types (P <0.026; Table 3).

Pooled emission factors for CH_4, VOC and PM (Σ(CH_4, VOC, PM); Table 4) were significantly different among sites (P = 0.002) and fuel types (P <0.001). Emission factors for Σ(CH_4, VOC, PM) for fuel collected from Upper Tambo were significantly less than fuels of other sites (P <0.049). As a consequence, the average proportion of carbon lost to the atmosphere as Σ(CH_4, VOC, PM) from the four sites ranged widely (13-23%). Differences in emission factors among fuel types was due to lesser emission factors for decomposing matter relative to all other fuel types and greater emission factors for leaf litter relative to understorey and overstorey biomass (P <0.017).

Carbon content of the different fuel types and ash (from the calorimeter) (Table S1), initial bulk density and residual mass fractions (Table S2), excess CO/CO_2 and excess Σ(CH_4, VOC, PM)/CO_2 ratios (Table S3) used to calculate the emission factors, on both a mass of compound released per unit of fuel mass burnt and on a carbon mass balance basis, can be found in the supplementary material.

3.3 Emission estimates

Results of the Monte-Carlo simulations of estimated emissions from the four sites, using seven different calculation scenarios, are shown in Fig. 2. Scenario 1 produced symmetrically distributed estimates, with median estimates ranging from close to 20 Mg $CO_{2\text{-}e}$ ha^{-1} for Oliver to 139 Mg $CO_{2\text{-}e}$ ha^{-1} for Upper Tambo. If CWD was omitted (Scenario 2), distributions were narrower and median estimates were reduced. The reduction in the median estimate varied among sites; for Oliver the reduction was 3%, Pettmans 34%, South Boundary 38% and Upper Tambo 71%.

Scenario 3 produced positively skewed distributions for all sites and reduced median estimates (by 40-54% from Scenario 1). Outputs of Scenario 4 (Scenario 3 excluding CWD) were similarly positively skewed, but more narrowly distributed. Relative to Scenario 1, excluding CWD lowered median estimated emissions by 53-83%. Relative to Scenario 3, such exclusion lowered median estimates by 4-69%. Scenario 5 produced the most positively skewed distributions for Oliver, Pettmans and South Boundary (Fig. 2). Consequently, the median estimate for Oliver was 90% greater than that of Scenario 1. Median estimates for other sites were between 16 and 76% less. When the same calculation method (Scenario 5) was applied, but excluding CWD data (Scenario 6), the distribution was still positively skewed but with a much narrower range (Fig. 2). The omission of CWD data in Scenario 6 resulted in a median estimate (relative to Scenario 1) being reduced by between 36 and 91% across all sites.

Simulations for sites using default fuel load, carbon content and emission factors from the Australian National Greenhouse Accounts, National Inventory Report 2011 (AUSNIR; DIICCSRTE, 2013; hereafter referred to as Scenario 7) were highly positively skewed, with a median estimate of 4.5 Mg ha^{-1}. This is some 77-97% less than median estimates for the four sites from Scenario 1. The 95% confidence range of Scenario 7 ranged from 0.05 Mg ha^{-1} to more than 35 Mg ha^{-1} with a mean value of close to 8 Mg ha^{-1}. Using default values in AUSNIR, estimated mean total emission across all sites was 13.3 Mg ha^{-1}. This is in the upper quartile of estimates for Scenario 7.

Based on Scenario 7, the probability that emissions are less than the median calculated using Scenario 1 was 88% for Oliver, 96% for Pettmans and 97% for South Boundary. For Upper Tambo, emission estimates based on Scenario 1 were outside the range of those calculated under Scenario 7.

4 Discussion

There were large differences in mass (biomass plus litter) lost among the four sites due to prescribed fire. These differences were due to the differing abundances, and consumption during fire, of the different fuel types. Given planned burning aims especially to reduce the loads of fine fuels (e.g. twigs, decomposing matter, ground layer vegetation and leaf litter), the fires studied here achieved this goal with only small changes in understorey and overstorey biomass. Losses of mass from CWD accounted for much of the variation among

sites, especially when considered in proportion to losses from finer fuels. When expressed in terms of carbon content, losses of carbon from CWD at Pettmans and South Boundary (18-24%) were greater than from *Eucalyptus obliqua* forests of south-east Australia (Volkova and Weston, 2013), but consistent with the model results of Hollis et al. (2011). In contrast, fine fuel and CWD accounted for 79% of the C lost at the Upper Tambo site. The site east of Orbost (Oliver) lost the least amount of mass (and C), retaining most of its fine fuels and showing no appreciable change in CWD. Estimation of fuel load is a major source of uncertainty in any estimation of potential or actual fire emissions, and the large variability in burning efficiency across the sites used in this study is consistent with variability described by Stropiana et al. (2010) and Urbanski et al. (2011).

Across the four sites, the mean proportion of fuel carbon lost to the atmosphere relative to the total amount of carbon ($\Sigma C_{emit}/C_{fuel}$) was 86%. This is less than the 97% suggested by Hurst et al. (1996) for the one planned burn they measured in a south-east Australian forest. However, a direct comparison of this study with the Hurst et al. (1996) study cannot be made due to the significantly different methodological approaches taken that they may bias either study. These methodological differences include factors such as: the measurement of aggregated emissions from naturally structured fuels taken using an aircraft, compared to individual fuel components measured at a very small scale in the laboratory; and neither study measures the same range of compounds. Indeed, these methodological differences also prevent direct comparison of emission factors, not just with Hurst et al. (1996) but also the recent work of Paton-Walsh et al. (2014) who made ground-based emission measurements from planned fires in temperate south-east Australian fires. There are large variabilities in emission factors for certain compounds among different ecosystems (see reviews by Andreae and Merlet (2001) and Akagi et al. (2011)). This demonstrates the need for more comprehensive emissions measurements for specific ecosystems and regions, including south-east Australian forests. If these measurements are conducted in a manner similar to those for the south-eastern and south-western US (e.g. Yokelson et al., 2013), field and laboratory measurements may be reconciled.

Monte-Carlo simulations clearly demonstrated the significance of availability of data to accurate calculations of likely emissions. If only fuel load (before and after burning) is known and default values from AUSNIR are used, estimated emissions could vary from true emissions by as much as 100%. One characteristic common across all simulations was that

when data for CWD is included, the range of emissions increased strongly, as a result of large variation in mass of CWD among sites. In addition, there was wide variation among sites in consumption of CWD during prescribed fires . Emissions estimated using Scenarios 1, 2, 3 and 4, where fuel loads were known before and after burning, had greatly reduced variance. Distributions of estimated emissions were more positively skewed as the amount of data available declined. In other words, the probability of an estimate being in the low portion of the distribution is greatly increased, in addition to the diminished probability that the estimate matches the true emission. This is amply demonstrated by the distribution of estimates calculated using Scenario 7 (AUSNIR default values) which encompassed the median emissions estimate of Scenario 1 for three of the four sites. There was, at most, only 12% probability of matching values. For the fourth site, Scenario 7 could not produce a distribution that overlapped with that calculated using Scenario 1.

This study has shown that even within a single, well-defined vegetation type, there is wide variability in emissions principally because of different burning efficiencies among sites and fuel types. In order to improve both the accuracy and precision of estimated emissions from planned burning, the use of a single efficiency factor, as described in AUSNIR, is clearly insufficient. The methodology used to predict emissions from savanna and grassland, where burning efficiencies are described as a function of fuel type and fire severity (Russell-Smith et al., 2009), is only effective if fuel loads are accurately known (Stropiana et al., 2010; Urbanski et al., 2011). Spatial variability in fuel loads (Burgan et al., 1998; Keane et al., 2001) and the spatio-temporal variability in fuel conditions (Clinton et al., 2006) mitigate against such a scenario. We have shown that in addition to the mass of different fuel types, their carbon content plays a significant role in potential emissions. The Australian National Greenhouse Gas Inventory Report 2011 (DIICCSRTEE, 2013) assumes a 50% default value for carbon content of forest fuels. Fuel types in this study, with the exception of decomposing matter, had carbon contents ranging between 45% and 56%, mostly close to the default value. However decomposing matter had a much lower C content (average 30%). Combustion of fuels with low carbon contents could lead to overestimation of carbon loss. Considerable improvements in emissions estimates from temperate forests in south-eastern Australia could be made if a greater number of emission factors were available for different fuel types. This would eliminate current reliance on site-aggregated values and would aid in the development of predictive models for emission factors, particularly if different combustion conditions such as fuel moisture content, fuel load, fuel arrangement and fire intensity could be incorporated

(Yokelson et al., 1999; Andreae and Merlet, 2001; Possell and Bell, 2013). Field studies are still required to verify laboratory determined emission factors.

5 Summary

Planned fires in a temperate *Eucalyptus* forest in south-east Australia released between 20 to 139 Mg $CO_{2\text{-}e}$ ha^{-1}. Variability in the range of emissions was a consequence of different burning efficiencies among investigated fuel types, with greater emissions when appreciable amounts of CWD were burnt. Simulation of emissions showed that as the amount of information available to calculate emissions is reduced, the probability of estimating true emissions greatly diminishes. Ideally, measurement of fuel load and carbon content of different fuel types should be made before and after fire. In conjunction with emission factors for a greater range of fuel types and conditions, our ability to estimate of carbon loss from forests via prescribed burns would be greatly improved and would provide invaluable data on carbon apportionment for the calibration of fuel models.

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1 Table 1. Summary of parameters and range of values used to calculate emission estimates for seven different scenarios by Monte Carlo
2 simulation. Max. to min. refers to the maximum and minimum values recorded for each fuel type and site. CWD is coarse woody debris. See
3 text for further details of the equations used.
4

Scenario	Calculation equation	Parameters					
		Carbon content (%)	Emission factors (g C g C^{-1})	Mass loss (Mg ha^{-1})	Pre-burn fuel loads (Mg ha^{-1})	Burning efficiency factor	CWD included?
1	4	Max. to min.	Max. to min.	Max. to min.	-	-	Yes
2	4	Max. to min.	Max. to min.	Max. to min.	-	-	No
3	4	0 – 100	0 – 1	Max. to min.	-	-	Yes
4	4	0 – 100	0 – 1	Max. to min.	-	-	No
5	5	0 – 100	0 – 1	-	Max. to min.	0 – 1	Yes
6	5	0 – 100	0 – 1	-	Max. to min.	0 – 1	No
7	5	0 – 100	0 – 1	-	17.9	0 – 1	No

Table 2. Fuel load and pre-burn carbon content of a range of fuel types measured before and after fire in four forest sites in East Gippsland, south-eastern Australia. Values are mean ± standard deviation (n = 3).

Fuel type	Oliver			Pettmans		
	Fuel load (Mg ha^{-1})		Carbon content (% dry weight)	Fuel load (Mg ha^{-1})		Carbon content (% dry weight)
	Pre-burn	Post-burn	Pre-burn	Pre-burn	Post-burn	Pre-burn
Twigs	7.75 ± 1.65	3.70 ± 1.58	49.67 ± 0.15	5.23 ± 1.31	0.01 ± 0.01	48.78 ± 0.88
Decomposing matter	3.11 ± 0.57	2.03 ± 2.01	29.79 ± 6.04	5.69 ± 1.36	0.02 ± 0.01	23.87 ± 7.05
Ground layer	3.31 ± 1.57	0.02 ± 0.03	46.68 ± 0.08	0.62 ± 0.33	0	46.74 ± 1.36
Leaf litter	1.85 ± 0.59	1.25 ± 0.17	54.95 ± 0.31	2.80 ± 0.29	0.27 ± 0.13	52.35 ± 1.92
Coarse woody debris	75.91 ± 19.64	76.43 ± 21.73	49.67 ± 0.15	61.14 ± 55.33	53.11 ± 58.08	48.78 ± 0.88
Understorey	1.78 ± 1.50	1.69 ± 1.48	53.53 ± 0.36	0.80 ± 0.54	0.76 ± 0.49	53.53 ± 0.36
Overstorey	14.87 ± 4.32	12.08 ± 3.17	54.95 ± 0.31	3.73 ± 1.40	3.38 ± 1.80	54.95 ± 0.31
	South Boundary			Upper Tambo		
Twigs	5.32 ± 0.67	0.07 ± 0.03	49.59 ± 0.42	5.91 ± 0.68	0.06 ± 0.02	49.14 ± 1.26
Decomposing matter	6.89 ± 0.23	0.05 ± 0.02	32.13 ± 2.69	5.94 ± 1.05	0.03 ± 0.01	35.42 ± 2.06
Ground layer	0.33 ± 0.18	0	47.72 ± 1.85	0.11 ± 0.06	0	47.57 ± 0.94
Leaf litter	4.25 ± 0.82	0.37 ± 0.11	53.55 ± 2.45	9.49 ± 10.56	0.30 ± 0.18	53.70 ± 1.69
Coarse woody debris	41.66 ± 16.39	33.35 ± 15.00	49.59 ± 0.42	83.70 ± 37.29	14.56 ± 5.99	49.14 ± 1.26
Understorey	0.52 ± 0.37	1.01 ± 0.22	53.53 ± 0.36	0.10 ± 0.17	0.29 ± 0.49	53.53 ± 0.36
Overstorey	2.78 ± 1.41	2.12 ± 0.91	54.95 ± 0.31	6.07 ± 1.95	4.89 ± 1.40	54.95 ± 0.31

Table 3. Proportion of the fuel carbon burned emitted into the atmosphere from different fuel types from forest sites in East Gippsland, south-eastern Australia. C_{emit} is the total carbon emitted into the atmosphere through combustion and C_{fuel} is the initial carbon content of fuel. Coarse woody debris was assumed to have the same values as twigs. Values are mean ± standard deviation (n = 3).

	Oliver	Pettmans	South Boundary	Upper Tambo
Fuel type	$\Sigma C_{emit}/C_{fuel}$			
Twigs	0.882 ± 0.015	0.819 ± 0.043	0.844 ± 0.026	0.857 ± 0.060
Decomposing matter	0.710 ± 0.177	0.558 ± 0.342	0.751 ± 0.136	0.632 ± 0.090
Ground layer	0.978 ± 0.009	0.960 ± 0.017	0.948 ± 0.058	0.986 ± 0.009
Leaf litter	0.957 ± 0.013	0.975 ± 0.025	0.956 ± 0.035	0.915 ± 0.019
Understorey	0.859 ± 0.054	0.859 ± 0.054	0.859 ± 0.054	0.859 ± 0.054
Overstorey	0.942 ± 0.014	0.942 ± 0.014	0.942 ± 0.014	0.942 ± 0.014

Table 4. Emissions factors for CO_2, CO and pooled CH_4, volatile organic compounds (VOC) and particulate matter (PM) for different fuel types from forest sites in East Gippsland, south-eastern Australia, that were combusted in a mass-loss calorimeter. Coarse woody debris was assumed to have the same values as twigs. Values are mean ± standard deviation (n = 3).

	Oliver			Pettmans		
	Emission factor [g C g C^{-1}]			Emission factor [g C g C^{-1}]		
Fuel type	CO_2	CO	ΣCH_4, VOC, PM	CO_2	CO	ΣCH_4, VOC, PM
Twigs	0.59 ± 0.03	0.02 ± 0.01	0.28 ± 0.05	0.58 ± 0.03	0.02 ± 0.01	0.23 ± 0.04
Decomposing matter	0.87 ± 0.13	0.06 ± 0.02	0.05 ± 0.08	1.00 ± 0.08	0.06 ± 0.01	0
Ground layer	0.62 ± 0.02	0.03 ± 0.01	0.35 ± 0.02	0.58 ± 0.04	0.03 ± 0.01	0.37 ± 0.05
Leaf litter	0.53 ± 0.03	0.02 ± 0.01	0.42 ± 0.02	0.56 ± 0.07	0.03 ± 0.01	0.40 ± 0.06
Coarse woody debris	0.59 ± 0.03	0.02 ± 0.01	0.28 ± 0.05	0.58 ± 0.03	0.02 ± 0.01	0.23 ± 0.04
Understorey	0.66 ± 0.19	0.02 ± 0.02	0.20 ± 0.15	0.66 ± 0.19	0.02 ± 0.02	0.20 ± 0.15
Overstorey	0.79 ± 0.06	0.03 ± 0.01	0.14 ± 0.06	0.79 ± 0.06	0.03 ± 0.01	0.14 ± 0.06
	South Boundary			Upper Tambo		
Twigs	0.47 ± 0.02	0.02 ± 0.01	0.36 ± 0.02	0.70 ± 0.06	0.02 ± 0.01	0.15 ± 0.07
Decomposing matter	0.68 ± 0.05	0.03 ± 0.01	0.07 ± 0.08	0.89 ± 0.17	0.05 ± 0.01	0
Ground layer	0.69 ± 0.18	0.04 ± 0.01	0.23 ± 0.12	0.74 ± 0.03	0.05 ± 0.01	0.22 ± 0.03
Leaf litter	0.65 ± 0.07	0.02 ± 0.01	0.29 ± 0.10	0.68 ± 0.04	0.03 ± 0.01	0.22 ± 0.04
Coarse woody debris	0.59 ± 0.03	0.02 ± 0.01	0.28 ± 0.05	0.58 ± 0.03	0.02 ± 0.01	0.23 ± 0.04
Understorey	0.66 ± 0.19	0.02 ± 0.02	0.20 ± 0.15	0.66 ± 0.19	0.02 ± 0.02	0.20 ± 0.15
Overstorey	0.79 ± 0.06	0.03 ± 0.01	0.14 ± 0.06	0.79 ± 0.06	0.03 ± 0.01	0.14 ± 0.06

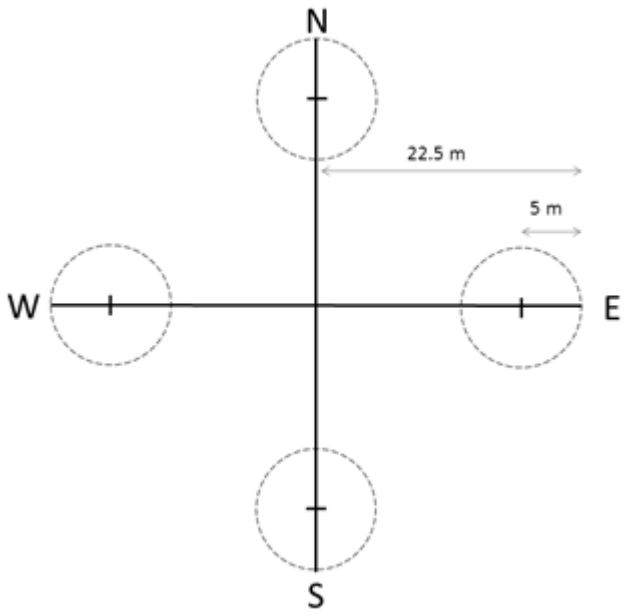

Figure 1. Plot layout for data and sample collection.

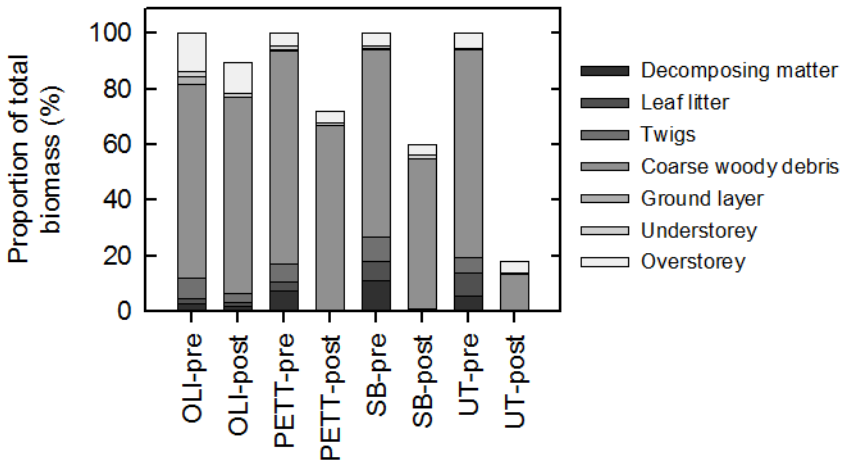

Figure 2. Proportion of the total biomass for each fuel type, at each site, before and after planned burning. The sites are: Oliver (OLI), Pettmans (PETT), South Boundary (SB) and Upper Tambo (UT). Each section of each bar represents the mean proportion measured from three plots within each site. 'Pre' and 'post' refer to measurements made before and after the planned burn.

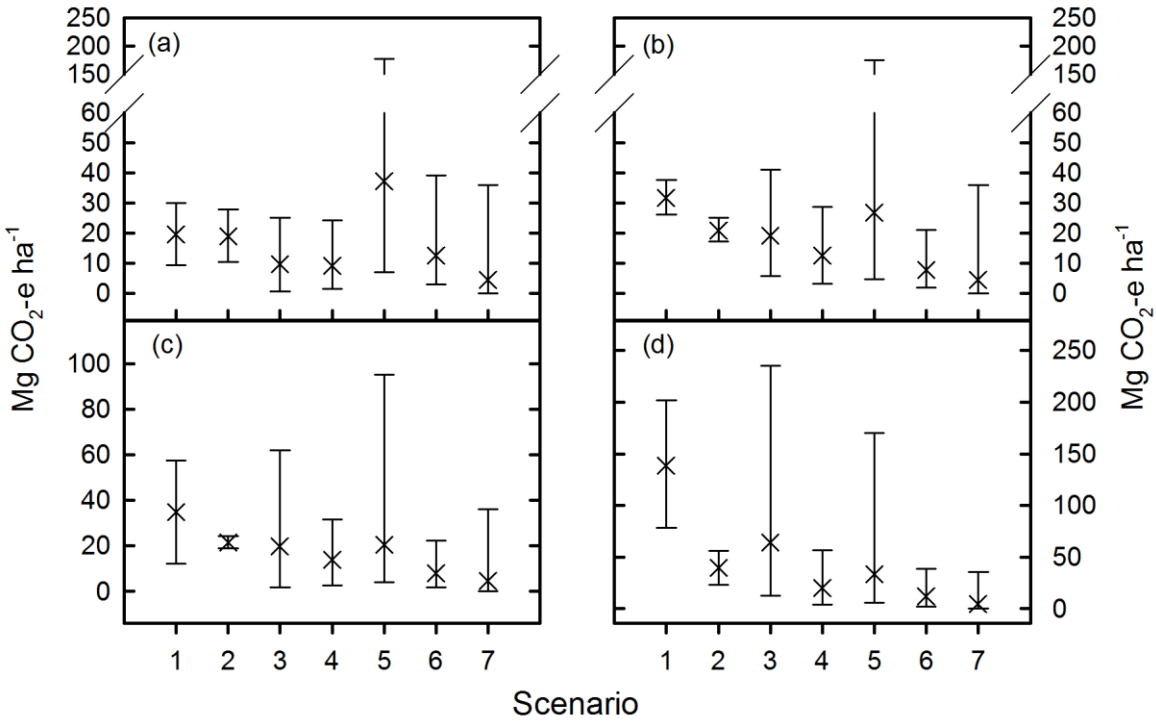

Figure 3. Estimates of CO_2-equivalent emissions from four forest sites in East Gippsland, south-eastern Australia using Monte-Carlo simulations of seven different scenarios. Sites are (a) Oliver, (b) Pettmans, (c) South Boundary, and (d) Upper Tambo. See Table 1 for description of the seven scenarios. Crosses represent the median emission as determined by the Monte Carlo simulations (n ≤71,233). The error bars represent the 95% confidence intervals of the Monte Carlo simulations.