Fibre wall and lumen fractions drive wood density variation across 24 Australian angiosperms

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Abstract. Wood density is considered a key plant trait, affecting mechanical and physiological performance, yet its biological meaning is still rather unclear. Accordingly we investigated the anatomical underpinnings of wood density in trees and shrubs. We measured wood density and anatomical traits in distal stems 4–10 mm diameter under bark in 24 Australian species. Proportions of wood components that are functionally distinct were analysed, including fibre wall and lumen, vessel wall and lumen, and axial and ray parenchyma. Wood density was mainly driven by the density of wood outside vessel lumens (density\textsubscript{NV}) rather than by vessel lumen fraction. In turn, density\textsubscript{NV} variation was chiefly affected by fibre wall and lumen fractions. Considerable anatomical variation was observed at a given density\textsubscript{NV}, especially among medium-density\textsubscript{NV} species (0.60–0.85 g cm\textsuperscript{-3}); this range of medium density\textsubscript{NV} roughly translates to 0.50–0.75 g cm\textsuperscript{-3} of overall density. The anatomy of these species formed a continuum from low fibre lumen and medium parenchyma fractions to medium fibre lumen and low parenchyma fractions. Our data suggest that wood density is an emergent property influenced by a complex anatomy rather than an unambiguous functional trait, particularly in medium-density species. With much anatomical variation, they likely represent a wide range of ecological strategies.

Keywords: Ecological strategies; fibres; parenchyma; rays; tissue fraction/proportion/percentage/volume; vessels; wood anatomy.

Introduction

Plants vary significantly in their ecological, physiological and mechanical properties or ‘traits’ both across climate and even within a site (Westoby et al. 2002; Wright et al. 2004; Chave et al. 2009). This indicates that there are multiple solutions to the problem of how to be a successful plant and maintain species continuity. Different solutions can be called plant ecological strategies (Westoby et al. 2002).

Wood density has been suggested as a key player in plant ecological strategies (Chave et al. 2009). Firstly, wood density has been linked with hydraulic strategies. Denser woods tend to operate at more negative water potentials (Ackerly 2004; Bucci et al. 2004; Santiago et al. 2004; Jacobsen et al. 2007b, 2008; Gotsch et al. 2010) and to have greater cavitation resistance than low-density woods (Hacke et al. 2001; Jacobsen et al. 2005; Pratt et al. 2007; Lens et al. 2011). Wood density has been studied in relation to hydraulic conductivity but the results are inconclusive, showing either a negative relationship between the two traits (measured conductivity, Stratton et al. 2000; Bucci et al. 2004; Santiago et al. 2004) or no relationship (theoretical conductivity, Poorter et al. 2010; Fan et al. 2011).
Also, density has been found to correlate negatively with capacitance (Meinzer et al. 2003, 2008; Pratt et al. 2007; Scholz et al. 2007). Secondly, wood density has been associated with plant mechanical strategies where denser woods tend to be stiffer and more resistant to breakage at a given wood diameter (Chave et al. 2009). However, it has also been suggested that plants can build thicker stems to compensate for their lower density (Anten and Schieving 2010; Larjavaara and Muller-Landau 2010; Butler et al. 2011). Thirdly, denser woods might be more resistant to pathogen attacks (Augspurger and Kelly 1984; Romero and Bolker 2008). Wood density has also been widely discussed in relation to life-history strategies. For example, species with denser wood tend to experience lower stem mortality rates (Putz et al. 1983; Kraft et al. 2010; Poorter et al. 2010). Growth rate is another important component of life history. Growth rate can be expected to negatively relate to wood density on the basis that higher investment in mass per volume should slow down growth (Enquist et al. 1999) and, while generally true, the correlation is not always strong (Poorter et al. 2008, 2010; Chave et al. 2009; Wright et al. 2010; Fan et al. 2012). Across species, wood density can vary with environmental factors such as temperature (Wiemann and Williamson 2002; Swenson and Enquist 2007; Martinez-Cabrera et al. 2009) and precipitation (Barajas-Morales 1985; Wiemann and Williamson 2002; Swenson and Enquist 2007; Martinez-Cabrera et al. 2009; Zhang et al. 2011), although not in all studies (Ter Steege and Hammond 2001; Wiemann and Williamson 2002; Muller-Landau 2004). Despite this broad climate-related patterning, wood density also tends to vary quite widely among co-occurring species (Wiemann and Williamson 2002; Muller-Landau 2004).

Thus there are many potential functional roles for wood density but also a number of unresolved questions about each potential role. Larjavaara and Muller-Landau (2010) argued that some observed correlations may not be causal, but rather may reflect correlated selection on other traits. Further, if wood density has multiple functions, then it might not be a very good predictor of any one of them. In any event, a useful step towards resolving this problem is to ask what are the structural underpinnings of wood density variation. The premise of the work reported here was that if structural underpinnings of wood density variation were rigorously quantified and better understood, this might help to explain complexities in the functional implications of wood density.

Most studies of anatomical components of angiosperm wood density span only one or a few species (Schulz 1957; Taylor 1971; Taylor and Wooten 1973; Ezell 1979; Vurdu and Bensend 1980; Fukuzawa 1984; Bosman et al. 1994; Stokke and Manwiller 1994; McDonald et al. 1995; Lei et al. 1996; Denne and Hale 1999; Rana et al. 2009) or focus on commercial woods (French 1923 as cited in Panshin and de Zeeuw 1980; Manwiller 1973 as cited in Koch 1985). Fewer studies make comparisons across a broad number of species (Fujisawa et al. 1991; Jacobsen et al. 2007a; Martinez-Cabrera et al. 2009; Poorter et al. 2010; Fichtler and Worbes 2012).

Wood is a complex tissue composed of three main cell types: vessels that transport water, fibres responsible for mechanical strength, and parenchyma that stores and transports nutrients. These tissues have different structural characteristics and their relative proportions within wood influence wood density. Vessel lumens have essentially zero density; fibre and vessel walls and parenchyma have positive density. Vessel fraction has variously shown either negative or no correlation with wood density (Preston et al. 2006; Jacobsen et al. 2007a; Mitchell et al. 2008; Martinez-Cabrera et al. 2009; Poorter et al. 2010; Zanne et al. 2010; Gleason et al. 2012). Parenchyma is another commonly occurring tissue, which has been reported to have positive, negative or no relationship with density (Taylor 1969; Fujisawa 1992; Jacobsen et al. 2007a; Martinez-Cabrera et al. 2009; Rana et al. 2009; Poorter et al. 2010). Wood density is generally well correlated with fibre properties, especially fibre wall fraction (Fujisawa et al. 1991; Jacobsen et al. 2007a; Martinez-Cabrera et al. 2009). However, it is unclear how these different fibre traits are interrelated with each other and consequently how these interrelations influence wood density.

Most previous work linking anatomy with density has concentrated on vessels with relatively little attention given to the other tissues. Furthermore, among the studies investigating all the major tissues (vessels, parenchyma and fibres) only one focused on the wood of twigs in 17 species studied by Jacobsen et al. (2007a). Twigs are important, being in direct spatial and functional contact with leaves and having been commonly subjected to physiological and ecological measurements. In this paper, we investigate wood from twigs from a wide range of angiosperm tree and shrub species growing in various environments (24 species from four sites in eastern Australia). We address two main unresolved issues: (i) Which fibre properties have the most decisive effect on wood density and how are those properties interrelated with each other? (ii) How do vessel and parenchyma proportions influence wood density?

Methods

Plant material and sites

Four sites were chosen that spanned a wide range of temperature and aridity in eastern Australia [see Supporting
Information]. The objective of site selection was to generate a broad range of trait values rather than to enable site comparisons. All carried natural, undisturbed vegetation growing on oligotrophic soils on flat to slightly sloping terrain. Two locations in Tasmania, at ~43°S, represented low mean annual temperature (MAT; 10.6 °C), and two locations in Queensland near 18°S represented higher MAT (c. 22.5 °C). Within each latitude, two locations were chosen so as to differ markedly in aridity index (AI; Willmott and Feddema 1992), the ratio of mean annual precipitation (MAP) to potential evapotranspiration (PET). In both Queensland and Tasmania the wetter site had an AI c. 1.0 and the drier site c. 0.6. MAP, MAT and PET were obtained from GIS (geographic information system) layers from the Australian Bureau of Meteorology.

At each of the four sites, six abundant and phylogenetically distinct woody eudicot species were chosen for sampling (species listed in Table 1). One species was sampled at two sites, yielding a total of 23 species from eight families. Distal, sun-exposed twigs of trees and shrubs were collected from three replicate individuals per species. The diameter under bark of twigs varied from 4 mm in plants with little pith to 10 mm in plants with higher pith content. Consequently the diameter of wood, excluding bark and pith, was 4–5 mm. This plant material is referred to here as twigs, although in several small shrub species ‘twigs’ were the main stems. Plant material was cut into segments 10–15 cm long and kept wet in sealed plastic bags in the refrigerator (4 °C). Wood density was measured within a week from collection; other parts of the same twigs were placed in fixative for later measurement of anatomical properties (details below).

Wood density

Wood density was measured on segments 3–5 cm long for each twig sample. Bark and pith were removed and measurements were carried out on xylem only. In this paper we refer to xylem as ‘wood’. After removing bark and pith, wood pieces were soaked in water for at least 48 h prior to volume measurement. Then a beaker filled with water was placed on a balance (0.0001 g, Mettler AE 160). A thin wire platform was suspended in water so that it did not touch the side or bottom of the beaker. The balance was tared before each measurement and a sample was gently placed on the platform. The mass of displaced water was read from the balance. From standard water density of 1 g cm\(^{-3}\) and knowing the mass of displaced water, we calculated sample volumes applying Archimedes’ buoyancy principle (e.g. 1 g of displaced water equals 1 cm\(^3\) volume). Samples were then placed in paper envelopes and dried at 70 °C for at least 72 h. Wood density was calculated as the dry mass divided by water saturated volume (g cm\(^{-3}\)).

Table 1. Sampled sites, species names and families.

<table>
<thead>
<tr>
<th>Site</th>
<th>Species name</th>
<th>Family</th>
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<tr>
<td>Cool-wet</td>
<td>Allocasuarina monilifera</td>
<td>Casuarinaceae</td>
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<td></td>
<td>Aotus ericoides</td>
<td>Fabaceae</td>
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<td></td>
<td>Banksia marginata</td>
<td>Proteaceae</td>
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<td></td>
<td>Eucalyptus amygdalina</td>
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<td></td>
<td>Leptospermum scoparium</td>
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<td></td>
<td>Leucopagon ericooides</td>
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<tr>
<td>Cool-dry</td>
<td>Bossiaea cinerea</td>
<td>Fabaceae</td>
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<td></td>
<td>Daviesia latifolia</td>
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<td>Epacris impressa</td>
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<td>Eucalyptus tenuiramis</td>
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<td>Alphitonia excelsa</td>
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<td></td>
<td>Persoonia falcata</td>
<td>Proteaceae</td>
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Anatomy

To obtain anatomical cross-sections the material was first fixed in formalin-acetic acid-alcohol (FAA) for 4 weeks. The FAA was prepared in proportions of 5 : 5 : 90 (formalin : glacial acetic acid : 70 % ethanol; Gerlach 1972). After 4 weeks, the fixative was replaced with 70 % ethanol. This 70 % ethanol was then replaced two more times within 10 days to further wash the fixative out. Final replacement of alcohol was used as a long-term storage medium. Segments for image analysis were rehydrated by immersion in 50 % ethanol and after 2–5 days in 30 % ethanol. Cross-sections were cut with a sledge microtome (Reichert, Vienna, Austria) at 10–20 μm thickness using disposable blades (Model A35, Feather Safety Razor Co. Ltd, Japan). For better contrast and tissue identification, sections were stained with safranin O (Gurr Microscopy Materials, BDH Chemicals Ltd, UK) for lignified regions.
cell walls (10 min) and with Janus green B (Gurr’s, London, UK) for cytoplasm (10 min). Safranin O solution used 2 g of stain in 100 mL of distilled water (Ruzin 1999) and Janus green B used 0.1 g of stain and 1 mL of glacial acetic acid in 100 mL of distilled water (Conn et al. 1960). Sections were rinsed in distilled water after each staining session. Afterwards, they were mounted in glycerol on a slide, covered with a cover slip and sealed with nail polish. Measurements were made only on cross-sections, but to assist in interpreting and identifying cell types, longitudinal tangential and radial sections and macerations were also made. For macerations, small shavings were placed in vials filled with Franklin’s solution: glacial acetic acid and 6 % hydrogen peroxide in proportions of 1 : 1 (Franklin 1945). These vials were loosely covered with Parafilm tape and heated in the oven at 60 °C for 1–2 days. Tissues were then rinsed with distilled water, stained with safranin O (10 min) and gently squashed onto a microscope slide.

Microphotographs of cross-sections were taken at ×100 and ×400 magnifications using a digital camera (Scion Corporation, CFW-1310C, USA) attached to a light microscope (Olympus BX 50F, Olympus Co. Ltd, Japan) and image capturing software Scion Visicapture, version 1.4 (Scion Corporation). Two to three images of the same area at different focal planes were taken and stacked in Photoshop CS4 (Adobe Systems Incorporated, USA). Cross-sections were bigger than the field of view; therefore, dozens (for ×100) or around 10 (for ×400) images per cross-section were taken and then merged in Photoshop, giving rise to images of whole cross-sections at ×100 and of one narrow transect at ×400. Tissue cross-sectional areas and vessel traits were measured on one wedge-shaped transect per replicate (× 100; Fig. 1) and fibre characteristics (× 400) on one rectangular transect, both stretching from pith to cambium. The radial transects were chosen to be the most representative for a section and the tension wood was avoided where possible. The transect borders were approximately parallel to the rays and followed middle lamella so that no open cells were positioned on the borders. Tissue areas, vessels and fibre walls were manually coloured in Photoshop using a Cintiq 21UX graphic tablet (Wacom Co., Ltd, Japan). Proto-xylem and newly produced xylem were excluded from analysis. Larger regions were inspected in species with larger vessels or with more variable structure in the tangential direction. The measured area ranged from 0.21 to 0.69 mm² for species from cool sites and from 0.31 to 1.9 mm² for species from hot sites. On average, for each image there were 170 vessels measured (+145) of all sizes including vessel tails. Fibre, fibre wall and lumen areas were measured for an average of 170 fibres per sample (+57) lying in two parallel rows from pith to cambium. Colour-coded images were analysed with Image-Pro Plus version 2.0.0.260 (Media Cybernetics, Inc., USA; Fig. 1).

Proportions of all major wood cell types (vessel wall and lumen, fibre wall and lumen, axial parenchyma, rays and tracheids) and their properties (mean vessel lumen area and mean fibre lumen and wall area) were quantified. Here, mean vessel lumen area is also called ‘mean vessel area’ for brevity. Mean vessel area, fibre lumen area and wall area were calculated as the arithmetic mean across all relevant cells (vessels or fibres) measured within a given radial sector of a sample. In addition to these traits we calculated theoretical maximum hydraulic

**Figure 1.** A twig cross-section of *Grevillea parallela*, Proteaceae. The radial sector of the stained section is shown on the left side and the processed image on the right. Colours in the processed image denote different tissue types: blue, vessel lumen; purple, vessel wall; green, rays; orange, axial parenchyma; brown, fibres. The scale bar corresponds to 100 μm.
conductivity (called here ‘theoretical conductivity’; also known as ‘potential conductivity’). Conductivity for the lumen of each conduit was calculated from the Hagen–Poiseuille expression \((\pi r^4)/(8\mu)\), where \(r\) represents mean vessel lumen radius and \(\mu\) represents water dynamic viscosity (assuming a standard viscosity value of 1 at 20 °C). The sum of all vessel lumen conductivities per unit cross-section is theoretical conductivity. This quantity expresses conductivity variation across species due to conduit diameters, but it should be thought of as the theoretical maximum since it does not capture effects of end-wall resistances, blockage by tyloses, effects of emboli, etc.

Traditionally, wood is considered a complex tissue composed of several cell types (Evert 2006). However, we refer to those cell types as ‘tissues’ for brevity and also because they perform distinctly different functions. Anatomical terminology follows the ‘IAWA list of microscopic features for hardwood identification’ (IAWA Committee 1989). Vascular and/or xylem tracheids occurred in 13 species and are referred to hereafter as tracheids. Tracheids were first determined in macerated wood and then identified on a cross-section on the basis of the number of pits and size of the pit border (both resembling that of vessel pits) and cell size (IAWA Committee 1989; Sano et al. 2011). Axial apotracheal and paratracheal parenchyma were collectively measured as axial parenchyma. None of the species studied here had storied rays. Tissue types were expressed as the fraction of a tissue per cross-sectional area (Fig. 1). Tissue fractions of the area outside vessel lumen (non-vessel area) were calculated as tissue fraction multiplied by non-vessel area fraction. The term ‘non-vessel’ area is used for brevity and it includes vessel walls, fibre walls and lumens, axial and ray parenchyma, and tracheids. Non-vessel-lumen quantities are hereafter denoted by subscript ‘NV’, e.g. fibre wall fractionNV, wood densityNV, etc. Fibre wall proportion in a given fibre was expressed as a proportion of the total fibre area. Fibre wall fraction was obtained by multiplying mean fibre wall proportion in a fibre by the fibre fraction per cross-section. Fibre lumen fraction was similarly calculated from mean fibre lumen proportion in a fibre multiplied by fibre fraction.

Statistical analysis
We collected measurements for 23 species, one of which occurred at two sites and was considered as two entities, giving a total of 24 data points analysed. Measurements were carried out on three replicate individuals per species and the trait values were averaged for comparisons across species. Wood density, vessel lumen, sum of ray and axial parenchyma, ray parenchyma and fibre wall fractions of total wood area as well as non-vessel wood area were all approximately normally distributed (Shapiro–Wilk test, \(P < 0.05\)). Vessel wall, axial parenchyma, tracheids and fibre lumen fractions of total wood area as well as of non-vessel wood area were right-skewed and were transformed to generate approximately normally distributed variables. Log10 transformations normalized all distributions with the exception of tracheid fraction and tracheid non-vessel fraction. We used ordinary least-squares regression to assess bivariate relationships (SigmaPlot, Systat, San Jose, CA, USA).

Results
Wood density and tissue proportions varied significantly across species as illustrated in Fig. 2 [see Supporting Information]. Wood density varied more than 2-fold, from 0.37 to 0.83 g cm\(^{-3}\). Figure 2 shows tissue fractions averaged across all species (bar at the top) and for each individual species separately (the remaining bars). The mean fibre fraction was 0.52 ± 0.09 (hereafter numbers represent average fraction ± one standard deviation). Fibre varied approximately 2-fold across species and was the most abundant tissue type. Fibre fraction could be partitioned into fibre walls (0.45 ± 0.08, 2-fold variation; brown bars in Fig. 2) and fibre lumens (0.08 ± 0.07, c. 60-fold variation; yellow bars). On average, parenchyma occupied 0.25 of wood cross-sectional area and varied almost 3-fold across species. It consisted of axial parenchyma (0.10 ± 0.05, 6-fold variation) and ray parenchyma (0.15 ± 0.05, 4.5-fold variation). Vessels occupied 0.20, where 0.15 ± 0.03 consisted of lumens (varying c. 2-fold) and 0.05 ± 0.03 of vessel walls (varying 4.5-fold). Tracheids occurred in just 13 of 24 species and occupied only small fractions (0.02 ± 0.03 averaged across all 24 species, 8.5-fold variation across the 13 species that had tracheids).

Alternatively, wood can simply be divided into two components: vessel lumen fraction and non-vessel fraction, which encompasses all tissues other than vessel lumens. The density of the non-vessel fraction and tissue fractions within the non-vessel fraction are indicated hereafter by the subscript ‘NV’, e.g. densityNV, fibre fractionNV. Since vessel lumen has zero density, overall wood density is (by definition) the product of the non-vessel fraction density (densityNV) and the non-vessel fraction itself (fractionNV): density = densityNV × fractionNV (Preston et al. 2006; Zanne et al. 2010). These three quantities can be log transformed and the equation then becomes a sum: \(\log(\text{density}) = \log(\text{density}_{\text{NV}}) + \log(\text{fraction}_{\text{NV}})\). Hence, when densityNV is plotted against fractionNV on log–log axes (Fig. 3), iso-lines of resulting overall wood density can be constructed and used to aid interpretation. Variation across species in densityNV (y-axis in Fig. 3) was four times greater than variation in fractionNV (x-axis), and thus in this species set was a far stronger determinant of variation in overall wood density.
Density (direction across the isolines). Not surprisingly then, density$_{NV}$ and overall density were tightly correlated with each other ($r^2 = 0.95, P < 0.001$). Vessel lumen fraction was only loosely (negatively) correlated with overall wood density ($r^2 = 0.20, P = 0.027$). Therefore, the following analyses concentrate entirely on density$_{NV}$ and its anatomical components [see Supporting Information].

Total fibre fraction$_{NV}$ (fibre lumen fraction$_{NV}$ plus fibre wall fraction$_{NV}$) was unrelated to density$_{NV}$ (Fig. 4). Species with the same fibre fraction$_{NV}$ varied widely in wall proportion relative to lumen within a fibre (as indicated by the width of the ‘donut’ rings in Fig. 4). Figure 4 shows that lowest-density (<0.5 g cm$^{-3}$) species had high fibre fraction$_{NV}$ but their fibres had low wall proportion (lower right of the graph). High-density$_{NV}$ species (>0.85 g cm$^{-3}$) also had high fibre fraction$_{NV}$ but their fibres had large wall proportion (upper right of the graph). A substantial number of species located in the middle of the graph with medium density$_{NV}$ (0.60–0.85 g cm$^{-3}$) had variable fibre fraction$_{NV}$ and fibre wall proportion within a fibre. Also, fibre wall proportion within a fibre was positively correlated with wood density$_{NV}$ ($r^2 = 0.62, P < 0.001$).

Density$_{NV}$ was positively correlated with fibre wall fraction$_{NV}$ ($r^2 = 0.40$; Fig. 5A), and negatively with fibre lumen fraction$_{NV}$ ($r^2 = 0.56$; Fig. 5B). Since the majority of species had low fibre lumen fraction$_{NV}$ (i.e. the data were right-skewed), the two species with highest fibre lumen fraction$_{NV}$ had a strong influence on these relationships (lower left of Fig. 5A, lower right of Fig. 5B). That said,
the correlations were still present across the other species considered on their own ($r^2 = 0.26$ and $P = 0.014$, $r^2 = 0.18$ and $P = 0.049$, respectively). The species with the lowest fibre wall fraction$_{NV}$ (upper left in Fig. 5A) was Daviesia latifolia, which had a high amount (fraction of 0.07) of thick-wall tracheids. Presumably, this high fraction of tracheid wall contributed to the quite high density$_{NV}$ of this species (high, given its very low fibre fraction).

Next we asked how fibre wall and lumen fractions$_{NV}$ were related to one another. Figure 6 shows an approximately triangular relationship between fibre wall fraction$_{NV}$ and fibre lumen fraction$_{NV}$. The highest-density$_{NV}$ species (>0.85 g cm$^{-3}$; four large symbols, upper left of Fig. 6) had high fibre wall fraction$_{NV}$ and low fibre lumen fraction$_{NV}$. Medium-density$_{NV}$ species (0.60–0.85 g cm$^{-3}$; 18 medium-sized symbols in Fig. 6) had variable fibre wall fraction$_{NV}$ and variable fibre lumen fraction$_{NV}$. The two lowest-density$_{NV}$ (<0.50 g cm$^{-3}$) species had the highest fibre lumen fraction$_{NV}$ and low fibre wall fraction$_{NV}$ (two smallest symbols, lower right of Fig. 6). The categories of high-, medium- and low-density species are used here for easy reference, but in fact, the trait values are continuous and no clear boundaries can be indicated. The species with lowest fibre wall fraction$_{NV}$ (lower left in Fig. 6) was D. latifolia (see the comment about tracheids above).

The second most abundant tissue, parenchyma, was not correlated with density$_{NV}$ nor were its components, rays and axial parenchyma (all $P > 0.7$). Similarly, vessel wall fraction$_{NV}$ was unrelated to density$_{NV}$. Nevertheless, both parenchyma and vessel wall fractions$_{NV}$ indirectly affected density$_{NV}$. Figure 6 illustrates that there was considerable variation in density$_{NV}$ (indicated by symbol size) at a given fibre wall fraction$_{NV}$, especially at lower wall fraction$_{NV}$ (lower half of the graph), and considerable variation in density$_{NV}$ at a given fibre lumen fraction$_{NV}$, especially at lower lumen fraction$_{NV}$ (left half of the graph). These variations in density$_{NV}$ could be partially explained by parenchyma and vessel wall fractions$_{NV}$. At a given fibre wall fraction$_{NV}$, density$_{NV}$ was positively correlated with parenchyma and vessel wall fractions$_{NV}$ ($r^2 = 0.15$, $P = 0.064$ and $r^2 = 0.29$, $P = 0.007$, respectively) and negatively with fibre lumen fraction$_{NV}$ ($r^2 = 0.45$, $P < 0.001$; all relationships tested against residuals from a regression of wood density$_{NV}$ on fibre wall fraction$_{NV}$). Conversely at a given fibre lumen fraction$_{NV}$ (i.e. tested against

Figure 6. Relationship between non-vessel density (wood density$_{NV}$) and fibre fraction in non-vessel fraction (fibre fraction$_{NV}$). Each ‘donut’ circle symbolizes one species. The width of the donut border (black) represents fibre wall proportion within an individual fibre and the width of the hole (white) represents fibre lumen proportion. These proportions were estimated from individual fibres (as fibre wall area—or lumen area—divided by total fibre area), for 75–314 fibres per replicate (mean 170), and then across three replicates per species.

Figure 5. Relationships between non-vessel density (wood density$_{NV}$) and (A) fibre wall fraction in non-vessel fraction (fibre wall fraction$_{NV}$) and (B) fibre lumen fraction in non-vessel fraction (fibre lumen fraction$_{NV}$). Each circle represents a different species (mean value from three replicates). ***$P < 0.001$. 

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residuals from a regression of wood density_{NV} on fibre lumen fraction_{NV}, wood density_{NV} was negatively correlated with parenchyma fraction_{NV} ($r^2 = 0.28$, $P = 0.008$), not correlated with vessel wall fraction_{NV} ($P = 0.29$) and positively correlated with fibre wall fraction_{NV} ($r^2 = 0.40$, $P < 0.001$). Additionally, parenchyma fraction_{NV} was tightly negatively correlated with total fibre fraction_{NV} ($r^2 = 0.74$, $P < 0.001$). The only remaining tissue, tracheids, occupied on average a very small fraction ofNV and was not subjected to detailed analysis.

Mean vessel area and theoretical conductivity [see Supporting Information] were negatively correlated with overall wood density across the species studied ($r^2 = 0.225$, $P = 0.019$ and $r^2 = 0.33$, $P = 0.0036$, respectively, Fig. 7A and B). However, these relationships were mainly driven by a few species with particularly large vessel lumens and theoretical conductivity (Fig. 7A and B). Among the majority of species there was considerable variation in mean vessel area and conductivity at a given wood density, and little relationship between the two (Fig. 7A and B).

**Discussion**

This study aimed to describe the anatomical components of wood density in twigs of 24 Australian tree and shrub species. Properties of fibres, the most abundant tissue, had the strongest effect on wood density variation, as has been shown in some previous studies (Fujiwara et al. 1991; Jacobsen et al. 2007; Martinez-Cabrera et al. 2009). However, our results contribute to a more comprehensive understanding of wood structure and its influence on density in twigs. Here, we discuss in detail the properties of fibres and other tissues as components of density.

**Wood density and its anatomical components**

Density of tissue outside vessel lumens (density_{NV}), rather than vessel lumen fraction, was the main driver of overall wood density variation. This result agrees with a comparison made across 584 species that considered main stem wood (Zanne et al. 2010), implying that density_{NV} determines overall density in both twigs (this study) and main stems similarly. Accordingly, the discussion here is directed towards density_{NV} and individual tissue fractions within the non-vessel part of the wood (indicated by subscript ‘NV’, e.g. fibre wall fraction_{NV}). We compare these results with the results for overall wood density reported by other studies, on the basis that overall wood density and density_{NV} are closely correlated ($r^2 = 0.95$, $P < 0.001$, this study).

All tissue fractions_{NV} influence density_{NV} but fibre wall and lumen fractions_{NV} are the most important. The
strong influence of fibre wall fraction_{NV} was due both to its high proportion and its variability, while the influence of fibre lumen fraction_{NV} was associated chiefly with its high variability (c. 60-fold). Other studies have consistently found a positive relationship between fibre wall fraction and density: in trunk wood among 50 Japanese trees (Fujiwara et al. 1991) and 61 North and South American shrubs (Martínez-Cabrera et al. 2009); and in twig wood of 17 South African shrubs (Jacsobsen et al. 2007a). Fibre lumen fraction has received less attention but has also been shown (in concordance with our study) to have a negative relationship with wood density (Martínez-Cabrera et al. 2009; Rana et al. 2009). The second most abundant tissue, parenchyma, did not correlate with density_{NV} in our study nor in tree and shrub trunks (Martínez-Cabrera et al. 2009; Poorter et al. 2010; Fichtler and Worbes 2012) but correlated negatively with twig wood density across 17 species (Jacsobsen et al. 2007a). These discrepancies might be caused by variable densities of parenchyma tissue itself (Taylor 1969; Fujiwara 1992; Guilley and Nepveu 2003) or by varied correlations between parenchyma and fibre wall and lumen fractions (see below). Plausibly, these discrepancies could also stem from different relationships between ray and axial parenchyma. We did not find any cross-correlation between those two components of parenchyma nor were they individually related to density. In contrast, Martínez-Cabrera et al. (2009) reported that ray and axial parenchyma were negatively correlated with each other and individually correlated with density (axial parenchyma positively correlated, rays—negatively). In that study these links were strongly associated with environmental variables (MAP, MAT and AI). These findings imply that different functional trade-offs can be related to ray and axial parenchyma individually and also that their link is affected by climate. Possibly these trade-offs may be more pronounced in trunk wood, as opposed to the twig wood studied here.

We did not find a direct relationship between parenchyma and density_{NV}. Nevertheless, our results imply that parenchyma together with fibre lumens can influence density_{NV} variation in a less direct way. At a given fibre wall fraction_{NV}, density_{NV} depended on the parenchyma fraction_{NV} relative to fibre lumen fraction_{NV}. Density_{NV} was marginally positively correlated with parenchyma fraction_{NV} and negatively with fibre lumen fraction_{NV}. Parenchyma has higher density than fibre lumen, which has zero density. Therefore, higher parenchyma fraction_{NV} relative to fibre lumen fraction_{NV} increases density_{NV}, and conversely higher fibre lumen fraction_{NV} decreases density_{NV}. We note that parenchyma fraction_{NV} was only weakly correlated with density_{NV} at a given fibre wall fraction_{NV}. This weak correlation possibly stems from variable parenchyma tissue densities (Taylor 1969; Fujiwara 1992; Guilley and Nepveu 2003).

Fibre fraction_{NV} (sum of fibre wall and lumen fractions_{NV}) was not associated with density_{NV} because of wide variation in wall proportion within a fibre (top right to low right in Fig. 4). Poorter et al. (2010) suggested a similar explanation, but as far as we are aware, this issue has not been quantitatively clarified until now.

Anatomical traits that are not expressed as fractions of wood volume have less direct effects on density. We found that density_{NV} was correlated with fibre wall proportion within a fibre. Previous studies have also shown that density can be related to fibre lumen diameter (Jacsobsen et al. 2007a; Martínez-Cabrera et al. 2009), fibre wall thickness (Fujiwara et al. 1991; Martínez-Cabrera et al. 2009) and fibre wall to lumen ratio (Martínez-Cabrera et al. 2009). We believe those traits would deliver a more insightful understanding of wood density when combined with information about fractions, e.g. the relationship among density_{NV}, fibre fraction_{NV} and fibre wall proportion in a fibre, as described above. Similarly for mean vessel area, which in this study was weakly negatively correlated with wood density. Mean vessel area per se should not affect wood density. Rather, vessel fraction (vessel area multiplied by vessel number per area) should be causally linked with density variation.

### Variability of anatomical structures

The discussion so far has focused on wood density variation. However, we also found considerable anatomical variation within a given range of density. Hereafter, we use the arbitrary categories of ‘medium’, ‘high’ and ‘low’ density and tissue fractions. However, we observed a continuum of trait values and the categories are used only for convenience. Species with medium density_{NV} (0.60–0.85 g cm$^{-3}$) showed broader structural variability than high- and low-density_{NV} species (>0.85 and <0.5 g cm$^{-3}$, respectively). The concept is illustrated in Fig. 8 and examples of cross-sections approximately corresponding to Fig. 8 are shown in Fig. 9. Species with high density_{NV} (large symbols in Fig. 6, top corner in Figs 8 and 9A) had the highest fraction_{NV} of fibre wall and small fibre lumen fraction_{NV}. Their total fibre fraction_{NV} was high and parenchyma fraction_{NV} was low. In contrast, medium-density_{NV} species had more variable fibre wall, fibre lumen and parenchyma fractions_{NV} (medium symbols in Fig. 6, middle in Figs 8 and 9B and C). Consequently, a spectrum of possible architectures may be outlined where the two ends of the spectrum are (i) low fibre, low fibre lumen and high parenchyma fraction_{NV} (middle left in Figs 8 and 9B) and (ii) high fibre, medium fibre lumen and low parenchyma fraction_{NV} (middle right in Figs 8 and 9C). The lowest-density species (<0.5 g cm$^{-3}$, small...
Ecological and physiological considerations

It has been shown that denser woods tend to operate at more negative minimum water potentials (Ackerly 2004; Bucci et al. 2004; Santiago et al. 2004; Jacobsen et al. 2007b, 2008; Gotsch et al. 2010) and have higher cavitation resistance (Hacke et al. 2001; Jacobsen et al. 2005; Pratt et al. 2007; Lens et al. 2011). However, we and previous literature have shown that wood density is most strongly related to fibre properties, not vessel properties. Thus, the relationship among wood density, minimum water potential and cavitation resistance does not appear to be directly influenced by wood density, but rather by a third, unexplained factor. Another element of hydraulic strategies is capacitance, which tends to be negatively correlated with wood density (Meinzer et al. 2003, 2008; Pratt et al. 2007; Scholz et al. 2007). Our results imply that capacitance water could be stored in fibre lumen or in parenchyma but the exact mechanism of this water storage and release is not known, nor is it clear what might be the advantage or disadvantage of fibre lumen capacitance compared with parenchyma capacitance. Density also plays a role in the mechanical behaviour of wood. It is a good predictor of mechanical strength and elasticity at a given wood diameter (Chave et al. 2009). However, it has also been shown that plants can compensate for mechanically weak wood by building thicker stems (Anten and Schieving 2010; Larjavaara and Muller-Landau 2010; Butler et al. 2011). Thick, low-density stems could be mechanically strong but their maintenance costs would be higher due to large surface area (Larjavaara and Muller-Landau 2010). Additionally, our results suggest that in similar density woods maintenance costs per sapwood cross-sectional area could vary depending on wood anatomical structure. Namely, woods with high fraction of living parenchyma would presumably incur higher maintenance costs than woods with high fraction of non-living fibres. What then would be the advantage of high parenchyma fraction? Hypothetically, high parenchyma fraction could ensure high nutrient storage capacity, which may be beneficial in some ecological strategies. This issue and other questions about causal links between wood anatomical structure and ecological strategies require further investigation and open interesting pathways for future research.

Twig and main trunk comparisons

Caution is needed when comparing results from twigs with results from main trunks. It is not proven that the relationships of wood density and anatomy are the same at the twig level as at the trunk level. Taylor and Wooten (1973) found that in five species vessel and fibre fractions shifted in the same direction with plant height, but this was not the case for ray fraction. Other studies examining association between vessel lumen fraction and wood density across a large number of species showed no relationship in the main trunks (Martinez-Cabrera et al. 2009; Poorter et al. 2010; Zanne et al. 2010) but a negative relationship in twigs (Preston et al. 2006; Jacobsen et al. 2007a; Ziemiańska et al. — Anatomical underpinnings of wood density
Mitchell et al. 2008; Gleason et al. 2012). Such disparities indicate that the relationship between wood density and tissue fractions may conceivably be different in main trunks than in twigs, yet it is unclear how generally this is so.

**Conclusions**

Wood density has been proposed as a key plant functional trait and is related to ecological strategies (Chave et al. 2009) but relatively little is known about the underpinnings of these relationships. This study clarifies the anatomical basis for wood density variation across species. It shows that wood density depends on anatomical structure, but also that a range of very different structures can result in very similar wood density, especially among species with medium density (here, 0.60–0.85 g cm\(^{-3}\)). This conclusion suggests that there may be a wider range of ecological strategies among such species. Taken together, these
findings imply that twig wood density should not be considered as an unambiguous indicator of plant ecological strategies. We hope this research will enhance the interpretation and design of ecological studies related to wood density.

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Contributions by the Authors
All authors contributed to the analysis of the results, writing and editing of the manuscript. D.W.B devised site selection. K.Z., D.W.B and S.M.G. carried out fieldwork. K.Z. performed anatomical work.

Conflicts of Interest Statement
None declared.

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Supporting Information
The following Supporting Information is available in the online version of this article –
File 1. Table. Details of the four sites sampled in this study.
File 2. Table. Species traits: wood density and tissue fractions.
File 3. Table. Species traits: wood density and tissue fractions of non-vessel fraction.
File 4. Table. Species traits: mean vessel area and theoretical conductivity.

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