

## Reviewer Report

**Title: A chromosomal-scale genome assembly of *Tectona grandis* reveals the importance of tandem gene duplication and enables discovery of genes in natural product biosynthetic pathways**

**Version: Original Submission**    **Date: 12/10/2018**

**Reviewer name: Nicolas Delhomme, Ph. D. rer. nat.**

### Reviewer Comments to Author:

In their manuscript, Dongyan Zhao et al., present the genome assembly of *Tectona grandis* realised using the most recent sequencing technologies, followed by the identification and validation of genes important to wood formation, a trait of interest in teak. The manuscript is well written and the analyses appear to have been robustly conducted, but their lack of details prevents me from being more convinced. This is my only significant comment to the manuscript as it stands, it is otherwise very well written and should be a good resource for the community. As a note, I do find that the title is too boldly written considering the content presented and that the readership would gain from a more fitting title (i.e. the pathways discussed in the manuscript are well known and there is not proof as of yet that their update knowledge in teak will lead to a more sustainable teak production).

#### Major comment

1) The description of the transcriptome analysis is completely missing in the main text, as are the details of which datasets were retrieved. Overall, I would wish for the supplementary document to contain the details of all analyses, including the software used, their versions and any non-default parameters - as was done for the WGD analysis. The supplementary information available from the FTP hints that more comprehensive analyses were done that is reported in the main text (e.g. classification of the gene models in different confidence bins, etc.). Such details should be made more readily visible as they would improve the manuscript's impact.

#### Minor comments

- 1) p. 3 l.38 The number of scaffolds of the released assembly could also be given. As a stand alone figure, an N50 value is not particularly informative.
- 2) p.4 l. 83 Detail the BUSCO categories (write them in full)
- 3) p.5. l.88 Briefly describe the annotation process. Further in the same paragraph detail which evidences were used for Augustus (maybe discuss why Maker-P was not used) and which datasets and parameters were used for PASA2. Also provide any custom scripts in a public repository that were used for the manual curation of the genes and gene models.
- 4) p5. l. 104 provide more details about what a SiZer analysis is. In general, extend the methods to contain the parameters used by the different tools, when non default (e.g. those for DupPipe line 100).
- 5) Detail how the phenylpropanoid pathway genes were identified. Similarly, detail how this was achieved for the TPSs, including tools, versions, non default parameters.
- 6) p. 12 l. 259 "asterisks" or "stars" rather than "dots"
- 7) Figure 2, how was the expression calculated and what metrics is represented? Same for figure 5 and 6
- 8) In Figure 2, use the gene name described in the text in addition to the gene IDs.

9) p. 6 second paragraph and Figure 3. Discussing the gene family expansion in the light of the WGD would be of interest.

10) What do the red and black bar represent in Figure 5? Add the information to the legend.

11) In Figure 6, the coordinates as well as the scaffold should be indicated in the schematic gene representation

12) Supplementary Table 1 should contain the BUSCO results for the released assembly (Illumina + nanopore)

13) Supplementary Table 3 and 4; add the expression unit in the column header or as a caption.

This is a signed review by Dr. Nicolas Delhomme, researcher at the Swedish University for Agricultural Sciences, and manager of the bioinformatics facility at the Umeå Plant Science Centre.

## **Methods**

Are the methods appropriate to the aims of the study, are they well described, and are necessary controls included? Choose an item.

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Are the conclusions adequately supported by the data shown? Choose an item.

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