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Comment on “Critical wind speed at which trees break”

Axel Albrecht,1 Eric Badel,2 Vivien Bonnesoeur,2,3 Yves Brunet,4 Thiéry Constant,3 Pauline Défossez,4 Emmanuel de Langre,5 Sylvain Dupont,6 Meriem Fournier,7 Barry Gardiner,5,6,8 Stephen J. Mitchell,7 John R. Moore,8 Bruno Moulia,2 Bruce C. Nicoll,6 Karl J. Niklas,9 Mart-Jan Schelhaas,10 Hans-Christof Spatz,11 and Frank W. Telewski12

1Forest Research Institute Baden-Wuerttemberg, Wonnhaldestrasse 4, 79100 Freiburg, Germany
2PIAF, INRA, Université Clermont Auvergne, 63000 Clermont-Ferrand, France
3LERFoB, INRA, AgroParisTech, F-54000 Nancy, France
4ISPA, INRA, Bordeaux Sciences Agro, F-33140 Villenave D’Ornon, France
5Ladhyx, École Polytechnique, 91128 Palaiseau, France
6Forest Research, Northern Research Station, Roxlin, EH25 9SY, Scotland, United Kingdom
7The University of British Columbia, 304-1-2424 Main Mall, Vancouver, British Columbia, Canada V6T 1Z4
8Scion, Private Bag 3020, Rotorua 3046, New Zealand
9Plant Biology Section, School of Integrative Plant Science, Cornell University, Ithaca, New York 14853, USA
10Alterra, Wageningen UR, 6700 AA Wageningen, Netherlands
11Institut für Biologie III, Universität Freiburg, Freiburg D-79104, Germany
12W. J. Beal Botanical Garden, Department of Plant Biology, Michigan State University, East Lansing, Michigan 48824, USA

(VIROT ET AL. [E. VIROT ET AL., PHYS. REV. E 93, 023001 (2016)] assert that the critical wind speed at which ≥50% of all trees in a population break is ≈42 m/s, regardless of tree characteristics. We show that empirical data do not support this assertion, and that the assumptions underlying the theory used by Virot et al. are inconsistent with the biomechanics of trees.

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Virot et al. [1] report results on (i) the breakage of brittle rods of dry wood with various geometry and (ii) extend these results to predict wind speeds for trunk breakage in forests. Our comments deal with part (ii). In this part of their work they developed minimal scaling arguments combining linear theory of rod brittle breakage under static bending (small perturbations and Griffith’s criterion) with the “elastic similarity” scaling of tree slenderness (Ref. [39] in [1]). This predicts that the critical wind speed for trunk breakage is ≈42 m/s, with low if any dependency on tree height and material properties. This prediction was compared to reprocessed data from storm Klaus (24th January, 2009). If correct, this work would have important implications about the key factors controlling the critical strength of trees against the wind, as well as for forest and urban tree management (as shown by numerous reports on [1] in the popular science and arboricultural press).

However, there are several physical and mechanical reasons why we think that the claim of Virot et al. [1] cannot be substantiated.

1. Given the meteorological variability of storms and the biological variability of trees, observations after a single storm event do not provide a sound basis for generalization and validation of stem breakage models. Moreover, an evaluation of the model using the data from storm Klaus (24th January, 2009) has serious problems because only 16% of all damage in this storm was due to trunk breakage (see Ref. [26] in [1]). Thus Virot et al. [1] evaluated their model against an inadequate data set. It would have been beneficial if the authors had investigated storm damage from forests with high levels of stem breakage, for example from the large body of data on forest wind damage accumulated by the plant biomechanics research community (e.g., [2,3]).

2. The experiments of Virot et al. [1] employed dry wooden rods and the authors use the properties of dry wood in their calculations. Unfortunately the material properties of dry wood, being stiff and brittle, differ significantly from those of green, living wood, which is more flexible and less brittle [4]. Wood in tree stems also has a large radial variation in mechanical properties, which is an essential adaptation of trees living in a windy environment [5]. Therefore, Eq. (4) in Virot et al. [1], which was derived for dry wood, cannot be applied to the green, living wood in tree trunks.

3. Virot et al. [1] treat the wind loads on trees as those on tree trunks modeled as point-loaded, cylindrical nontapered rods with uniform material properties. This model is not consistent with the biomechanics of trees [6]. In particular, Virot et al. [1] made the unrealistic assumption that tree trunks are nontapered, homogeneous in composition, and entirely pruned by high winds before stem breakage. Each of these assumptions is unfounded; e.g., total branch pruning is a very rare event (in the vast majority of cases a tree’s branched crown is the major source of wind drag) [7].

4. The conclusion reached in Virot et al. [1] depends on the assumption of the interspecific scaling relationship \( H \propto D^{2/3} \) (wherein \( H \) = height and \( D \) = trunk diameter), which emerges from the hypothesis of elastic self-similarity [6,8]. However, elastic self-similarity is based on buckling under self-weight, which is almost unheard of in the natural world except in the case of extremely slender trees growing under unusual circumstances [9]. In addition, comparisons with broad datasets have shown that this scaling holds true only for trees of exceptionally tall stature, many of which emerge above the canopy of the surrounding forest (i.e., tree champions). Specifically, this scaling relationship does not hold for even-aged
single species stands [6,10], which are typical of the maritime pine forest dataset presented by Virot et al. [1]. Indeed, the size, shape, and material properties of all plants including trees are responsive to wind-induced mechanical loads during their growth, a phenomenon called thigmomorphogenesis [11–13]. Thus, trees growing in sheltered conditions differ in height, trunk diameter, and wood mechanical properties from their counterparts growing in windy sites [14]; and tree form is also influenced by the amount of competition for growing space and by silvicultural practices [15]. Consequently, no single scaling relationship for tree height in relationship to trunk diameter holds true even for trees belonging to the same species.

(5) The model by Virot et al. [1] is static. However, previous comparisons between static models and experimental data have revealed that dynamic effects linked to wind turbulence (wind gusts) cannot be neglected [16]. This led Virot et al. [1] to apply an additional “gust factor” of 2 for the wind loading. However, wind-tunnel and field measurements (e.g., [16]) have obtained values many times greater than 2 (typically 6–10) and have shown that these values depend on the ratio of tree height to tree spacing.

These and other concerns (e.g., damage propagation during storms [17], growth prestresses [18], and mechanical fatigue [19]) are not compatible with the assertion of a constant critical wind speed for tree failure. However, we do believe that the most rapid advance in understanding the interaction between trees and the wind will come from close collaboration among scientists from diverse disciplines such as physics, meteorology, biomechanics, forestry, and plant biology and we hope our commentary on Virot et al. [1] will encourage debate and further work on this complex subject.