Significance testing testate amoeba water table reconstructions

Richard J Payne1,2, Kirill V Babeshko2, Simon van Bellen3, Jeffrey J Blackford4, Robert K Booth5, Dan J Charman6, Megan R Ellershaw7, Daniel Gilbert8, Paul D M Hughes9, Vincent EJ Jassey8,10 Łukasz Lamentowicz11, Mariusz Lamentowicz11, Elena A Malysheva2, Dmitri Mauquoy12, Yuri Mazei2,13, Edward A D Mitchell14,15, Graeme T Swindles16, Andrey N Tsyganov2, T Edward Turner16, Richard J Telford17

1 Environment, University of York, Heslington, York, YO105DD, United Kingdom.
2 Department of Zoology and Ecology, Penza State University, Krasnaya str. 40, 440026 Penza, Russia.
3 GEOTOP-Université du Québec à Montréal, 201 Avenue Président-Kennedy, Montréal, Québec, H2X 3Y7, Canada.
4 Department of Geography, Environment and Earth Sciences, University of Hull, Cottingham Road, Hull, HU6 7RX, United Kingdom.
5 Department of Earth and Environmental Sciences, Lehigh University, 1 West Packer Avenue, Bethlehem, PA 18015-3001, USA.
6 Geography, College of Life and Environmental Sciences, University of Exeter, Amory Building, Rennes Drive, Exeter, EX4 4RJ, United Kingdom.
7 Natural England, Northminster House, Peterborough, PE1 1UA, United Kingdom.
8 Laboratoire Chrono-Environnement, UMR 6249 Université de Franche-Comté/CNRS, UFR Sciences et Techniques, 16 route de Gray, 25030 Besançon, France.
9 Geography and Environment, University of Southampton, University Road, Southampton, SO17 1BJ, United Kingdom.
10 Swiss Federal Institute for Forest, Snow and Landscape Research (WSL), Site Lausanne, Station 2, 1015 Lausanne, Switzerland.
11 Department of Biogeography and Palaeoecology, Faculty of Geographical and Geological Science, Adam Mickiewicz University, Poznan, ul. Dziegielowa 27, 61-680 Poznan, Poland.
12 School of Geosciences, University of Aberdeen, Elphinstone Road, Aberdeen AB24 3UF, United Kingdom.
13 Department of Hydrobiology, Lomonosov Moscow State University, Leninskiye gory, 1, Moscow 119991, Russia.
14 Laboratory of Soil Biology, University of Neuchâtel, Rue Emile-Argand 11, CH-2000 Neuchâtel, Switzerland.
15 Jardin Botanique de Neuchâtel, Pertuis-du-Sault 58, CH- 2000 Neuchâtel, Switzerland.
16 School of Geography, University of Leeds, Leeds, LS2 9JT, United Kingdom.

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ABSTRACT

Transfer functions are valuable tools in palaeoecology, but their output may not always be meaningful. A recently-developed statistical test (‘randomTF’) offers the potential to distinguish among reconstructions which are more likely to be useful, and those less so. We applied this test to a large number of reconstructions of peatland water table depth based on testate amoebae.

Contrary to our expectations, a substantial majority (25 of 30) of these reconstructions gave non-significant results (P>0.05). The underlying reasons for this outcome are unclear. We found no significant correlation between randomTF P-value and transfer function performance, the properties of the training set and reconstruction, or measures of transfer function fit. These results give cause for concern but we believe it would be extremely premature to discount the results of non-significant reconstructions. We stress the need for more critical assessment of transfer function output, replication of results and ecologically-informed interpretation of palaeoecological data.

KEYWORDS: Testate amoeba; Protist; Palaeoecology; Palaeohydrology; Transfer function; randomTF

INTRODUCTION

Testate amoebae are widely-used proxies in palaeoecological studies; in particular for the reconstruction of water table depth in peatlands (Charman, 2001; Mitchell et al., 2008). Over the last 25 years palaeoecology has been revolutionised by the use of statistical models (transfer functions) to quantitatively reconstruct environmental variables. However, questions are increasingly being raised about the reliability and robustness of transfer function results (Belyea, 2007; Juggins, 2013).

A transfer function will always give an output but that output may not always be meaningful. The only way to establish whether the output of a transfer function is ‘true’ is by comparing the results to independent data, but such data are not always available and even in such cases correlations are complicated by temporal autocorrelation and the limitations of the chronology.

Although we cannot realistically assess whether all reconstructions are correct we can conceivably test whether they are potentially useful. Telford and Birks (2011) propose a pragmatic solution: that a reconstruction can be considered statistically significant if it explains more of the variance in the fossil data than those of transfer functions trained on randomly-generated data. Telford and Birks (2011) propose a method, ‘randomTF’, in which:

1. The transfer function is applied to the fossil data to derive a reconstruction (using any commonly-applied method).

2. The proportion of variance in the fossil data explained by the reconstruction is determined using constrained ordination.

3. Multiple new transfer functions are derived using the established modern species data but with the environmental data replaced by uniformly distributed random variables.

4. These transfer functions are applied in turn to the fossil data and the variance they explain tested. This is repeated a large number of times, typically 999.

5. A reconstruction is considered statistically significant when the proportion of variance explained is greater than that of 95% of the transfer functions based on randomly-generated data.
We would expect reliable reconstructions to explain more variance in the fossil data than transfer functions trained on random data, and therefore to give significant results. However, a significant randomTF value is not proof of accuracy and a non-significant result does not necessarily imply inaccuracy. Non-significant results do however give cause for concern and suggest that transfer function output should be treated with caution. randomTF tests can potentially tell us which reconstructions we should trust more, which less, and whether we can predict more than one environmental variable from the same fossil dataset. Telford and Birks (2011) also propose an alternative test (‘obs.cor’) based on the correlation of optima values with axis species scores from a constrained ordination of the fossil data. This test is not applicable to all transfer functions methods and is not considered here. The randomTF test has been applied in a few studies (Amesbury et al., 2013; Lamarre et al., 2013; Swindles et al., 2015a) but is not yet routinely used in testate amoeba palaeoecology. Here we apply this test to a large number of published and unpublished records with the aim to identify the characteristics which are likely to lead to better reconstructions, giving better randomTF results.

METHODS

We identified 30 published and unpublished testate amoeba palaeoecological records (Table 1). These records span a large range of regions, mire types, analysts, time periods, and sampling resolutions, and form a large and reasonably representative sample of testate amoeba palaeoecological research. Reconstructions of water table depth were produced using either the transfer function used in the original study, the most geographically-appropriate model where a transfer function was not previously applied, or in a few cases transfer functions which have been produced since the data were originally published. Taxonomy was harmonised between the fossil data and training set, which in many instances required the grouping or deletion of some taxa (performance statistics may therefore differ slightly from those previously published). Transfer functions were applied based on the model selected by the original authors with sample specific errors calculated by bootstrapping (1000 cycles). All transfer functions were based on either weighted averaging, weighted averaging with tolerance downweighting or weighted average-partial least squares (Birks, 1995). We applied randomTF using 999 permutations with redundancy analysis as the ordination method. Analyses were conducted in R3.1.2 (R Development Core Team, 2014) using the packages analogue (Simpson, 2007), rioja (Juggins, 2009) and palaeoSig (Telford, 2011).

RESULTS and DISCUSSION

Only five of the 30 tests yielded a significant P-value (P<0.05; Table 1). While we expected that some reconstructions would give non-significant results this proportion is much higher than we anticipated. While a few reconstructions fail to reach P=0.05 by a relatively narrow margin (Tørvesø 1, Staroselsky Moch, Dot Lake B), many more have P-values which substantially exceed this value. Another two records published in the literature have given significant P-values: those of Swindles et al. (2015a) for Stordalen, Sweden and Lamarre et al. (2013) for Lac Le Caron, Canada. Amesbury et al. (2013) found a significant result for the Nordans Pond site of Hughes et al. (2006) using an extended transfer function, whereas here we find a non-significant result using the transfer function used in the original study (Charman and Warner, 1997). In the latter three cases multiple model structures were tested with some producing significant reconstructions, and some not. We note that in these instances a correction for multiple comparisons (such as a Bonferroni correction) would
probably have meant that the reconstructions did not reach significance. However, even if these  
results are included, eight significant P-values out of 32 reconstructions remains a surprisingly low  
proportion.

Telford and Birks (2011) identify four factors which might make the randomTF test prone to type II  
error (“false negative”): low numbers of effective species; small numbers of fossil samples; limited  
variability in the reconstruction and poorly-performing or poorly-fitting transfer functions. All of  
these factors apply to some of the reconstructions we examine but it is not clear that any are a  
consistent cause of non-significant P-values. Overall, P-value was not significantly correlated with  
properties of the training set (mean/standard deviation/range of WTD) or fossil data (species  
richness, Hill’s N2 or number of samples), performance metrics of the transfer function (leave one  
out RMSEP or $R^2$), properties of the reconstruction (mean/standard deviation/range of predications,  
ratio of range to RMSEP or training set range, mean boot-strapped error estimates) or measures of  
transfer function fit (proportion of shared taxa, proportion of fossil samples with poor modern  
analogues, squared residual length) (Spearman Rs; $P>0.05$). P-value was strongly correlated with the  
proportion of variance in the fossil data explained by the reconstruction (Spearman Rs=-0.89,  
$P<0.001$), suggesting (unsurprisingly) that where a high proportion of variance is explained this is  
unlikely to be exceeded by transfer functions trained on random data.

The five reconstructions yielding significant results were three short records from the Elatia Forest of  
northern Greece (Dexameni; Krya Vrissi 1&2; Payne and Pates (2009)), the high-resolution  
Mauntschas record from the Swiss Alps (Lamentowicz et al., 2010; van der Knaap et al., 2011) and a  
record from Frasne in the Jura Mountains of eastern France (Jassey et al. unpublished). These five  
records have little obvious similarity. The transfer functions used for the Dexameni, Krya Vrissi and  
Mauntschas reconstructions all included samples from the same sites and for Frasne the closest  
training set site was only c.10 km distant. However, ten of the sites with non-significant  
reconstructions were also included in their respective training sets. The three short records from  
Greece (Payne and Pates, 2009) are all characterised by a single large change—a shift to drier  
conditions in the recent past but this is not a feature of the Frasne or Mauntschas records and some  
non-significant reconstructions are similar (e.g. Andorra: van Bellen et al., in press).

Non-significant reconstructions include studies where it is difficult to see any a priori reason to  
suspect problems: sites with high resolution, good numbers of samples and species, transfer  
functions which perform well in cross-validation, include samples from the same site with good  
overlap in assemblage and with modern and fossil samples counted by the same analyst (e.g. Dead  
Island: (Swindles et al., 2010), Minden: Booth and Jackson (2003)). Non-significant results in these  
instances are a real surprise.

Our results provide some evidence that where there is a choice of transfer function this can affect  
randomTF significance level. With the Nordans Pond record (Hughes et al., 2006) the transfer  
function used in the original publication yields a non-significant P-value (P=0.83; Table 1) while a  
more recent transfer function with a larger training set and better performance statistics gives a  
significant P-value (Amesbury et al., 2013). With the Frasne record (Jassey and Gilbert unpublished)  
marginally better results are found with the smaller and weaker-performing, but geographically  
closer, Jura transfer function (Mitchell et al., 1999) than the larger, better-performing, north-west  
Europe model (Charman and Blundell, 2007) although both are P<0.05. It is also probable that in
So what should we take from these results?

Interpreting these findings is a challenge and among the authors of this paper there is a considerable range of viewpoints. In the original paper Telford and Birks (2011) state that ‘reconstructions that fail this test have limited credibility and should be treated with considerable caution’. On this basis these results could be taken to question the reliability of a substantial proportion of published testate amoeba water table reconstructions and thus raise questions about the approach as a whole.

However there are also strong arguments for a more cautious interpretation. Unlike many proxies the ecological underpinnings of testate amoeba palaeoecology are strong. The thickness of water films, for which water table depth is a surrogate, determine an amoeba’s ability to move and feed, (although seasonal variability is an important area of uncertainty: (Marcisz et al., 2014b)). Numerous modern studies have found significant correlations between amoeba communities and water table depth (Mitchell et al., 2008) and studies have begun to support this link experimentally (Marcisz et al., 2014a; Mulot et al., 2014). While there undoubtedly are both practical and fundamental issues which can complicate palaeoecological reconstruction, many of these are common to other proxies and archives. Most testate amoeba analysts would expect our reconstructions to satisfy the fundamental requirements for quantitative palaeoecology laid out by Birks (1995).

A substantial proportion of all applications of the randomTF test, with a variety of proxies in a variety of settings, have produced non-significant results (Cwynar et al., 2012; Luoto et al., 2014; Salonen et al., 2013; Wooller et al., 2012). If non-significant results are so common it is arguable that the test may be overly pessimistic. In the case of the records we consider here it can be argued that there are reasons to accept many of the non-significant reconstructions based on correlations between proxies and with independent data sources. There are also an increasing number of studies which show transfer functions to have acceptable performance when tested with independent data (Payne et al., 2012; Swindles et al., 2015b). The increasing number of pitfalls and caveats which have been identified in transfer functions over recent decades should be a warning to palaeoecologists of the dangers of uncritical acceptance of new statistical methods.

It is important to reiterate that even from the most sceptical viewpoint a non-significant randomTF P-value does not prove that a reconstruction is invalid. It would be very premature to discount the results of reconstructions identified as non-significant here; even non-significant reconstructions may still be useful. However the unexpected finding that many reconstructions fail this test clearly shows the requirement for a more detailed and critical assessment of reconstructions and a better understanding of the factors which cause non-significant results. Transfer function reconstructions should always be accompanied by thorough ecological interpretation of the record. For instance where a dry shift is reconstructed by the transfer function on the basis of a switch in dominance from *Archerella flavum* to *Trigonopyxis arcula* this can probably be considered robust given the well-understood hydrological preferences of these two taxa. However a similar reconstructed change should be treated with much greater caution if it is based on a change in dominance from (for instance) *Heleopera petricola* to *Cryptodifflugia sacculus*; taxa with much less well-understood ecological preferences.
Replication of results among cores, sites, proxies and archives, and informed ecological interpretation of the primary data remain the best ways for palaeoecologists to ensure that our results are valid and useful.

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Author contributions: The study was conceived by RJP and RJT. RJP conducted statistical analyses and wrote the first draft of the manuscript. Other authors contributed data, advised on taxonomy, and provided interpretation.

References


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<th>Mean bootstrap SE (cm)</th>
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1 Note that using a larger alternative training set and a WMAT model (but not an ML or WA-tol(Inv) model) Amesbury et al. (2013) did find a significant result for this record, suggesting that choice of training set can have a strong influence on significance level.

2 A significant result but with slightly lower significance level (P=0.04) is produced if using the Northwest Europe transfer function (Charman and Blundell, 2007).

3 RandomTF results previously presented by van Bellen et al. (submitted), recalculated here.