

# Phylogenomic analyses of lophophorates (brachiopods, phoronids and bryozoans) confirm the Lophotrochozoa concept

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Based on embryological and morphological evidence, Lophophorata was long considered to be the sister or paraphyletic stem group of Deuterostomia. By contrast, molecular data have consistently indicated that the three lophophorate lineages, Ectoprocta, Brachiopoda and Phoronida, are more closely related to trochozoans (annelids, molluscs and related groups) than to deuterostomes. For this reason, the lophophorate groups and Trochozoa were united to Lophotrochozoa. However, the relationships of the lophophorate lineages within Lophotrochozoa are still largely unresolved. Maximum-likelihood and Bayesian analyses were performed based on a dataset comprising 11 445 amino acid positions derived from 79 ribosomal proteins of 39 metazoan taxa including new sequences obtained from a brachiopod and a phoronid. These analyses show that the three lophophorate lineages are affiliated with trochozoan rather than deuterostome phyla. All hypotheses claiming that they are more closely related to Deuterostomia than to Protostomia can be rejected by topology testing. Monophyly of lophophorates was not recovered but that of Bryozoa including Ectoprocta and Entoprocta and monophyly of Brachiozoa including Brachiopoda and Phoronida were strongly supported. Alternative hypotheses that are refuted include (i) Brachiozoa as the sister group of Mollusca, (ii) ectoprocts as sister to all other Lophotrochozoa including Platyzoa, and (iii) ectoprocts as sister or to all other protostomes except chaetognaths.

**Keywords:** Brachiopoda; Bryozoa; Lophophorata; Metazoa; Phoronida; phylogeny

## 1. INTRODUCTION

Phylogenetic analyses of molecular markers have substantially changed our view of animal evolution in the past two decades (Halanych 2004). The new subdivision of Protostomia into two main groups, Lophotrochozoa and Ecdysozoa, originally based on 18S rDNA sequences (Halanych *et al.* 1995; Aguinaldo *et al.* 1997), has been corroborated by sequences of single nuclear protein-encoding genes (e.g. Ruiz-Trillo *et al.* 2002; Anderson *et al.* 2004), datasets combining multiple nuclear protein-encoding sequences (Peterson *et al.* 2004; Helmkamp *et al.* 2008) and phylogenomic approaches (Philippe *et al.* 2005; Philippe & Telford 2006; Baurain *et al.* 2007; Hausdorf *et al.* 2007).

However, the relationships within Lophotrochozoa could not be resolved robustly so far, neither with a large dataset of combined small and large subunit rDNAs (Passamanek & Halanych 2006), nor with a dataset including several nuclear protein-encoding sequences (Helmkamp *et al.* 2008). Phylogenomic data were able to resolve some disputed relationships within Lophotrochozoa (Hausdorf *et al.* 2007), but such data are still missing for some phylogenetically important phyla such as Brachiopoda and Phoronida.

The placement of the lophophorate taxa within Lophotrochozoa as indicated by molecular phylogenetic

studies is particularly inconsistent with the morphological evidence (Lüter & Bartolomaeus 1997). As originally defined based on morphology, Lophophorata consists of Ectoprocta, Brachiopoda and Phoronida, taxa that share a ciliated, tentacular feeding apparatus around the mouth opening called lophophore. Based on embryological and morphological characters, Lophophorata was traditionally considered the sister or paraphyletic stem group of Deuterostomia (Hennig 1979; Schram 1991; Ax 1995; Lüter & Bartolomaeus 1997; Sørensen *et al.* 2000; Brusca & Brusca 2003). However, Nielsen (2001) argued that the lophophore of Bryozoa is not homologous to that of Brachiopoda plus Phoronida, and considered Lophophorata diphyletic. He suggested that ectoprocts are more closely related to entoprocts within Spiralia, whereas he still considered Brachiopoda + Phoronida as the sister group of Deuterostomia *sensu stricto* (his Neorealia). By contrast, the molecular phylogenetic studies have shown Ectoprocta as well as Brachiopoda and Phoronida to be more closely related to Trochozoa, i.e. Annelida, Mollusca and related groups, than to Deuterostomia; these include analyses that used rDNA (Halanych *et al.* 1995; Mackey *et al.* 1996; Littlewood *et al.* 1998; Cohen 2000; Giribet *et al.* 2000; Peterson & Eernisse 2001; Mallatt & Winchell 2002; Passamanek & Halanych 2006), *Hox* genes (de Rosa *et al.* 1999; Passamanek & Halanych 2004), mitochondrial protein genes (Stechmann & Schlegel 1999; Helfenbein & Boore 2004; Waeschenbach *et al.* 2006), single nuclear protein genes (e.g. Ruiz-Trillo *et al.* 2002; Anderson *et al.* 2004) and sets of multiple nuclear

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protein genes (Helmkamp *et al.* 2008). For this reason, Halanych *et al.* (1995) united the lophophorate groups and Trochozoa into Lophotrochozoa.

Yet the morphological similarities between Brachiozoa (Brachiopoda + Phoronida = Phoronozoa) and Deuterostomia seem so strong that they affect the topology of the trees even in some analyses considering both 18S rDNA sequences and morphological characters. In the total-evidence analysis of Zrzavý *et al.* (1998), Brachiozoa clustered with Deuterostomia, while in the analysis of Eernisse & Peterson (2004) deuterostomes were the sister group of Lophotrochozoa. There, the brachiozoans were sister to the remaining lophotrochozoan groups. However, this was not the case in some other total-evidence analyses (Giribet *et al.* 2000; Peterson & Eernisse 2001) in which Brachiopoda and Phoronida were part of Lophotrochozoa (or Trochozoa), and Deuterostomia did not appear as the sister group of Lophotrochozoa. As a caveat to these findings, the above-mentioned studies did not include many genes at all.

To provide a more robust resolution of the relationships of Brachiopoda, Phoronida and Bryozoa, we supplemented a previously compiled dataset of 79 sequences encoding ribosomal proteins with new expressed sequence tag (EST) sequences of a brachiopod and a phoronid.

## 2. MATERIAL AND METHODS

### (a) EST generation and processing

Specimens of the brachiopod *Novocrania anomala* (Müller 1776) and the phoronid *Phoronis muelleri* Selys-Longchamps 1903 were collected in the Gullmarsfjord near Kristineberg, Sweden. To minimize potential contamination sources, care was taken to remove epibionts growing on the shells and tubes, respectively. Total RNA was isolated from pools of 20 living adult individuals each with the TRIzol Plus purification system (Invitrogen, Karlsruhe, Germany). The mRNA of *Novocrania* was purified by the Dynabeads mRNA Purification Kit (Invitrogen) before it was transcribed by primer extension. The products were size fractionated and cloned directionally using CloneMiner technology (Invitrogen) to construct a cDNA library. In *Phoronis*, the PolyATtract mRNA Isolation System III (Promega, Mannheim, Germany) was used, followed by transcription and long-distance PCR amplification, size fractionation and directional cloning employing the Creator SMART cDNA Library Construction Kit (Clontech, Heidelberg, Germany).

From these libraries, ESTs were generated by sequencing 2247 (*Novocrania*) and 2315 (*Phoronis*) clones from the 5' end on the automated capillary sequencer system ABI 3730XL (Applied Biosystems, Darmstadt, Germany) using BIGDYE chemistry (Applied Biosystems). EST processing was accomplished as described previously (Hausdorf *et al.* 2007), with the addition of a second clustering step after quality clipping to improve contig assembly. The final number of contigs acquired from each organism amounted to 1699 (*Novocrania*) and 1467 (*Phoronis*).

### (b) Extraction and alignment of ribosomal protein sequences

Ribosomal protein sequences were retrieved from the new datasets using 79 human ribosomal protein sequences as local BLAST search queries. A total of 42 and 54 (at least partial) ribosomal protein sequences were identified in *Novocrania*

and *Phoronis*, respectively. These sequences, available in GenBank under the accession nos. EU558289–EU558330 (*Novocrania*) and EU558331–EU558384 (*Phoronis*), were individually aligned to orthologous riboprotein sequences of 36 additional taxa compiled previously (Hausdorf *et al.* 2007) and of a nemertean (Struck & Fisse 2008) using the CLUSTALW algorithm (Thompson *et al.* 1994). The resulting single-gene alignments were inspected and adjusted manually, and concatenated into a single multiple sequence alignment. Ambiguously aligned positions were automatically removed by GBLOCKS (Castresana 2000) applying low stringency parameters. The resulting alignment included 5458 amino acids of *Novocrania* (47.8% of the total alignment length minus gap positions) and 7922 amino acids of *Phoronis* (69.3%). More extensive information about the number of genes and amino acids present per taxon is reported in the electronic supplementary material. The final alignment has been deposited at TREEBASE (<http://www.treebase.org>, study accession no. S2050).

### (c) Phylogenetic analyses

Maximum-likelihood analyses were conducted with TREEFINDER (Jobb *et al.* 2004; Jobb 2007). The rtRev+G+F model of protein evolution was used for the maximum-likelihood analyses because its fit to the present dataset was superior to other models according to the Akaike information criterion with a correction term for small sample size. Confidence values for the edges of the maximum-likelihood tree were computed by bootstrapping (100 replications; Felsenstein 1985).

To test predefined phylogenetic hypotheses, we used constrained trees and the 'resolve multifurcations' option of TREEFINDER to obtain the maximum-likelihood tree for a specified hypothesis. Then we investigated whether the maximum-likelihood trees for these hypotheses are part of the confidence set of trees applying the approximately unbiased test (Shimodaira 2002) and the expected likelihood weights method (Strimmer & Rambaut 2002).

Bayesian analyses were performed using PHYLOBAYES v. 2.3 (Blanquart & Lartillot 2006) based on the site-heterogeneous CAT model (Lartillot & Philippe 2004). Four independent Markov chains, starting from random points of the parameter space, were run simultaneously for 20 000 cycles each. Chain stationarity was evaluated by monitoring key parameters for long-term trends (e.g. log likelihood, alpha parameter). The first 2000 points were consequently discarded as burn-in. Both runs reached convergence, indicated by the maximal and mean difference of split frequencies amounting to 0.21 and 0.007, respectively. Subsampling every tenth tree from each chain, a 50% majority rule consensus tree was finally computed. We accept Bayesian posterior probabilities larger than 95% and bootstrap values larger than 70% as significant.

## 3. RESULTS AND DISCUSSION

### (a) Deuterostome versus lophotrochozoan relationships of lophophorates

The results of our maximum-likelihood (figure 1a), as well as Bayesian analyses (figure 1b), based on concatenated sequences of 79 ribosomal proteins encompassing 11 445 amino acid positions from 39 taxa, demonstrate that the three lophophorate lineages, Ectoprocta, Brachiopoda and Phoronida, are more closely related to trochozoan

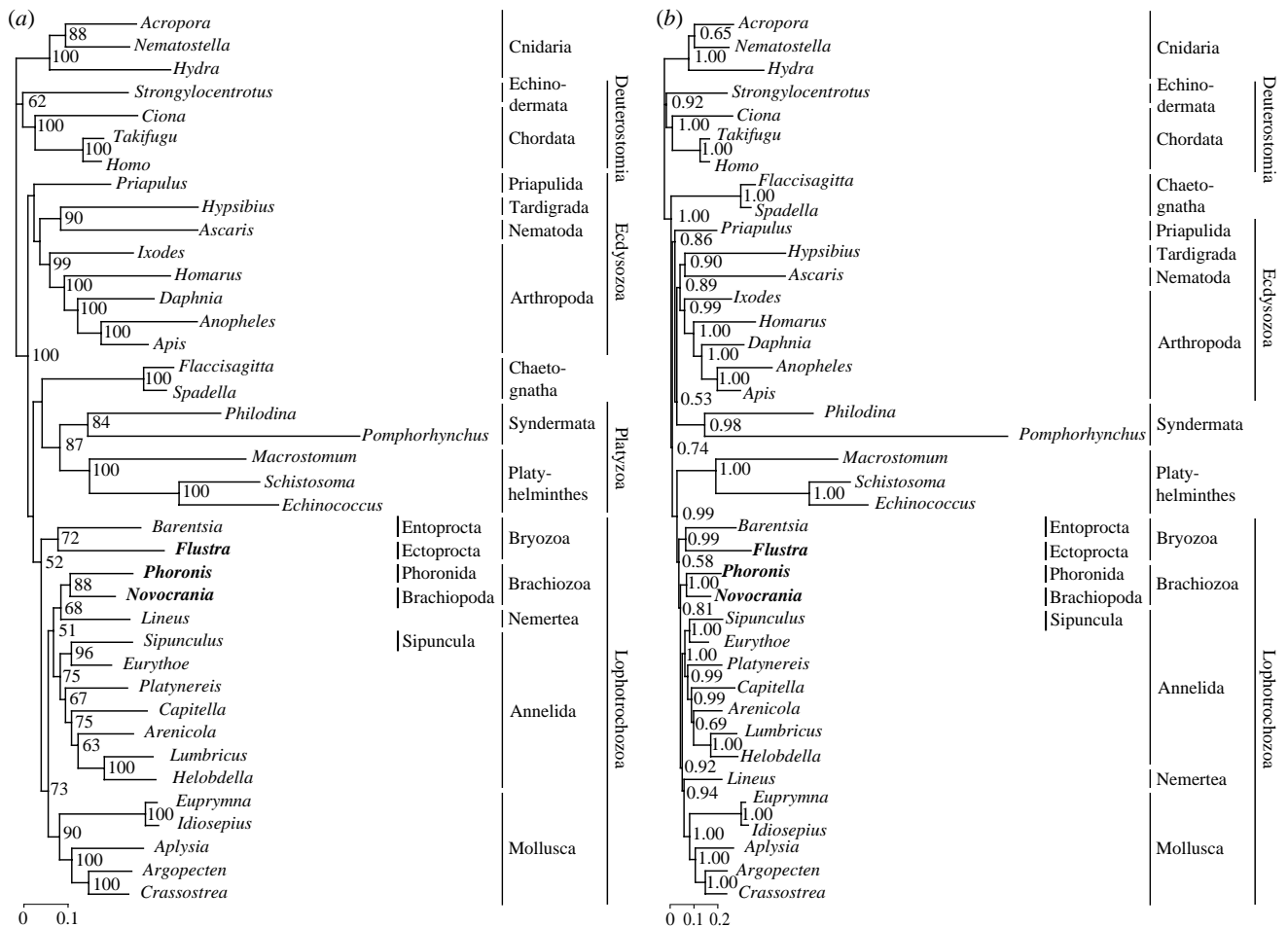


Figure 1. Phylogenetic analyses of lophophorate relationships based on 11 445 amino acid positions derived from 79 concatenated ribosomal proteins. Lophophorate lineages appear in bold. (a) Maximum-likelihood tree. Bootstrap support values larger than 50% are shown to the right of the nodes. (b) Bayesian inference reconstruction. Bayesian posterior probabilities are shown to the right of the nodes.

phyla than to deuterostomes. They do not form a monophyletic group. Thus, our analyses confirm the results of previous molecular phylogenetic studies (see §1).

Characters that were traditionally used to unite Lophophorata with deuterostomes include the following: a body organization with three distinct coelomic cavities, namely protocoel, mesocoel and metacoel (archimery); a mesosomal tentacular apparatus; entomesoderm derived from the archenteron by enterocoely, larvae with upstream-collecting ciliary bands; and heterogeneously assembled metanephridia (Hennig 1979; Schram 1991; Ax 1995; Lüter & Bartolomaeus 1997; Sørensen *et al.* 2000; Brusca & Brusca 2003). However, the hypothesis that all lophophorate lineages are more closely allied to Deuterostomia than Protostomia can be rejected by topology tests based on our ribosomal protein data (table 1, hypothesis 1). Nielsen (2001) argued that ectoprocts show no trace of archimery and that only Brachiopoda and Phoronida form a monophyletic group with Deuterostomia *sensu stricto* (his Neorennalia). Lüter (2000) suggested that the origin of the coelomic anlage from differentiated archenteral epithelium, which he defined as enterocoely, is a synapomorphy of Brachiopoda and Deuterostomia; he therefore considered these two taxa as sister groups. Consequently, we tested the hypotheses that Brachiozoa or Brachiopoda alone are the sister

groups of Deuterostomia. Both possibilities were rejected (table 1, hypotheses 2–3).

The conflicting results concerning the phylogenetic relationships of the lophophorates is a major incongruity between morphological and molecular phylogenetic approaches. However, in the last decade, the morphological evidence for a close relationship between the lophophorate groups and the deuterostomes has become weaker by careful re-examinations of the characters. It has been shown that neither brachiopods nor phoronids possess three coelomic cavities, because a protocoel is lacking in all lophophorate groups (Lüter 2000; Bartolomaeus 2001). Thus, the archicoelomate concept (Siewing 1980) uniting Lophophorata and Deuterostomia, founded on the similarities of three distinct coelomic cavities, lost its basis. Additionally, the finding that Pterobranchia may nest within the enteropneusts (Cameron *et al.* 2000; Peterson & Eernisse 2001; Winchell *et al.* 2002) suggests that the ancestral deuterostome more closely resembled a mobile worm-like enteropneust than a sessile colonial pterobranch. This means that the similar tentacular feeding apparatuses of lophophorates and pterobranchs are not a synapomorphy of lophophorates and deuterostomes as supposed previously (Hennig 1979; Schram 1991; Ax 1995; Lüter & Bartolomaeus 1997), but evolved independently as convergent adaptations to the sessile

Table 1. Topology test results.

no. <sup>a</sup>	phylogenetic hypothesis	references claiming the hypothesis	likelihood	$\Delta$ likelihood <sup>b</sup>	AU <sup>c</sup>	ELW <sup>d</sup>
1	ML tree (figure 1a) Lophophorata + Deuterostomia	Hennig (1979), Schram (1991), Ax (1995), Lüter & Bartolomaeus (1997), Sørensen <i>et al.</i> (2000) and Brusca & Brusca (2003)	-273 512 -273 957	0 445	0.6421 0.0000*	0.5010 0.0000*
2	Brachiozoa + Deuterostomia	Nielsen (2001)	-273 977	465	0.0000*	0.0000*
3	Brachiopoda + Deuterostomia	Lüter (2000)	-273 805	293	0.0000*	0.0000*
4	Ectoprocta sister to all other Lophotrochozoa inclusive Platyzoa	Halanych <i>et al.</i> (1995), Halanych (2004) and Passamanek & Halanych (2006)	-273 599	87	0.0000*	0.0052*
5	Ectoprocta sister to all other protostomes except chaetognaths	Giribet <i>et al.</i> (2000)	-273 624	112	0.0000*	0.0002*
6	Trochozoa (Entoprocta + Eutrochozoa)	Zrzavý <i>et al.</i> (1998), Giribet <i>et al.</i> (2000) and Peterson & Eernisse (2001)	-273 592	80	0.0634	0.0107*
7	Eutrochozoa (Neotrochozoa + Nemertea)	Zrzavý <i>et al.</i> (1998), Giribet <i>et al.</i> (2000) and Peterson & Eernisse (2001)	-273 535	23	0.4114	0.1817
8	Neotrochozoa (Annelida + Mollusca)	Zrzavý <i>et al.</i> (1998), Giribet <i>et al.</i> (2000) and Peterson & Eernisse (2001)	-273 520	8	0.5929	0.2777
9	Conchozoa (Brachiozoa + Mollusca)	Cavalier-Smith (1998) and Halanych (2004)	-273 571	59	0.0000*	0.0015*
10	Lophophorata monophyly	Emig (1984)	-273 571	59	0.1111	0.0219*

<sup>a</sup> Numbers refer to the order of appearance in the text.

<sup>b</sup>  $\Delta$ likelihood, differences between the likelihood of a constrained tree and the maximum-likelihood tree.

<sup>c</sup> AU, approximately unbiased test (*p*-values). Values for topologies significantly rejected at the 0.05 level are indicated by an asterisk.

<sup>d</sup> ELW, expected likelihood weights. Values for topologies not included in the 0.95 confidence set are indicated by an asterisk.

lifestyle (Halanych 1996). Moreover, Lüter (2000) argued that the mesoderm does not originate by enterocoely in Ectoprocta and Phoronida, but that this is the case in Brachiopoda and Deuterostomia only. What is more, whether the mesoderm of brachiopods originates by enterocoely is also in dispute. Jenner (2004) tentatively concluded that reports of true enterocoely, i.e. mesoderm origin by epithelial folding, in brachiopods appear unsupported and that no fundamental difference in the source of mesoderm and mode of coelomogenesis exists between brachiopods and various protostomes. To conclude, there are fewer morphological characters arguing against protostome affiliations of brachiopods and phoronids than traditionally assumed.

### (b) Relationships of lophophorates within Lophotrochozoa

The phylogenetic analyses of our ribosomal protein dataset (figure 1) strongly indicate that Brachiopoda and Phoronida constitute a monophyletic group, Brachiozoa (=Phoronozoa; bootstrap support 88%, Bayesian posterior probability 1.00). This corroborates previous results based on rDNA (Mackey *et al.* 1996; Cohen *et al.* 1998; Littlewood *et al.* 1998; Cohen 2000; Mallatt & Winchell 2002; Halanych 2004; Cohen & Weydmann 2005; but see Passamanek & Halanych 2006), sodium-potassium ATPase  $\alpha$ -subunit (Anderson *et al.* 2004), morphology (Nielsen 2001) and a combination of morphological and 18S rDNA datasets (Zrzavý *et al.* 1998; Giribet *et al.* 2000; Peterson & Eernisse 2001).

Our previous study (Hausdorf *et al.* 2007) recovered Ectoprocta as the sister group of Entoprocta, but this finding was considered tentative until a phoronid and a brachiopod could be added to the analysis, which was done here (figure 1). Indeed Ectoprocta and Entoprocta remain strongly united (bootstrap support 72%, Bayesian posterior probability 0.99). This agrees with the hypothesis that Bryozoa *sensu lato* is monophyletic (Nielsen 1971, 1985, 2001; Cavalier-Smith 1998). Alternative hypotheses concerning the phylogenetic position of ectoprocta, namely that they are sister to all other Lophotrochozoa including Platyzoa, i.e. Platyhelminthes, Syndermata and related groups (Halanych *et al.* 1995; Littlewood *et al.* 1998; Halanych 2004; Passamanek & Halanych 2006), or that they are sister to all other protostomes except chaetognaths (Giribet *et al.* 2000) could be rejected (table 1, hypotheses 4–5).

Peterson & Eernisse (2001) defined several nested clades within Lophotrochozoa, namely (i) Neotrochozoa, which unites Mollusca and Annelida (with the annelids including Echiura and Sipuncula; see Hausdorf *et al.* (2007) and Struck *et al.* (2007)), (ii) Eutrochozoa, which includes Neotrochozoa and Nemertea, and (iii) Trochozoa, which comprises Eutrochozoa and Entoprocta. This last hypothesis, which we could not rule out with the previous dataset (Hausdorf *et al.* 2007), is now rejected by the expected likelihood weights method relying on the enlarged dataset (table 1, hypothesis 6). Although the more conservative approximately unbiased test is still marginally insignificant, this strengthens the evidence for the monophyly of Bryozoa *sensu lato*. On the other hand,

neither the Neotrochozoa hypothesis nor the Eutrochozoa hypothesis is rejected by either test method (table 1, hypotheses 7–8).

Brachiopods plus phoronids appear as the sister group of nemerteans in the maximum-likelihood tree (figure 1*a*). By contrast, the Bayesian inference analysis shows a sister-group relationship of Brachiozoa and Eutrochozoa (figure 1*b*). The relationships of Brachiozoa within Lophotrochozoa thus remain uncertain. However, we can dismiss the Conchozoa hypothesis (Cavalier-Smith 1998; Mallatt & Winchell 2002), according to which Brachiozoa is the sister group of Mollusca (table 1, hypothesis 9).

As mentioned earlier, the three traditional lophophorate lineages, Ectoprocta, Phoronida and Brachiopoda, did not join into a monophyletic clade in our trees (figure 1). The monophyly of Lophophorata was rejected with the expected likelihood weights method, but not with the approximately unbiased test (table 1, hypothesis 10). If we constrain the monophyly of Lophophorata, it becomes the sister group of Eutrochozoa in the resulting maximum-likelihood tree (not shown). In this tree, Entoprocta is the sister group of Lophophorata plus Eutrochozoa. Even if this topology should prove correct, the radial cleavage of Lophophorata would be a secondary modification derived from spiral cleavage, given that the spiral cleavage of Entoprocta is homologous to that of Annelida and Mollusca.

When we constrain the monophyly of Eutrochozoa (table 1, hypothesis 7), then Brachiozoa and Bryozoa (including Ectoprocta and Entoprocta) form a monophyletic group in the resulting maximum-likelihood tree. The same maximum-likelihood tree results if we constrain the monophyly of Brachiozoa and Bryozoa. Thus, the test results (table 1, hypothesis 7) apply to this hypothesis as well. This extended version of ‘Lophophorata’ including Entoprocta is therefore part of the confidence set of trees, given our ribosomal protein dataset, a possibility that is especially interesting, because it is in better agreement with morphological data than topologies that suggest independent origins of Ectoprocta and Brachiozoa within Lophotrochozoa. Potential synapomorphies of Brachiozoa and Bryozoa are the transition to a sessile lifestyle accompanied by the evolution of a horseshoe-shaped, tentacular feeding apparatus and a hydrostatic skeleton consisting of a lophophore coelom and a trunk coelom. In this view, both coelomic cavities were connected in the common ancestor of the two bryozoan subgroups and then were lost in Entoprocta. Most potential synapomorphies of Brachiozoa and Bryozoa are characters that were once thought to support a sister-group relationship between Lophophorata and Deuterostomia, but in light of the present evidence that these two groups are unrelated, must have originated by convergence (see above). Hypotheses that suppose that Ectoprocta and Brachiozoa originated independently of each other from different lophotrochozoan ancestors would require additional convergences of these characters.

Despite the progress presented herein, the resolution achieved in our analyses is still insufficient to fully reconstruct the evolutionary history of Lophotrochozoa. This lack of resolution could neither be avoided by the inclusion of many riboprotein genes and all major lophotrochozoan taxa, nor by the use of the CAT

model, which has been shown often to overcome long-branch attraction artefacts when other models fail (Baurain *et al.* 2007; Lartillot *et al.* 2007). Actually, the grouping of taxa with the longest branches in the maximum-likelihood tree (figure 1*a*), namely Syndermata and Platyhelminthes, is dissolved in the Bayesian inference reconstruction calculated with the CAT model (figure 1*b*). Further systematic errors unaccounted for by the present tree reconstruction methods, aggravated by the presumably rapid radiation of the lophotrochozoan taxa in the Late Precambrian and the limited taxon sampling within many phyla, might be responsible for the lack of resolution within Lophotrochozoa, which has been observed both here and in other studies (Halanych *et al.* 1995; Giribet *et al.* 2000; Peterson & Eernisse 2001; Mallatt & Winchell 2002; Ruiz-Trillo *et al.* 2002; Anderson *et al.* 2004; Passamaneck & Halanych 2006; Helmkamp *et al.* 2008). Improved models of molecular evolution and further taxonomic sampling within lophophorates and other lophotrochozoans will hopefully solve these issues in the future.

*Added in preparation.* While our manuscript was submitted, Dunn *et al.* (2008) published an important phylogenomic analysis of a huge number of new metazoan EST data. Regarding the relationships of brachiopods and phoronids, our maximum-likelihood tree (figure 1*a*) corresponds closely with the results presented by Dunn *et al.* (2008). In both analyses, brachiopods and phoronids form a clade with nemerteans (clade A in Dunn *et al.* 2008) that is the sister group of annelids (including sipunculans). These groups together (clade B in Dunn *et al.* 2008) are sister to the molluscs (together called clade C in Dunn *et al.* 2008). However, the results of our analyses differ from those of Dunn *et al.* (2008) with regard to the relationships of ectoprocts and entoprocts. Whereas these two groups form a well-supported clade in our analyses, their position is unstable in the analyses of Dunn *et al.* (2008). In the 77-taxon analysis of Dunn *et al.* (2008; figure 1), ectoprocts are sister to Platyzoa and entoprocts are sister to clade C.

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