

# When Three Trees Go to War

Leo van Iersel, Mark Jones, Mathias Weller

# ▶ To cite this version:

Leo van Iersel, Mark Jones, Mathias Weller. When Three Trees Go to War. 2023. hal-04013152v2

# HAL Id: hal-04013152 https://hal.science/hal-04013152v2

Preprint submitted on 17 Aug 2023 (v2), last revised 19 Sep 2023 (v3)

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution 4.0 International License

# When Three Trees Go to War

Leo van Iersel<sup>1</sup>, Mark Jones<sup>1</sup>, and Mathias Weller<sup>2,3</sup>

<sup>1</sup> Delft Institute of Applied Mathematics, Delft University of Technology – The Netherlands

<sup>2</sup> Technische Universität Berlin – Germany

<sup>3</sup> CNRS, LIGM (UMR 8049), Université Gustave Eiffel – France

Correspondence: mathias.weller@cnrs.fr

#### Abstract

How many reticulations are needed, in the worst case, for a rooted phylogenetic network to display a given set of k rooted binary phylogenetic trees on n leaves? For k = 2, Baroni, Semple, and Steel [Ann. л Comb. 8, 391–408 (2005)] showed that the answer is n-2. Here, we show that, for  $k \geq 3$  the answer 5 is at least  $(3/2 - \epsilon)n$ . Concretely, we prove that, for each  $\epsilon > 0$ , there is some  $n \in \mathbb{N}$  such that three 6 n-leaf caterpillar trees can be constructed in such a way that any network displaying these caterpillars contains at least  $(3/2 - \epsilon)n$  reticulations. The case of three trees is interesting since it is the easiest case that cannot be equivalently formulated in terms of agreement forests. Instead, we base the result on a 9 surprising lower bound for multilabelled trees (MUL-trees) displaying the caterpillars. Indeed, we show 10 that one cannot do (more than an  $\epsilon$ ) better than the trivial MUL-tree resulting from a simple concatenation 11 of the given caterpillars. The results are relevant for the development of methods for the Hybridization 12 Number problem on more than two trees. This fundamental problem asks to construct a rooted phylo-13 genetic network with a minimum number of reticulations displaying a given set of rooted phylogenetic trees. 14

2

15

16

Keywords: Phylogenetic Networks, Tree Containment, Caterpillar Tree, Hybridization Number

#### 1 Introduction

A fundamental task of evolutionary analysis is to construct a phylogeny for a set of taxa depicting their 18 ancestral relations. While many biological studies are content with the simplification that this phylogeny is 19 a tree, there are circumstances, such as the presence of horizontal gene transfer (observed in many bacte-20 ria [DARM08]) or hybridization (common among plant species and also observed among animals [Mal05]), 21 that require constructing phylogenetic *networks* which, in contrast to trees, allow modeling such "reticulate" 22 evolution. Mathematically, a rooted phylogenetic network is a directed acyclic graph (DAG) with a single root 23 and leaves bijectively labelled by the elements of a set X, modeling the considered taxa. In this paper, we 24 will only consider phylogenetic networks that are rooted and binary. In such networks, the indegree-2 nodes 25 represent reticulate evolutionary events and are called *reticulations*. 26

The construction of the most parsimonious (that is, containing the least amount of reticulations) network 27 that is still "compatible" with a given set of trees is modeled by the Hybridization Number problem, which is 28 well understood for the special case of having exactly two input trees [BS07; BSS05; Bar+05; IK11; Kel+12]. To 29 formalize this problem, we say that a network *displays* a tree if this tree can be obtained from a subgraph of 30 the network by suppressing nodes with exactly one incoming and exactly one outgoing arc. The Hybridization 31 Number problem then asks for a smallest network (in terms of the number of reticulations) displaying all input 32 trees. Throughout this paper, we will focus on the simplified version of this problem where all input trees and 33 the output network are required to be binary and all have the same set of leaf labels. 34

Unfortunately, many observations made for this case do not generalize to more than two input trees. One 35 such observation is that for any two trees with n leaves, there is always a network with n-2 reticulations 36 displaying the two trees. This bound is tight because two "inverse" caterpillar trees need exactly n-2 reticulations [BSS05]. In this work, we show that, for three or more trees, at least  $(3/2 - \epsilon)n$  reticulations may 38 be required, even if the trees are caterpillars. See Fig. 1 for an example of our construction. This result rep-39 resents a first lower bound for more than two trees that improves upon the n-2 bound. In particular, it 40 refutes the tempting conjecture that n reticulations are sufficient to display any set of three phylogenetic 41 trees. If the bound of n had held, it would have had positive consequences for the development of methods 42 for the Hybridization Number problem, by bounding the worst-case complexity of subnetworks that need to 43 be considered inside an algorithm.

To prove the  $(3/2 - \epsilon)n$  bound, we fist derive a corresponding bound for "multilabelled trees" (MUL-trees), 45 that is, trees in which each leaf has one label, but each label may be used more than once. Again, the goal 46 is to find a smallest (in terms of the number of leaves) MUL-tree displaying all input trees. Surprisingly, we 47 show that, given at most three caterpillars, one cannot do better (up to an  $\epsilon$ ) than the trivial MUL-tree that 48 simply concatenates the given caterpillars. More precisely, we show that, for each  $\epsilon > 0$  and  $t \leq 3$ , there is 49 some  $n \in \mathbb{N}$ , and t caterpillars with n leaves, such that any MUL-tree displaying the caterpillars has at least 50  $(t-\epsilon)n$  leaves. This is very close to the upper bound of  $t \cdot n$  for any set of t trees, which holds because the t 51 trees can simply be concatenated into a single MUL-caterpillar with  $t \cdot n$  leaves. 52

Upper bounds on the number of reticulations needed to display a set of trees follow from results on "universal tree-based" networks [BS18], which are, roughly speaking, networks that can be obtained from *any tree* on the same set of leaves by subdividing arcs of the tree and adding arcs between subdividing nodes [FS15; Hay16; Zha16]. Bordewich and Semple [BS18] showed that such a network has  $\Theta(n \log n)$  reticulations. The lower bound does not (directly) carry over to our question, but the upper bound does. Concretely, Bordewich and Semple [BS18] proved that any network displaying all phylogenetic trees on n leaves needs at least  $O(n \log n)$ reticulations.

The structure of this paper is as follows. After the preliminaries in Section 2, we prove the bound for MULtrees in Section 3 and the bound for networks in Section 4, concluding with some open problems in Section 5.



**Figure 1.** Three caterpillar trees with 9 leaves, resulting from Construction 1 for  $n = 3^i = 9$ . Lemma 2 implies that any MUL-tree displaying these caterpillars needs at least 19 leaves. Lemma 3 further implies that any network displaying these caterpillars needs at least 2 reticulations. While this second number does not seem particularly surprising, the strength of Lemma 3 is in the *asymptotic bound* it provides for growing *n*.

# 2 Preliminaries

We will deal with sequences of letters over an unspecified alphabet. To differentiate such letters from variable names (even those referring to letters), we will typeset them in typewriter font such as a, 3, B. For all sequences s' that can be produced from s by removing zero or more letters, we say that s' is a *subsequence* of s and we write  $s' \leq s$ . We use  $\circ$  to denote the usual concatenation operator on sequences, where  $s \circ s'$  denotes the result of writing out s' after s. For letters a and b, we write a  $\leq_s$  b if the last occurence of a preceeds the first occurence of b in s, that is, some prefix of s contains all occurences of a but no occurence of b.

We also deal with (binary, phylogenetic) MUL-networks, which are directed, acyclic graphs (DAGs) with only 69 the following types of nodes: (1) a unique source (called the *root*) with out-degree zero or two; (2) sinks (called 70 *leaves*) with in-degree at most one, labeled using a function  $\mathcal{L}$  from the set of leaves to some set of labels; (3) in-71 degree one and out-degree two nodes (tree nodes); and (4) in-degree two, out-degree one nodes (reticulation 72 nodes or reticulations). A MUL-tree is a MUL-network without reticulations. A network is a MUL-network whose 73 labelling function  $\mathcal{L}$  is injective. A *tree* is a network without reticulations. A *caterpillar tree* (or just a *caterpillar*) 74 is a tree in which each node is either a leaf or the parent of a leaf. If X is a MUL-network, a subset of nodes in 75 a MUL-network, or a sequence, then  $\mathcal{L}(X)$  is the set (not the multiset) of labels/letters occurring in X. If X is a 76 MUL-network, then n(X) denotes the number of leaves in X. If X is a sequence, then n(X) denotes its length. 77

We use the following correspondence between sequences and caterpillars. We say that sequence s corresponds to caterpillar P if the elements of s are exactly the leaf labels of P and these labels are ordered responds to caterpillar P if the elements of s are exactly the leaf labels of P and these labels are ordered responding distance from the root in P. Observe that each caterpillar has exactly two corresponding sequences since it has exactly two leaves with the same distance from the root. Conversely, each sequence has exactly one corresponding caterpillar. See Fig. 1 for an example of three caterpillars with corresponding sequences abc123ABC, CABcab312 and 231BCAbca.

Let N be a MUL-network and let u be a node of N. If u has a descendant v in N (that is, N contains a 84 directed *u-v*-path), then we write  $v \leq_N u$  and we call *u* an *ancestor* of *v*. Note that the " $\leq_X$ "-relations for 85 MUL-networks and sequences X naturally extend to sets of nodes/letters, that is,  $P <_X Q$  if  $p <_X q$  for all 86  $p \in P$  and  $q \in Q$ . For a set L of nodes of N, the MUL-network of N induced by L (written N[L]) is the result of 87 removing all nodes u of N that have no descendant in L followed by the exhaustive contraction of indegree-88 one outdegree-one nodes onto their respective parents (this last operation is also called "supression"). If L89 is a set of labels, then N[L] is the MUL-network of N induced by all nodes with a label in L. The result of 90 removing all nodes x from N with  $x \leq N$  u is denoted by  $N_u$  and, if u is a reticulation and  $N_u$  does not contain 91 any reticulations of N, then u is called a *lowest reticulation*. If N is a MUL-tree and x and y are nodes in N, then 92 the *lowest common ancestor* (LCA) of x and y in N is the unique minimum with respect to " $\leq_N$ " of all nodes u 93 of N such that  $N_u$  contains both x and y. 94

An *embedding* of a MUL-network T into a MUL-network N is a function  $\phi$  that maps the nodes of T to nodes of N and the arcs of T to directed paths in N such that

- 1. the paths in the image of  $\phi$  are arc-disjoint;
- 2. for each arc uv of T,  $\phi(uv)$  starts in  $\phi(u)$  and ends in  $\phi(v)$  in N.

97 98

_	i	0	1	2	3
-	$X_i$	$\lambda$	a1A	abc123ABC	abcdefghi123456789ABCDEFGHI
	$Y_i$	$\lambda$	Aa1	CABcab312	IGHCABFDEighcabfde978312645
	$Z_i$	$\lambda$	1Aa	231BCAbca	564897231EFDHIGBCAefdhigbca

**Figure 2.** Example of Construction 1. For i = 1, the functions  $r_1$ ,  $r_2$ , and  $r_3$  map  $\lambda$  to a, 1, and A, respectively. For i = 2, they map  $\{a, 1, A\}$  to  $\{a, b, c\}$ ,  $\{1, 2, 3\}$ , and  $\{A, B, C\}$ , respectively. In particular, sequence  $X_2$  is given by  $r_1(a1A) \circ r_2(a1A) \circ r_3(a1A) = abc123ABC$ , while  $Y_2$  is given by  $r_3(Aa1) \circ r_1(Aa1) \circ r_2(Aa1) = CABcab312$ . For i = 3,  $r_1$  maps  $\{a, b, c, 1, 2, 3, A, B, C\}$  to  $\{a, b, c, d, e, f, g, h, i\}$ , and analogously for  $r_2$ ,  $r_3$ .

We say that MUL-network N displays MUL-network T if there is an embedding of T into N.

The *backbone* of a caterpillar is the path containing all edges not incident to a leaf. The *backbone* of an <sup>100</sup> embedding  $\phi$  of a caterpillar P in a MUL-network N is the path obtained by merging the paths  $\phi(e)$  for all <sup>101</sup> edges e on the backbone of P.

#### 3 Lower Bound on MUL-Trees

In this section, we construct a family C of triples of caterpillars such that for any  $\epsilon > 0$ , the familiy C 104 contains a triple  $(C_1, C_2, C_3)$  of *n*-leaf caterpillars (where *n* depends on the choice of  $\epsilon$ ) such that any MULtree displaying all three caterpillars has at least  $(3 - \epsilon)n$  leaves. As a byproduct, we show the existance of a family of *pairs* of caterpillars with a  $(2 - \epsilon)n$  lower-bound on the leaf-number in any displaying MUL-tree.

A *relabeling* is a function mapping a label to another label and we allow applying relabelings to sets of labels, sequences and (MUL-)trees in the natural way.

**Construction 1.** Let  $C_0 = (X_0, Y_0, Z_0)$  denote the triple of sequences on a single label  $\lambda$ . For each i > 0, we recursively construct a triple  $C_i = (X_i, Y_i, Z_i)$  of sequences of length  $3^i$  as follows: Let  $r_1, r_2$ , and  $r_3$  be relabelings defined on the labels of  $C_{i-1}$  with disjoint images. Then,

1. 
$$X_i := r_1(X_{i-1}) \circ r_2(X_{i-1}) \circ r_3(X_{i-1})$$
 113

2.  $Y_i := r_3(Y_{i-1}) \circ r_1(Y_{i-1}) \circ r_2(Y_{i-1})$ 

3.  $Z_i := r_2(Z_{i-1}) \circ r_3(Z_{i-1}) \circ r_1(Z_{i-1})$ 

Note that  $\mathcal{L}(r_1(X_{i-1}))$ ,  $\mathcal{L}(r_1(Y_{i-1}))$  and  $\mathcal{L}(r_1(Z_{i-1}))$  are identical and we refer to this set by  $A_i$ . Similarly, we abbreviate  $B_i := \mathcal{L}(r_2(X_{i-1}))$  and  $C_i := \mathcal{L}(r_3(X_{i-1}))$ .

It turns out that sequences constructed by Construction 1 have very short common subsequences.

**Proposition 1.** Let i > 0 and let  $(X_i, Y_i, Z_i)$  be a triple of sequences constructed by Construction 1. Let  $k \in \{1, 2, 3\}$  and let  $s_k$  be a common subsequence of any k of the three sequences. Then,  $|s_k| \le (4-k)^i$ .

*Proof.* Clearly, the claim trivially holds for k = 1 so we consider  $k \in \{2, 3\}$  in the following.

**Case 1:** k = 3. The proof is by induction on *i*. For i = 0, all three sequences contain a single label  $\lambda$  so the 122 claim is trivially true. Suppose in the following that the claim holds for i - 1. Let  $s_3$  be a common subsequence 123 of  $X_i$ ,  $Y_i$ , and  $Z_i$ . Since all labels in the image of  $r_1$  preceed all labels in the image of  $r_3$  in  $X_i$  and all labels 124 in the image of  $r_3$  preceed all labels in the image of  $r_1$  in  $Y_i$ , we know that  $s_3$  does not contain labels of the 125 images of both  $r_1$  and  $r_3$ . Similarly, it can be seen that  $s_3$  cannot contain labels of any two of  $r_1$ ,  $r_2$ , and  $r_3$ . Thus, without loss of generality,  $s_3$  consists only of labels of  $r_1$ , implying that  $s_3$  is a common subsequence 127 of the result of removing all labels of  $r_2$  and  $r_3$  from  $X_i$ ,  $Y_i$ , and  $Z_i$ , that is,  $s_3$  is a common subsequence of 128  $r_1(X_{i-1})$ ,  $r_1(Y_{i-1})$  and  $r_1(Z_{i-1})$ . But then,  $r_1^{-1}(s_3)$  is a common subsequence of  $X_{i-1}$ ,  $Y_{i-1}$  and  $Z_{i-1}$  and, 129 by induction hypothesis, the length of  $s_3$  is 1. 130

**Case 2:** k = 2. Again, the proof is by induction on i and the induction base case i = 0 is trivially true, so we will suppose that the claim holds for i-1. By symmetry, we can further suppose without loss of generality that s<sub>2</sub> is a common subsequence of  $X_i$  and  $Y_i$ . If  $s_2$  only uses labels from the image of one  $r \in \{r_1, r_2, r_3\}$ , then 133

103

114

115

118

121

 $r^{-1}(s_2)$  is a common subsequence of  $X_{i-1}$  and  $Y_{i-1}$  so the claim holds by induction hypothesis. Otherwise,  $s_2$  uses labels of at least two of  $r_1$ ,  $r_2$ , and  $r_3$ . Since all labels of  $r_3$  succeed all labels of  $r_1$  and  $r_2$  in  $X_i$  but preceed them in  $Y_i$ , we know that  $s_2$  uses labels of  $r_1$  and  $r_2$  but not of  $r_3$ . Thus,  $s_2$  admits two subsequences s'and s'' such that s' and s'' contain only labels of  $r_1$  and  $r_2$ , respectively, and  $|s'| + |s''| = |s_2|$ . Then, however, s' is a subsequence of the result of removing all labels of  $r_2$  and  $r_3$  from  $X_i$  and  $Y_i$ , that is, of  $r_1(X_{i-1})$  and  $r_1(Y_{i-1})$  (see Construction 1). Thus,  $r_1^{-1}(s')$  is a common subsequence of  $X_{i-1}$  and  $Y_{i-1}$  and, by induction hypothesis,  $|s'| \leq 2^{i-1}$ . An analogous argument shows that  $|s''| \leq 2^{i-1}$  and, thus,  $|s_2| = |s'| + |s''| \leq 2^i$ .  $\Box$ 

In the following, we prove lower bounds on the number of leaves in any MUL-tree displaying k of the the caterpillars in  $C_i$  for  $k \in \{1, 2, 3\}$  and  $i \in \mathbb{N}$ . We denote these bounds by  $N_i^{(k)}$  and we note that, by the concatenation argument,  $N_i^{(3)} \leq N_i^{(2)} + N_i^{(1)}$  and  $N_i^{(2)} \leq 2N_i^{(1)}$  and  $N_i^{(1)} = n(X_i) = n(Y_i) = n(Z_i) = 3^i$ .

**Lemma 1.** Let  $i \in \mathbb{N}$  and let T be any MUL-tree displaying  $X_i$  and  $Y_i$ . Then,  $n(T) \ge 2 \cdot 3^i - 2^i$ .

*Proof.* The proof is by induction on *i*. For the induction base, observe that all of  $X_0$ ,  $Y_0$  and T consist of a single leaf and n(T) = 2 - 1 = 1. For the induction step, suppose that the lemma holds for all j < i. Let  $T_A$ ,  $T_B$  and  $T_C$  denote the subtrees of T induced by labels in  $A_i$ ,  $B_i$  and  $C_i$ , respectively, and observe that their label multisets are a partition of the label multiset of T since  $A_i$ ,  $B_i$  and  $C_i$  are disjoint. In the following, we show that at least one of  $T_A$ ,  $T_B$  and  $T_C$  contains  $2N_{i-1}^{(1)}$  leaves. Since, by definition, the other two contain at least  $N_{i-1}^{(2)}$  leaves, we have

$$n(T) = n(T_A) + n(T_B) + n(T_C) \ge 2N_{i-1}^{(2)} + 2N_{i-1}^{(1)} \stackrel{\text{Ind. Hyp.}}{\ge} 2(2 \cdot 3^{i-1} - 2^{i-1}) + 2 \cdot 3^{i-1} = 2 \cdot 3^i - 2^i - 2^i \cdot 3^i - 2^i - 2^i \cdot 3^i - 2^i - 2^i$$

Towards a contradiction, assume that  $T_A$ ,  $T_B$  and  $T_C$  contain less than  $2N_{i-1}^{(1)}$  leaves. For all  $F \in \{A_i, B_i, C_i\}$ , 145 let  $\chi_F$  and  $\psi_F$  be respective embeddings of  $X_i[F]$  and  $Y_i[F]$  into  $T_F$  and note that, by assumption,  $T_F$  contains strictly less than  $2N_{i-1}^{(1)} = n(X_i) + n(Y_i)$  leaves, implying that some leaf  $\ell_F$  in  $T_F$  is mapped-to by both  $\chi_F$  and  $\psi_F$ . In particular, the label of each  $\ell_F$  occurs only once in  $T_F$  and, thus, in T, which allows us to use  $\ell_F$ and its label interchangeably. Now, since  $A_i <_{X_i} B_i <_{X_i} C_i$  we know that  $LCA(\ell_{A_i}, \ell_{C_i})$  is a strict ancestor of  $LCA(\ell_{A_i}, \ell_{B_i})$  in  $X_i$  and, since T displays  $X_i$ , this also holds in T. But since  $C_i <_{Y_i} A_i <_{Y_i} B_i$  we also know that  $LCA(\ell_{A_i}, \ell_{B_i})$  is a strict ancestor of  $LCA(\ell_{A_i}, \ell_{C_i})$  in T, which is clearly a contradiction.

**Corollary 1.** Let  $\epsilon > 0$ . Then, there is some  $n \in \mathbb{N}$  and two caterpillar trees of the same set of n labels, such that any MUL-tree displaying them has at least  $(2 - \epsilon)n$  leaves.

*Proof.* Let  $i \in \mathbb{N}$  such that  $(2/3)^i \leq \epsilon$  and, hence,  $2^i \leq 3^i \epsilon = n\epsilon$ . Let T be any MUL-tree displaying  $X_i$  and  $Y_i$ . Then, by Lemma 1,  $n(T) \geq 2 \cdot 3^i - 2^i \geq 2n - n\epsilon = (2 - \epsilon)n$ .

**Lemma 2.** Let  $i \in \mathbb{N}$  and let T be any MUL-tree displaying  $X_i$ ,  $Y_i$  and  $Z_i$ . Then,  $n(T) \ge 3^{i+1} - 2^{i+1}$ .

*Proof.* The proof is by induction on *i*. For the induction base, observe that all of  $X_0$ ,  $Y_0$ ,  $Z_0$ , and *T* consist of a single leaf and n(T) = 3 - 2 = 1. For the induction step, suppose that the lemma holds for all j < i. Let  $T_A$ ,  $T_B$  and  $T_C$  denote the subtrees of *T* induced by labels in  $A_i$ ,  $B_i$  and  $C_i$ , respectively, and observe that their label multisets are a partition of the label multiset of *T* since  $A_i$ ,  $B_i$  and  $C_i$  are disjoint. If any of  $T_A$ ,  $T_B$ , and  $T_C$  contains  $3N_{i-1}^{(1)} = 3^i$  leaves, then

$$n(T) = n(T_A) + n(T_B) + n(T_C) \ge 2N_{i-1}^{(3)} + 3N_{i-1}^{(1)} \stackrel{\text{Ind. Hyp.}}{\ge} 2(3^i - 2^i) + 3^i = 3^{i+1} - 2^{i+1}.$$

Further, if any two of  $T_A$ ,  $T_B$ , and  $T_C$  contain  $N_{i-1}^{(2)} + N_{i-1}^{(1)} \stackrel{\text{Lem 1}}{\geq} 2 \cdot 3^{i-1} - 2^{i-1} + 3^{i-1} = 3^i - 2^{i-1}$  leaves, then

$$n(T) = n(T_A) + n(T_B) + n(T_C) \ge N_{i-1}^{(3)} + 2(N_{i-1}^{(2)} + N_{i-1}^{(1)}) \stackrel{\text{Ind. Hyp.}}{\ge} 3^i - 2^i + 2(3^i - 2^{i-1}) = 3^{i+1} - 2^{i+1}$$

Thus, in the following, suppose that neither of the two cases holds. In particular, at least two trees among  $T_A$ ,  $T_B$ , and  $T_C$  contain strictly less than  $N_{i-1}^{(2)} + N_{i-1}^{(1)}$  leaves. By symmetry, suppose these are  $T_A$  and  $T_B$ . 158

156



**Figure 3.** Illustration of the five leaves handled in the proof of Lemma 2. The three parts depict possible embeddings of  $X_i[F]$ ,  $Y_i[F]$ , and  $Z_i[F]$  for all  $F \in \{A_i, B_i, C_i\}$ . We assume that the embedding  $\psi_A$  of  $Y_i[A_i]$  overlaps  $\chi_A$  in  $k_A$  and  $\phi_A$  in  $\ell_A$  (left part) and, likewise for B. For C, only  $\chi_C$  and  $\phi_C$  overlap, in  $\ell_C$  (right part). While T is not necessarily a caterpillar, drawing T linearly like that may help understand the situation.

For each  $F \in \{A, B, C\}$ , let  $\chi_F$ ,  $\psi_F$ , and  $\phi_F$  denote the respective embeddings of  $X_i[F_i]$ ,  $Y_i[F_i]$  and  $Z_i[F_i]$ 159 into  $T_F$  and let us say that two among them *overlap* if  $T_F$  has a leaf that is assigned to by both. Note that 160 removing the  $n(X_i[A_i]) = n(X_{i-1}) = N_{i-1}^{(1)}$  leaves of  $T_A$  that are mapped-to by  $\chi$  results in a MUL-tree with 161 strictly less than  $N_{i-1}^{(2)}$  leaves and, by Lemma 1, this MUL-tree cannot display both  $Y_i[A_i]$  and  $Z_i[A_i]$ ). Thus, 162  $\chi_A$  overlaps one of  $\psi_A$  and  $\phi_A$  and the analog holds for  $\psi_A$  and  $\phi_A$ . By pigeonhole principle, one among 163 the three embeddings overlaps both others (while the other two may not necessarily overlap). Let  $\ell_A$  and  $k_A$ 164 denote the corresponding leaves (possibly  $\ell_A = k_A$  if all three embeddings assign to  $\ell_A$ ). Since the same 165 argument holds for B, we define  $\ell_B$  and  $k_B$  analogously (see Fig. 3 for an illustration). Now, since  $T_C$  contains 166 strictly less than  $3N_{i-1}^{(1)}$  leaves by assumption, we know that two of  $\chi_C$ ,  $\psi_C$ , and  $\phi_C$  overlap in a leaf  $\ell_C$  of  $T_C$ . 167 Next, we consider the relative positions of these five leaves in T. 168

In the following, a leaf a in  $T_A$  with parent t is said to be *above* a leaf b in  $T_B$  (written  $a \rightsquigarrow b$ ) if  $b <_T t$ , and analogously for any pair chosen from the combined leaf-set of  $T_A$ ,  $T_B$  and  $T_C$ . A third leaf  $\ell$  is said to be *between* a and b if  $a \rightsquigarrow \ell \rightsquigarrow b$ . For all leaves a whose parent is an ancestor of b in  $T_A$ , we have that a is above both a and b and, likewise, for  $T_B$  and  $T_C$ . By symmetry, suppose that  $\ell_F \rightsquigarrow k_F$  for all  $F \in \{A, B\}$ 

Now, as  $C <_{Y_i} A$  and  $C <_{Z_i} A$ , all leaves of  $\psi_C$  and  $\phi_C$  are below all leaves of  $\psi_A$  and  $\phi_A$ . Since  $\ell_C$  is contained in the former and both  $\ell_A$  and  $k_A$  are contained in the latter, we have that  $\ell_A$  and  $k_A$  are both above  $\ell_C$ . Further, as  $B <_{X_i} C$  and  $B <_{Z_i} C$ , we have  $\ell_C \rightsquigarrow \ell_B$ . Thus,  $\ell_A \rightsquigarrow \ell_C \rightsquigarrow \ell_B \rightsquigarrow k_B$ , in particular both of  $\ell_A, k_A$  are above both of  $\ell_B, k_B$ . However, the caterpillar  $X_i$  contains at least one leaf mapped to  $\ell_A$  or  $k_A$ , and at least one leaf mapped to  $\ell_B$  or  $k_B$ . But since  $A <_{X_i} B$ , this implies that at least one of  $\ell_B$  and  $k_B$  is above one of  $\ell_A$  and  $k_A$ , contradicting  $\ell_A \rightsquigarrow k_A \rightsquigarrow \ell_B \rightsquigarrow k_B$ .

**Corollary 2.** Let  $\epsilon > 0$ . Then, there is some  $n \in \mathbb{N}$  and three caterpillar trees of the same set of n labels, such that any MUL-tree displaying them has at least  $(3 - \epsilon)n$  leaves.

*Proof.* Let  $i \in \mathbb{N}$  such that  $(2/3)^i \le \epsilon/2$  and, hence,  $2^{i+1} \le 3^i \epsilon = n\epsilon$ . Let T be any MUL-tree displaying  $X_i$ ,  $Y_i$  and  $Z_i$ . Then, by Lemma 2,  $n(T) \ge 3^{i+1} - 2^{i+1} \ge 3n - n\epsilon = (3 - \epsilon)n$ .

#### 4 Lower Bound on Networks

In this section, we build on the lower bound developed for MUL-trees in Section 3 to prove that, for any  $\epsilon > 0$  and large enough n, any single-labeled phylogenetic network displaying the three n-leaf caterpillars constructed in Construction 1 has at least  $(3/2 - \epsilon)n$  reticulations. To this end, we will give an algorithm that transforms any network displaying the caterpillars into a MUL-tree displaying the caterpillars by "unzipping" (or 187



**Figure 4.** Illustration of the operation of "unzipping"  $(N, \phi)$  at a lowest reticulation. The embedding of the three caterpillars  $X_i$ ,  $Y_i$  and  $Z_i$  is depicted as green solid, red dashed and blue dotted lines, respectively, within the network outlined in gray. Note that all leaves of  $N_z$  into which  $\phi$  embeds leaves of  $X_i$  as well as at least one of  $Y_i$  and  $Z_i$ , are duplicated in the process. Note also that not all three caterpillars are necessarily embedded in  $N_z$ , as previous unzip operations may have split a caterpillar off  $N_z$ .

"duplicating") subtrees. Then, we show that, if the network had fewer than  $(3/2 - \epsilon)n = (3/2 - \epsilon)3^i$  reticulations, then the resulting MUL-tree has fewer than  $3^{i+1} - 2^{i+1}$  leaves, contradicting Lemma 2.

190

212

213

#### 4.1 Transforming the Network into a MUL-tree

In the following, we present a transformation acting on a given embedding of the three caterpillars into a 191 multi-labeled network. Each application of our transformation rule will reduce the number of reticulations by 192 one at the cost of creating new leaves. Hence, exhaustive application will result in an embedding of the three 193 caterpillars into a MUL-tree. The rule acts on the subtree below a lowest reticulation and also manipulates a 194 reservoir of virtual "tokens" which will help in the amortized analysis of how many new leaves are created in 195 the process.

In the following, we work with pairs  $(N, \phi)$ , where N is a MUL-network with  $3^i$  distinct labels and  $\phi$  is an 197 embedding of the caterpillars  $X_i$ ,  $Y_i$  and  $Z_i$  (as constructed by Construction 1) into N such that all arcs of N are 198 used by the embedding  $\phi$ . We call such pairs *caterpillar embeddings*. Note that the assumptions that all arcs of 199 N are used is satisfied by any embedding of the caterpillars into a network with smallest reticulation number. 2000 We make use of the fact that no embedding of any caterpillar can use both arcs incoming to any reticulation r201 of N, so the caterpillars with leaves embedded below r can be divided into two groups, depending on which incoming arc of r is used in their embedding. We call this the *parity* of a caterpillar with respect to r. We say 203 that a caterpillar that does not have leaves below r has parity  $\perp$  with respect to r. Note that, since all arcs 204 of N are used by  $\phi$ , there are two caterpillars with different non- $\perp$  parity with respect to r. Let  $N_r$  be the 205 subnetwork of N rooted at r. We say that the backbone of a caterpillar Q is embedded in  $N_r$  (or below r) if  $\phi$ 206 maps a non-leaf of Q into  $N_r$ . Note that this is the case if and only if at least two leaves of  $N_r$  are used by the 207 embedding of Q into N. The central operation in the transformations "unzips"  $(N, \phi)$  at r (see Fig. 4). 208

**Definition 1.** Let  $(N, \phi)$  be a caterpillar-embedding, let r be a reticulation in N with child z such that the subnetwork  $N_z$  of N rooted at z does not contain reticulations. Let xr and yr denote the incoming arcs of r with  $x \neq y$ . The operation of unzipping N at r consists in the following steps:

- 1. Remove the node r from N.
- 2. Add a copy  $N_z^\prime$  of  $N_z$  with root  $z^\prime$  to N and add the arcs xz and  $yz^\prime.$
- 3. For each caterpillar Q such that  $\phi$  embeds Q using the arc yr, replace all nodes u of  $N_z$  by their copy u' in  $N'_z$  in the embedding of Q.
- 4. repeatedly remove all leaves of the resulting MUL-network that are not used by the embedding, and suppress indegree-one outdegree-one nodes. 217

**Observation 1.** Let  $(N, \phi)$  be a caterpillar embedding and let  $(N', \phi')$  be the result of unzipping N at a reticulation r. Then,  $(N', \phi')$  is a caterpillar embedding. In particular, all arcs of N' are used by  $\phi'$ .

The rest of this section depends on an arbitrary number  $q \in \mathbb{N}$ , which we will pick "sufficiently large" in the proof of the main theorem.<sup>1</sup> Further, we also suppose that our input caterpillars are "sufficiently large" with respect to q, that is, their length n satisfies

$$n > 12qn^{\log_3 2}$$
. (1)

**Transformation Rule 1.** Let  $(N, \phi)$  be a caterpillar-embedding, let r be a lowest reticulation in N, and let Q 220 denote the set of caterpillars whose backbone is embedded in  $N_r$ . Then, 221

- 1. unzip N at r,
- 2. create three tokens in the token reservoir, and
- 3. for each leaf  $\ell$  below r in N and each pair of different-parity caterpillars in Q whose embedding uses  $\ell$ , remove 224 tokens from the token reservoir. 225

As we will see, we never need to remove more tokens than are contained in the token reservoir.

**Lemma 3.** Let N be a network with n leaves, let  $(N, \phi)$  be a caterpillar embedding, let k be the number of reticulations of N, and let  $(T, \phi')$  be the result of applying Transformation Rule 1 exhaustively to  $(N, \phi)$ . Then, T has at most n + 4(q+1)k/3q leaves.

Proof.Intuitively, the proof is based on the observation that, whenever the transformation creates many new230leaves for only a single reticulation it removes, then we can use half of these leaves to construct a common231subsequence of two caterpillars. Then, Proposition 1 implies that this cannot happen too often.232

Formally, we consider a series of "configurations"  $C_0, C_1, \ldots, C_{\Omega}$ , each consisting of a caterpillar embedding and a token reservoir where  $C_0 := ((N, \phi), t_0 = 0)$  and  $C_{\Omega} = ((T, \phi'), t_{\Omega})$  for some  $t_{\Omega}$  and each  $C_j$ results from an application of Transformation Rule 1 to the previous configuration  $C_{j-1}$ . To show Lemma 3, we assign a "weight"  $\omega$  to each  $C_i$ . We prove that  $\omega$  is monotonically non-increasing with respect to Transformation Rule 1. This will imply an upper bound on the number of leaves of the MUL-tree T displaying all three caterpillars. For a configuration  $C := ((\Gamma, \psi), t)$ ,

1. let  $\#_r^{\Gamma}$  denote the number of reticulations in  $\Gamma$ ,

2. let  $\#_i^{\Gamma}$  denote the number of leaves of  $\Gamma$  that are used by the embedding of exactly *i* caterpillars, and define

$$\omega((\Gamma,\psi),t) := \sum_{i \in \{1,2,3,r\}} c_i \cdot \#_i^{\Gamma} + c_t \cdot t \tag{2}$$

where  $c_1 := c_3 := 3q$ ,  $c_2 := 4q$ ,  $c_r := 4(q+1)$ , and  $c_t := 1$ . In the following, we omit the superscript  $\Gamma$  when the context and we abbreviate the total number of leaves as  $\#^{\Gamma} := \sum_{i \in \{1,2,3\}} \#^{\Gamma}_i$ .

**Claim 1.**  $\omega$  is monotonically non-increasing with respect to Transformation Rule 1.

*Proof.* We consider the following cases:

**Case 1:** No caterpillar has its backbone embedded in  $N_r$ . Then  $N_r$  has at most three leaves.

**Case 1a:**  $N_r$  contains two or three leaves, each with a single caterpillar embedded into it. Then, the numbers  $\#_i$  do not change for any i, so  $\omega$  increases by  $\Delta \omega = 3c_t - c_r \le 0$ .

**Case 1b:**  $N_r$  contains two leaves, a leaf  $\ell_1$  with a single caterpillar embedded into it and a leaf  $\ell_2$  with two caterpillars embedded into it. If the two caterpillars whose embedding uses  $\ell_2$  have the same parity, then  $\#_i$  249 does not change for any i, see Case 1a. Otherwise,  $\#_1$  grows by two and  $\#_2$  decreases by one, implying that  $\omega$  grows by  $\Delta \omega = (2c_1 + 3c_t) - (c_2 + c_r) = 6q + 3 - 8q - 4 \le 0$ .

**Case 1c:**  $N_r$  contains a single leaf  $\ell$  with exactly two caterpillars embedded into it (their parity must differ 252 in this case). Then,  $\#_1$  grows by two and  $\#_2$  decreases by one, implying that  $\omega$  grows by  $\Delta \omega = (2c_1 + 3c_t) - 253$  $(c_2 + c_r) = 6q + 3 - 8q - 4 \le 0.$ 

243

244

245

239

240

222

223

<sup>&</sup>lt;sup>1</sup>We note that the proofs that follow also work for  $q \in \mathbb{R}$ , but for ease of presentation we assume  $q \in \mathbb{N}$ .

**Case 1d:**  $N_r$  contains a single leaf  $\ell$  with three caterpillars embedded into it. Then,  $\#_1$  and  $\#_2$  each grow by one and  $\#_3$  decreases by one, implying that  $\omega$  grows by  $\Delta \omega = (c_1 + c_2 + 3c_t) - (c_3 + c_r) = 7q + 3 - 7q - 4 \le 0$ . 256

**Case 2:** Exactly one caterpillar Q has its backbone embedded in  $N_r$ . Let  $L_Q$  denote the set of all (at least two) leaves below r that leaves of Q are embedded into.

**Case 2a:** All caterpillars with a leaf embedded into a leaf of  $L_Q$  have the same parity as Q. Then, the numbers  $\#_i$  do not change for any i, so  $\omega$  increases by  $\Delta \omega = 3c_t - c_r \leq 0$ .

**Case 2b:** Exactly one leaf  $\ell$  of  $L_Q$  is used to embed a leaf of a caterpillar with different parity than Q. If  $_{261}$   $\ell$  is used by exactly one caterpillar with different parity than Q, then  $\#_1$  grows by two and  $\#_2$  decreases by  $_{262}$  one, implying that  $\omega$  grows by  $\Delta \omega = (2c_1 + 3c_t) - (c_2 + c_r) = 6q + 3 - 8q - 4 \le 0$ . If  $\ell$  is used by all three  $_{263}$  caterpillars, at least one of which has different parity than Q, then  $\#_1$  and  $\#_2$  grow by one and  $\#_3$  decreases by  $_{264}$  by one, implying that  $\omega$  grows by  $\Delta \omega = (c_1 + c_2 + 3c_t) - (c_3 + c_r) = 7q + 3 - 7q - 4 \le 0$ .

**Case 2c:** Two leaves  $\ell$  and  $\ell'$  of  $L_Q$  are used to embed a leaf of a caterpillar with different parity than Q. Then,  $\#_1$  grows by four and  $\#_2$  decreases by two, implying that  $\omega$  grows by  $\Delta \omega = (4c_1 + 3c_t) - (2c_2 + c_r) = 267$  $12q + 3 - 12q - 4 \le 0.$ 

**Case 3:** Exactly two caterpillars Q and Q' have their backbone embedded in  $N_r$  and **their parity is the same**. Let  $L_Q$  and  $L_{Q'}$  denote the sets of leaves in  $N_r$  that leaves in Q and Q', respectively, are embedded into. 270

**Case 3a:** No leaf of the third caterpillar is embedded in any leaf in  $L_Q \cup L_{Q'}$ . Then, the numbers  $\#_i$  do not change for any i, so  $\omega$  increases by  $\Delta \omega \leq 3c_t - c_r \leq 0$ 

**Case 3b:** Exactly one leaf of the third caterpillar is embedded in a leaf  $\ell$  in  $L_Q \cup L_{Q'}$ . If  $\ell \notin L_Q \cap L_{Q'}$ , 273 then  $\#_1$  grows by two and  $\#_2$  decreases by one, implying that  $\omega$  grows by  $\Delta \omega = (2c_1 + 3c_t) - (c_2 + c_r) =$  274  $6q + 3 - 8q - 4 \le 0$ . If  $\ell \in L_Q \cap L_{Q'}$ , then  $\#_1$  and  $\#_2$  grow by one and  $\#_3$  decreases by one, implying that 275  $\omega$  grows by  $\Delta \omega = (c_1 + c_2 + 3c_t) - (c_3 + c_r) = 7q + 3 - 7q - 4 \le 0$ .

**Case 4:** Exactly two caterpillars Q and Q' have their backbone embedded in  $N_r$  and **their parity is differ ent**. Let  $L_Q$  and  $L_{Q'}$  denote the sets of leaves in  $N_r$  that leaves in Q and Q', respectively, are embedded into. Further, let  $m := |L_Q \cap L_{Q'}|$ .

**Case 4a:** The embedding of the third caterpillar uses no leaf in  $L_Q \cup L_{Q'}$ . Then  $\#_1$  grows by 2m,  $\#_2$  decreases by m, and the token reservoir shrinks by 2qm - 3 tokens. Thus,  $\omega$  grows by  $\Delta \omega = (2mc_1 + 3c_t) - 281$  $(mc_2 + c_r + 2mqc_t) = (6mq + 3) - (4mq + 4q + 4 + 2mq) = -4q - 1 \le 0.$ 

**Case 4b:** The embedding of the third caterpillar uses a leaf of  $L_Q \cap L_{Q'}$ . Then,  $\#_1$  grows by 2m - 1,  $\#_2$  decreases by m - 2,  $\#_3$  decreases by one, and the token reservoir shrinks by 2qm - 3 tokens. Thus,  $\omega$  grows by  $\Delta \omega = ((2m-1)c_1+3c_t) - ((m-2)c_2+c_3+c_r+2mqc_t) = (6mq-3q+3) - (4mq-8q+3q+4q+4+2mq) = 285 - 2q - 1 \le 0.$ 

**Case 4c:** The embedding of the third caterpillar uses a leaf of  $L_Q \setminus L_{Q'}$ . If the third caterpillar has the same parity as Q, then this is identical to Case 4a. Otherwise,  $\#_1$  grows by 2(m+1),  $\#_2$  decreases by m+1, and the token reservoir shrinks by 2qm - 3 tokens. Thus,  $\omega$  grows by  $\Delta \omega = (2(m+1)c_1 + 3c_t) - ((m+1)c_2 + c_r + 2mqc_t) = (6mq + 6q + 3) - (4mq + 4q + 4q + 4 + 2mq) = -2q - 1 \le 0.$ 

**Case 4d:** The embedding of the third caterpillar uses a leaf of  $L_{Q'} \setminus L_Q$ . This case is identical to Case 4c.

**Case 5:** All three caterpillars have their backbone embedded in  $N_r$ . Let  $L_2$  be the set of leaves below r such that each leaf of  $L_2$  is used by the embeddings of exactly two caterpillars and these caterpillars have different parity. Let  $L_3$  be the set of leaves below r that are used in the embeddings of all three caterpillars, and observe that each such leaf causes us to remove 4q tokens from the reservoir. Further, abbreviate  $m_2 := |L_2|$  and  $m_3 := |L_3|$ . Then,  $\#_1$  grows by  $2m_2 + m_3$ ,  $\#_2$  grows by  $m_3 - m_2$ ,  $\#_3$  shrinks by  $m_3$  and the token reservoir shrinks by  $2qm_2 + 4qm_3 - 3$ . Thus,  $\omega$  grows by  $\Delta \omega = ((2m_2 + m_3)c_1 + m_3c_2 + 3c_t) - (m_2c_2 + m_3c_3 + c_r + 207)(2qm_2 + 4qm_3)c_t) = (6qm_2 + 7qm_3 + 3) - (6qm_2 + 7qm_3 + 4q + 4) \le 0$ .

**Claim 2.** For all configurations  $C_j$ , the token reservoir  $t_j$  is non-negative.

*Proof.* We show that, when the first withdrawal from the token reservoir happens, the number of tokens 300 accumulated in the reservoir exceeds the total number of tokens ever to be removed from it. To this end, 301

consider what happens when Transformation Rule 1 is applied to a lowest reticulation r in  $N^j$  for some iteration  $((N^j, \phi_j), t_j)$ . Recall that tokens are removed only if, for some caterpillars P and Q whose backbones are embedded below r and that have different parity below r, P and Q share some leaf  $\ell$  below r. In such a case, we call r bad with respect to (P, Q), and  $\ell$  is called r-bad with respect to (P, Q) (we omit the prefix if r is unknown). Note that no leaf is r-bad with respect to (P, Q) for more than one r, since, after applying Transformation Rule 1 to r, all r-bad leaves are "unzipped" and no longer shared by P and Q.

Now, fix P and Q, and consider only those leaves and reticulations that are bad with respect to (P, Q). In the following, we simply refer to such leaves and reticulations as "bad". Note that, since the embeddings of P and Q are subgraphs of the network N and the backbones of P and Q are embedded below each bad reticulation, the ancestor relation between bad reticulations is the same in the embedding of P as in the embedding of Q (otherwise, N contains a cycle). Since P and Q are caterpillars, there is a unique linear ordering  $r_0, r_1, \ldots, r_m$  of the bad reticulations such that  $r_{i+1}$  is an ancestor of  $r_i$  in both P and Q for all i.

In the following, we construct a common subsequence of P and Q containing at least half of all bad leaves, 314 which will imply the claim through use of Proposition 1. To this end, for each  $i_i$  let  $s_i$  denote the sequence of  $r_i$ -bad leaves in P. Recalling that the  $r_i$  occur on the backbone of Q in the same order as they do in P, it 316 suffices to show that a subsequence of Q can be obtained from  $s_i$  by removing at most one leaf and retaining 317 at least one leaf. To this end, we conduct a closer inspection of the configuration  $C_i = ((N^j, \phi_i), t_i)$  in which 318 Transformation Rule 1 is applied to  $r_i$ . Suppose that there are at least two  $r_i$ -bad leaves since, otherwise,  $s_i$ 319 is already a subsequence of Q. Let u denote the lowest node of the tree  $N_{r_i}^j$  that still has the backbones of 320 both P and Q embedded in it. Clearly, all  $r_i$ -bad leaves that are not below u form a suffix of  $s_i$  that is also a 321 subsequence of Q. By definition of u, it is not a leaf of  $N^j$  and u has children  $v_P$  and  $v_Q$  in  $N^j$  such that at 322 most one leaf  $\ell_Q$  below  $v_P$  is used in the embedding of Q and at most one leaf  $\ell_P$  below  $v_Q$  is used in the 323 embedding of P. But then, removing  $\ell_P$  from  $s_i$  yields a subsequence of Q. Note that the removal of  $\ell_P$  is 324 necessary to form a subsequence of Q since the sequences corresponding to P and Q may disagree on the 325 relative ordering of  $\ell_P$  and  $\ell_Q$ . 326

Now, the concatenation s of all subsequences of Q corresponding to the  $s_i$  is a common subsequence of P and Q and it contains at least half of all bad leaves (recall that we only remove a leaf from  $s_i$  if it contains at least two leaves). Then, by Proposition 1, the number of bad leaves is at most  $2 \cdot 2^i = 2 \cdot 2^{\log_3 n} = 2 \cdot n^{\log_3 2}$ , where  $n = 3^i$  is the number of leaves in N. Summing over all three caterpillar pairs, we get an upper bound of  $6n^{\log_3 2}$  bad leaves overall.

Next, we show that the first token retraction is preceeded by the creation of enough tokens to compensate 332 for all retractions. To this end, consider the first configuration  $C_i = ((N^j, \phi_i), t_i)$  such that applying Transformation Rule 1 to a reticulation r in  $N^{j}$  incurs a withdrawal from the token reservoir. In particular, this implies 334 the existance of two different-parity caterpillars P and Q such that their backbone is embedded below r and 335 both their embeddings use a common leaf  $\ell$  below r. However, by construction of P and Q, none of the labels 336 occuring in the lowest third of P occurs in the lowest third of Q and, thus,  $\ell$  is preceded by at least n/3 leaves 337 in either P or Q; without loss of generality, suppose Q. Since the backbone of Q is embedded below r, there 338 are at least n/3 leaves below r used by the embedding of Q, but not that of P. However, since all leaves in N 339 are used by all three caterpillars, all these n/3 leaves were "unzipped" in previous operations. 340

Since  $C_i$  is the first configuration in which P and Q have different parity and "share" a leaf, we know that 341 in all previous "unzip" operations, either (a) P and Q have the same parity or (b) the embeddings of P and Q share no leaves below the corresponding reticulation or (c) the embedding of at least one of P and Q uses 343 only one leaf below the corresponding reticulation. Clearly, in cases (a) and (b), the unzip operation does not 344 "separate P from Q" in any leaf, that is, the unzip operation does not reduce the number of leaves used by the 345 embeddings of both P and Q. In case (c), each unzip operation can "separate P from Q" in at most one leaf, 346 implying that  $C_i$  is preceded by at least n/3 unzip operations, each creating 3 tokens in the reservoir. Thus, 347 by the time the first withdrawal is made from the reservoir, it contains at least n tokens which is sufficient 348 to cover all withdrawals (recall that we withdraw 2q tokens for each of the at most  $6n^{\log_3 2}$  bad leaves) since 349  $n > 12qn^{\log_3 2}$  by (1).

With Claim 1 and Claim 2 we can now prove the bound on the number of leaves  $\#^T$  in T in the number n of leaves in N and the number k of reticulations in N. To this end, note and recall that (a)  $t_0 = 0$ , (b) all leaves in N are used by the embeddings of all three caterpillars, and (c) T has no reticulations. Then,

$$c_{1} \cdot \#^{T} = c_{1} \sum_{i \in \{1,2,3\}} \#^{T}_{i} \leq \sum_{i \in \{1,2,3,r\}} c_{i} \cdot \#^{T}_{i} \stackrel{Claim \ 2}{\leq} \sum_{i \in \{1,2,3,r\}} c_{i} \cdot \#^{T}_{i} + c_{t} \cdot t_{\Omega} \stackrel{(2)}{=} \omega(C_{\Omega})$$

$$\overset{Claim \ 1}{\leq} \omega(C_{0}) \stackrel{(2)}{=} \sum_{i \in \{1,2,3,r\}} c_{i} \cdot \#^{N}_{i} + c_{t} \cdot t_{0} = c_{3} \cdot \#^{N}_{3} + c_{r} \cdot \#^{N}_{r} = 3qn + 4(q+1)k$$

$$= c_{1}(n + 4(q+1)k/3q).$$

Thus, T has at most  $n + \frac{4(q+1)k}{3q}$  leaves.

Lemma 3 tells us that if we can construct a network with few reticulations that displays our caterpillars, then we can also construct a MUL-tree with few leaves that displays our caterpillars. Since Corollary 1 says that such MUL-trees do not exist, we conclude that such networks do not exist. 353

**Theorem 1.** Let  $\epsilon > 0$ . Then, there are three caterpillar trees, each with  $n \in \mathbb{N}$  leaves, such that any network displaying all three caterpillars has at least  $(3/2 - \epsilon)n$  reticulations. 356

*Proof.* Let  $\delta := 2\epsilon/3$  and choose q large enough so that  $\beta := \delta - 1/q+1 > 0$ . Finally, choose i such that  $\beta \ge (1 - 1/q+1)(2/3)^i$ . Let N be a network displaying  $X_i$ ,  $Y_i$  and  $Z_i$ , that is, there is a caterpillar embedding  $(N, \phi)$ . Let  $(T, \phi)$  be the result of applying Transformation Rule 1 exhausively to  $(N, \phi)$  and note that, by Observation 1,  $(T, \phi)$  is a caterpillar embedding, that is, T displays  $X_i$ ,  $Y_i$ , and  $Z_i$ . Further, by Lemma 2, T has at least  $3^{i+1} - 2^{i+1}$  leaves. Let k denote the number of reticulations in N. Then,

$$3^{i+1} - 2^{i+1} \stackrel{Lemma \ 2}{\leq} \#_T \stackrel{Lemma \ 3}{\leq} n + \frac{4(q+1)k}{3q} = 3^i + \frac{4(q+1)}{3q} \cdot k$$

and, thus,

$$k \ge (2 \cdot 3^{i} - 2^{i+1}) \cdot 3^{q/4(q+1)} = 3/2 \cdot q/q + 1 \cdot (n - 2^{i})$$
  
=  $3/2(1 - 1/q + 1)(n - 2^{i})$   
 $\ge 3/2((1 - 1/q + 1)n - \beta n)$   
=  $3/2(1 - \delta)n = (3/2 - \epsilon)n.$ 

#### 5 Discussion and Open Problems

We have shown that, for each  $\epsilon > 0$  and  $t \le 3$ , there is some  $n \in \mathbb{N}$ , and t caterpillars with n leaves, such that any MUL-tree displaying the caterpillars has at least  $(t-\epsilon)n$  leaves. Whether this result can be generalized to  $t \ge 4$  remains an interesting open question, even more so for networks, where the question would be to generalize our lower bound of  $(3/2 - \epsilon)n$  reticulations required to display three caterpillars to  $t \ge 4$  caterpillars. 360

Note that Theorem 1 can be stated more precisely as "there is some function  $\alpha(n) \in o(1)$  such that, for each n, there are three caterpillars with n leaves that cannot be displayed by any network with fewer than  $(^3/_2 - \alpha(n))n$  reticulations". This raises the question how the "smallest" functions  $\alpha$  for which this statement still holds, may look like (other than being in o(1)). In the MUL-tree case, a closer inspection of our proofs gives  $\alpha(n) \leq 2 \cdot (^2/_3)^{\log_3 n}$ . If we force the MUL-tree displaying the t trees to be a caterpillar, the question is equivalent to the question of the shortest supersequence that any collection of t permutations over  $\{1, \ldots, n\}$  may have. This can be shown [Hun23] to be at least  $(t - O(n^{-1/2}))n$ , implying  $\alpha(n) \in O(n^{-1/2}) \subset o((^2/_3)^{\log_3 n})$ . If there is always a MUL-caterpillar among the optimal MUL-trees displaying the t caterpillars, then this would imply a stricter bound for both MUL-trees and networks than what we showed here. Apart from improving

350

351

the function  $\alpha(n)$ , it is interesting whether the factor of 3/2 can be improved. In other words, is there a family 371 of triples of phylogenetic trees for which more than 3n/2 reticulations are needed? We do know that all triples 372 of phylogenetic trees can have a network with 2(n-2) reticulations that display them: the network can be 373 obtained from a tree on two leaves by inserting each remaining leaf using two reticulations. However, the 374 same three trees might also be displayed by a network with strictly fewer reticulations. The best possible 375 bound for three trees is therefore between  $(3/2 - \epsilon)n$  and 2(n-2). More generally, the best possible bound 376 for  $t \ge 3$  trees is between  $(3/2 - \epsilon)n$  and (t - 1)(n - 2). Can this gap be closed or narrowed? As our results 377 are for caterpillars, we expect that trees of varying topology can be used to prove such tighter bounds. 378

The last two questions beg the, somewhat philosophical question of whether reticulations are strictly more powerful than multiple leaves? For MUL-trees, we know that we cannot do much better than the trivial upper bound of  $t \cdot n$ . Is the same true for networks, can we not do much better than the trivial upper bound of (t-1)(n-2), or are networks really more powerful than MUL-trees in this sense? 382

Finally, our original motivation for considering this problem came from the Hybridization Number problem. However, the bounds proven in this paper do not have direct formal consequences for that problem. Hence, another interesting direction for future research is to see if our bounds can be used to prove a negative result regarding exact algorithms for Hybridization Number, e.g. parameterized by treewidth (see [IJW22]). 380 380 380 380 380 380 380 380 380

### Acknowledgements

We thank Simone Linz, Guillaume Scholz, and Stefan Grünewald for their thorough review work and helpful comments. We also thank Zach Hunter for explaining the bound on supersequences of permutations.

## **Fundings**

Leo van Iersel is funded in part by Netherlands Organization for Scientific Research (NWO) grants OCENW.KLEIN•125 and OCENW.GROOT.2019.015. Mark Jones is funded by Netherlands Organization for Scientific Research (NWO) grant OCENW.KLEIN.125.

## Conflict of interest disclosure

The authors declare that they comply with the PCI rule of having no financial conflicts of interest in relation <sup>395</sup> to the content of the article. All authors are also recommenders of PCI Mathematical & Computational Biology. <sup>396</sup>

#### References

[BS07]	Bordewich M and Semple C. Computing the minimum number of hybridization events for a consistent evolutionary history. <i>Discrete Applied Mathematics</i> 155 (2007), 914–928.	398 399
[BS18]	Bordewich M and Semple C. A universal tree-based network with the minimum number of retic- ulations. <i>Discrete Applied Mathematics</i> 250 (2018), 357–362. issn: 0166-218X. https://doi.org/10. 1016/j.dam.2018.05.010.	400 401 402
[BSS05]	Baroni M, Semple C, and Steel M. A framework for representing reticulate evolution. <i>Annals of Combinatorics</i> 8 (2005), 391–408. https://doi.org/10.1007/s00026-004-0228-0.	403 404
[Bar+05]	Baroni M, Grünewald S, Moulton V, and Semple C. Bounding the number of hybridisation events for a consistent evolutionary history. <i>Journal of mathematical biology</i> 51 (2005), 171–182.	405 406
[DARM08]	Dagan T, Artzy-Randrup Y, and Martin W. Modular networks and cumulative impact of lateral transfer in prokaryote genome evolution. <i>Proceedings of the National Academy of Sciences</i> 105 (2008), 10039–10044.	407 408 409

390

387

394

[FS15]	Francis AR and Steel M. Which Phylogenetic Networks are Merely Trees with Additional Arcs? <i>Systematic Biology</i> 64 (June 2015), 768–777. issn: 1063-5157. https://doi.org/10.1093/sysbio/	410 411
	syv037.	412
[Hay16]	Hayamizu M. On the existence of infinitely many universal tree-based networks. <i>Journal of Theoretical Biology</i> 396 (2016), 204–206. issn: 0022-5193. https://doi.org/10.1016/j.jtbi.2016.02.023.	413 414
[Hun23]	Hunter Z. <i>Shortest Common Supersequence of Permutations – TCS StackExchange</i> . 2023. url: https://cstheory.stackexchange.com/questions/53153 (visited on 08/06/2023).	415 416
[IJW22]	Iersel L van, Jones M, and Weller M. Embedding Phylogenetic Trees in Networks of Low Treewidth. In: <i>30th Annual European Symposium on Algorithms (ESA 2022)</i> . Vol. 244. Leibniz International Pro- ceedings in Informatics (LIPIcs). 2022, 69:1–69:14. isbn: 978-3-95977-247-1. https://doi.org/10. 4230/LIPIcs.ESA.2022.69.	417 418 419 420
[IK11]	lersel L van and Kelk S. When two trees go to war. <i>Journal of theoretical biology</i> 269 (2011), 245–255.	421 422
[Kel+12]	Kelk S, Iersel L van, Lekic N, Linz S, Scornavacca C, and Stougie L. Cycle killer qu'est-ce que c'est? On the comparative approximability of hybridization number and directed feedback vertex set. <i>SIAM journal on discrete mathematics</i> 26 (2012), 1635–1656.	423 424 425
[Mal05]	Mallet J. Hybridization as an invasion of the genome. <i>Trends in ecology &amp; evolution</i> 20 (2005), 229–237.	426 427
[Zha16]	Zhang L. On tree-based phylogenetic networks. <i>Journal of Computational Biology</i> 23 (2016), 553–565.	428 429