

# TGIF: Topological Gap In-Fill for Vascular Networks

## A Generative Physiological Modeling Approach

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**Abstract.** This paper describes a new approach for the reconstruction of complete 3-D arterial trees from partially incomplete image data. We utilize a physiologically motivated simulation framework to iteratively generate artificial, yet physiologically meaningful, vasculatures for the correction of vascular connectivity. The generative approach is guided by a simplified angiogenesis model, while at the same time topological and morphological evidence extracted from the image data is considered to form functionally adequate tree models. We evaluate the effectiveness of our method on four synthetic datasets using different metrics to assess topological and functional differences. Our experiments show that the proposed generative approach is superior to state-of-the-art approaches that only consider topology for vessel reconstruction and performs consistently well across different problem sizes and topologies.

**Keywords:** vascular reconstruction, vascular connectivity, angiogenesis.

## 1 Introduction

Vascular image analysis plays a crucial role in many research areas and has various applications. In neuroscience, for instance, gaining further insight into the topological and morphological properties of the cortical microvasculature is fundamental to a better understanding of the neurovascular coupling and regulation in neurodegenerative diseases such as Alzheimer's [1]. Advances in high-resolution imaging modalities, e.g., synchrotron radiation micro-CT, have greatly facilitated the acquisition of highly detailed image data of complex vascular systems across different scales down to the capillary level [8]. However, the analysis and reconstruction of complete vascular networks is still an open problem despite extensive ongoing research in vascular image analysis. Most of the research addresses the problem of *vessel detection* and *segmentation* [4] while mostly neglecting *vascular connectivity* [9]. Incomplete connectivity dramatically complicates or even precludes the analysis of hemodynamic and metabolic properties [1, 8]. Topological defects are commonly caused by limitations of the reconstruction methodology (e.g. invalid model assumptions) or problems related to image acquisition (image noise, artifacts) and sample preparation (insufficient dye filling, air bubbles, clotting).

There is only very little previous work on reconstruction and correction of vessel connectivity [4, 9]. A popular principle is the extraction of a minimum spanning tree (MST) w.r.t. different distance metrics based on Euclidean distance or physiological principles [2, 4]. Risser et al. [10] proposed a gap filling approach based on tensor voting. The

authors use local directional and morphological information to define a tensor field supporting likely vessel pathways to be tracked to fill gaps. More recently, Kaufhold et al. [3] presented a learning-based algorithm to recover missing connectivity by threshold relaxation relying on last traces of information in the image data that have been missed before. Though in case of total signal loss at a gap (e.g. caused by incomplete filling with dye), their method is no longer applicable.

The slightly different and more fundamental problem of synthetic formation of functionally adequate vascular networks has attracted considerable attention in the field of computational physiology [5, 12, 13]. Here, physiological principles are applied to generate synthetic vascular models that satisfy prescribed physiological conditions.

In this work, we combine different key ideas of both aforementioned problem classes to reconstruct consistent 3-D arterial trees from partially incomplete image data. We utilize artificially generated, yet physiologically plausible, vascular structures for the correction of vascular connectivity. To this end, we adapt a simulation framework based on a simplified angiogenesis model to iteratively complete the defective vasculature taking into account the topological and morphological evidence extracted from the image data. The primary objective of this work is first and foremost the reconstruction of a physiologically meaningful vasculature rather than the recovery of the exact topology.

## 2 Methods

**Generating New Vascular Structures.** The proposed approach for topological gap in-fill (TGIF) is primarily based on our generative model for the construction of synthetic 3-D arterial tree models as introduced in [12]: The physiologically motivated method combines a simplified model for sprouting angiogenesis stimulated by a growth factor with structural remodeling and degeneration in response to morphological and metabolic stimuli to iteratively generate an arterial tree model until the (prescribed) metabolic demand of the embedding tissue is met, i.e., no hypoxic regions left. The vascular system is modeled as a set of binary trees where each vessel branch is approximated by a sequence of edges (vessel segments) each representing a cylindrical tube of fixed length and radius. Additional morphological constraints are enforced at construction time with regard to the morphometrical configuration at bifurcations, i.e., relationship of radii (Murray's law) and optimal bifurcation angles derived from Murray's minimum work principle (MWP). The iterative growth proceeds in four steps [12]:

- (S1) Angiogenesis model: reaction-diffusion process to estimate tissue perfusion and oxygenation as well as the corresponding concentration map of vascular endothelial growth factors (VEGFs) emitted by hypoxic-ischemic cells.
- (S2) Vascular growth forming new sprouts in chemotactic response to VEGF.
- (S3) Structural remodeling and degeneration of redundant vascular structures.
- (S4) Repeat steps (S1)–(S3) until prescribed tissue metabolism is met.

To apply this concept [12], we identify feeding vessels (root nodes) entering the image volume and isolated vascular components with no connection to any root. As these unrooted parts are lacking blood supply (stagnant hypoxia), they do not contribute to the perfusion of the surrounding tissue estimated in step (S1). The resulting hypoxic areas lead to locally increased VEGF secretion attracting new sprouts. Thus, step (S2) iteratively grows new sprouts originating from rooted (perfused) segments following the local VEGF gradient (constant step length  $\delta$ ), and the sprouting tip is connected to an

unrooted segment when it gets closer than 4  $\mu\text{m}$ . Still, due to the sparse vascularization of 3-D tissues ( $\approx 2\%$  vascular volume fraction [8]), a sprouting tip would rarely hit another unrooted vessel if purely driven by angiogenesis [13]. Thus, we propose to amend the generative model by an “attractor” to enforce such connections between vessel segments if supported by the topology and morphology, which we discuss next.

**Connecting Unrooted Segments.** We allow rooted sprouts to link with nearby unrooted vascular structures in step (S2). Similar to [13], we define the set of potential connection candidates at a sprouting tip as unrooted vessel segments lying within a spherical-cone-shaped sector along the sprout direction (maximum distance  $R_{\max} = 100 \mu\text{m}$ , angular deviation  $\theta_{\max} = 60^\circ$  [13, Tbl. 1]). To find the optimal connection candidate for a sprouting node  $v_s$  with proximate connection candidates  $v_c \in \mathcal{V}_c$ , we first rank the connecting edges  $e = (v_s, v_c)$  w.r.t. the heuristic cost function (HCF) proposed by Reichold [8] considering directional and spatial information. This parametric model favors short over long infills, but also a minimal bending angle of the new connection:

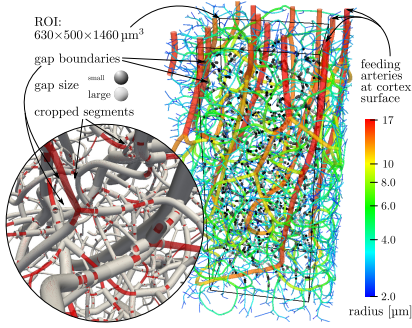
$$C_{\text{HCF}}(e) = \lambda_d \cdot \frac{l(e)}{d_{\max}} + (1 - \lambda_d) \cdot \left[ 1 - \exp\left(-\frac{\alpha}{\alpha_0}\right) \right] \left[ 1 - \text{erf}\left(\frac{l(e)}{d_0}\right) \right], \quad (1)$$

where  $l(e) \leq d_{\max}$  denotes the edge length, and  $\lambda_d \in [0, 1]$  is a weighting factor to balance the distance-based and angle-based term. The bending angle  $\alpha$  is defined as the sum of the bending angles relative to the vessel direction at  $v_s$  and  $v_c$ , respectively (see supplementary material for illustrations of  $C_{\text{HCF}}$ ). The parameter  $d_0$  defines the maximum distance beyond which the influence of  $\alpha$  is negligible. The factor  $\alpha_0$  is used to shape the angle-dependent term. As the ranking of the candidate edges w.r.t.  $C_{\text{HCF}}$  is not symmetric, i.e., the top-ranked edge  $(v_s, v_t)$  of  $v_s$  can be ranked very low at  $v_t$ , we apply a k-reciprocal neighborhood search [7]: From the top  $k_{\text{rNN}}$  ranked candidate edges of  $v_s$ , we pick the highest ranked edge  $\hat{e} = (v_s, \hat{v}_t)$  for which  $\check{e} = (\hat{v}_t, v_s)$  is among the  $k_{\text{rNN}}$  highest ranked edges of  $\hat{v}_t$ . If there is no such “ $k_{\text{rNN}}$ -optimal” edge  $\hat{e}$ , we follow the homing mechanism proposed by Secomb et al. [13, Eq. 10–14] that makes sprouting tips susceptible to nearby vessel segments, i.e., the growth direction of the evolving sprout is computed as the weighted sum over the set of potential connection candidates as defined above (sensitivity  $k_V = 10$  [13, Tbl. 1]). Finally, if the sprouting tip does not sense any proximate vessel segments, i.e., empty set of connection candidates, we fall back to the VEGF-based strategy in step (S2) to determine the growth direction.

**Remodeling.** Adding new vascular connections modifies the network topology and – under consideration of global geometrical constraints, such as continuity of radii along connections and radii at bifurcations following Murray’s law – we have to update the vessel diameters in step (S3) to match the new conditions. To this end, we adjust the diameters of newly generated vessels (*synthetic* nodes) and maintain estimated vessel radii at *static* nodes with strong evidence in the image data.

Every synthetic subtree generated at an infill location has a well-defined static root node  $n_r$  at the proximal transition from static to synthetic data. Also, each terminal node at the distal end of the subtree is either (1) non-static, i.e., newly generated sprouting tip not yet linked with a connection candidate, or (2) static, i.e., transition from synthetic to static data with prescribed radius. We iteratively remodel the subtree as follows:

- (R1) Prescribe vessel radius at static terminal nodes according to image evidence.
- (R2) Propagate radii from all subtree terminals to root  $n_r$  according to Murray’s law [12].



**Fig. 1.** Synthetic vascular tree model with artificial gaps ( $\mathcal{B}_2$ ,  $\gamma_{\text{crop}} = 32\%$ ).

**Table 1.** Examined parameter ranges for different reconstruction approaches. Default HCF parameters used for TGIF are underlined.

Method	Parameter	Value/Range	Text/Reference
MWP	$\lambda_{\text{MWP}}$	{0.0, 0.1, 0.2, 0.3, 0.4, 0.5}	[2, Eq. 13]
TV	$L$	see $d_{\text{max}}$	[10, Eq. 6]
	$\theta$	{15°, 30°, 45°, 60°}	[10, Eq. 6]
	$l_{\text{max}}$	{0.5, 1.0, 1.5, 2.0} $\times s_{\text{gap}}$	$\mathcal{B}_1$ [10, Sec. III.K.4]
		{12, 48, 96, 144} [ $\mu\text{m}$ ]	$\mathcal{B}_2$
	$s_{\text{min}}$	{ $10^{-4}$ , $10^{-2}$ , $10^{-1}$ , 0.5}	[10, Sec. III.K.4]
HCF	$d_{\text{max}}$	{0.5, 0.75, 1.0, 1.5, 2.0} $\times s_{\text{gap}}$ [ $\mathcal{B}_1$ ] Eq. (1)	
		{12, 24, 48, 96} [ $\mu\text{m}$ ]	$\mathcal{B}_2$
	$\lambda_d$	{ <u>0.4</u> , 0.5, 0.6, 0.7}	
	$\alpha_0$	{ <u>100°</u> , 120°, 140°, 160°} / $\log(2)$	
	$d_0$	{0.5, 1.0, 2.0, <u>4.0</u> } $\times d_{\text{max}}$	
	$k_{\text{rNN}}$	{ <u>1</u> , 2, 4, 6}	
TGIF	$\lambda_g$	{1, 2, 4, 8}	[12, Eq. 12]
	$\delta$	{8, 16, 32, 64} [ $\mu\text{m}$ ]	
	$\xi$	{1.00, 1.05, 1.10, 1.15}	(R3)

(R3) STOP if for updated and static vessel radius at  $n_r$ :  $\frac{T_{\text{updated}}}{T_{\text{static}}} < \xi$  for constant  $\xi \geq 1$ .

(R4) Randomly choose a non-static terminal node  $n_t$  of the subtree (FAIL if none is left), shrink the terminal vessel branch at  $n_t$  ( $r' = 0.9r$ ), and prune it if  $r' \leq 2 \mu\text{m}$ .

(R5) Repeat steps (R1)–(R4) until STOP or FAIL.

(R6) Reinforce static radius at  $n_r$  and propagate radii down to subtree terminals [12].

If the tree cannot be remodeled properly (FAIL), we discard the new connection edge and continue with steps (S1)–(S4). Note that we prevent static vessel branches from undergoing any kind of remodeling, degeneration or cropping, except for the adjustment of vessel radii at short isolated branches following the concept of “islands” in [10]. The simulation is ultimately stopped if the surrounding tissue is sufficiently perfused [12].

### 3 Experiments and Results

We evaluate the performance of the proposed algorithm for TGIF on four synthetic datasets and compare it to different baseline approaches.

**Experimental Data.** Following the experimental setup of Schneider et al. [12, Fig. 4–6], two artificial arterial tree models are generated for different random seeds. For each of these models we derive a vessel size imaging (VSI) map that is in turn used to synthesize a functionally equivalent but structurally different arterial tree following [5]. The vessel radii range from (2–17)  $\mu\text{m}$  in all four cases. We then create artificial gaps by introducing simulated air bubbles (occlusions) at randomly sampled locations within the vessel lumen and cropping the occluded parts of the vasculature. We examine two different configurations for the choice of the size and number of artifacts: In a first set of experiments ( $\mathcal{B}_1$ ), we vary the bubble size  $s_{\text{gap}} \in \{3, \dots, 384\} [\mu\text{m}]$ , while the number of artifacts is chosen such that the cumulative length of the cropped vessel segments amounts to  $\gamma_{\text{crop}} = 5\%$  of the total segment length of the ground truth (GT) vasculature. Conversely, in the second configuration ( $\mathcal{B}_2$ ), we consider different crop factors  $\gamma_{\text{crop}} \in \{1, \dots, 32\} [\%]$  for a fixed ensemble of bubble artifacts ( $s_{\text{gap}} = (3, 6, 12, 24, 48, 96) [\mu\text{m}]$ ). In order to avoid boundary artifacts, we only crop vessel segments within a centered box-shaped region of interest (ROI) (see Figure 1).

Likewise, we exclude large vessels ( $r > 10 \mu\text{m}$ ) that are rarely affected by incomplete filling in real data. From each of the deteriorated vasculatures, we derive an observable counterpart to be used in our infill experiments by modeling the forward imaging and inverse segmentation process as follows: First, we synthesize 3-D image data by sequentially discretizing each vessel segment via a Gaussian point spread function and additive Gaussian noise (isotropic spatial resolution:  $1 \mu\text{m}$ ). Next, we apply vessel segmentation [11] and skeletonization based on distance-ordered homotopic thinning (DOHT) to reconstruct the vascular structure. Vessel calibers along the skeletonized centerline are determined based on distance transformation. Finally, we apply spline smoothing to each vessel branch and assign a constant radius that preserves the total lumen of the original branch to make the estimation of vessel direction and diameter more robust.

**Alternative Methods.** On these data we test four alternative gap filling approaches: (1) Prim’s algorithm was applied to extract the *minimum spanning tree (MST)* with each candidate edge connecting a rooted with an unrooted node ( $e = (v_r, v_c)$ ) weighted by the edge length  $l(e)$ . The radius of candidate edges added to the tree is chosen as the arithmetic mean of the maximum adjacent segment radius at  $v_r$  and  $v_c$ . (2) Jiang et al. [2] proposed a different weighting scheme based on Murray’s *minimum work principle (MWP)*. The authors derive a simplified cost function for each candidate edge as a mixture of the shortest path tree (SPT) and  $r^2$ -weighted MST [2, Eq. 13]:

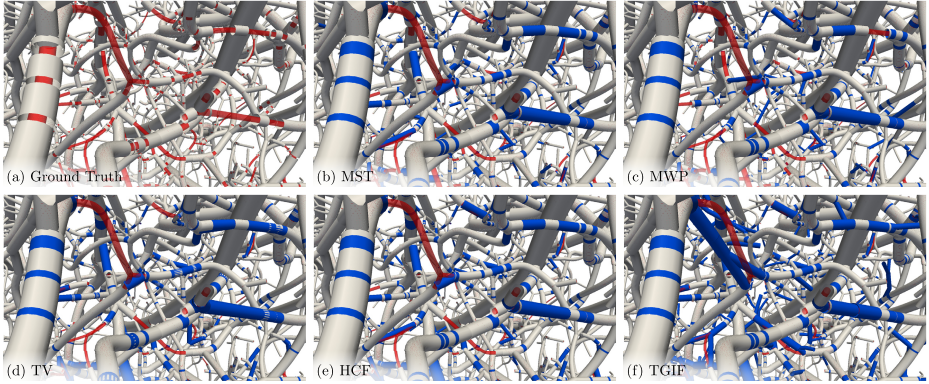
$$C_{\text{MWP}}(e = (v_r, v_c)) = \underbrace{l(e) \cdot r^2(e)}_{r^2\text{-MST}} + \underbrace{\lambda_{\text{MWP}} \frac{r_{\min}^3}{2} \cdot \sum_{e_i \in \mathcal{P}(v_c, \hat{v}_r)} l(e_i)/r(e_i)}_{\text{simplified SPT}}, \quad (2)$$

where  $r_{\min} = 2 \mu\text{m}$  denotes the minimum vessel radius and  $\mathcal{P}(v_c, \hat{v}_r)$  the shortest path from candidate node  $v_c$  to the closest root node  $\hat{v}_r$  of  $v_r$ . The radius of candidate edges is chosen as in MST. Following [2], we apply Prim’s algorithm for different values of  $\lambda_{\text{MWP}} < 1$  (see Table 1) to find the optimal tree minimizing the MWP cost.

(3) The gap filling approach by Risser et al. [10] based on *tensor voting (TV)* was applied as follows: First, we compute the tensor and saliency maps from the vessel network using an isotropic spatial resolution of  $1.5 \mu\text{m}$ . For the generation of missing segments we found that the tracking scheme proposed by the authors [10, Eq. 8], i.e., greedily following the maximum saliency in a local neighborhood, can easily fail due to the limited spatial resolution. Thus, we adapt a fiber tracking scheme also used for diffusion tensor imaging to directly track the tensor field rather than the derived saliency measure [6] and use it as a fallback strategy. We perform an exhaustive grid search for each dataset to find the optimal configuration of the four main parameters defining the tensor field and the break criteria during tracking (see Table 1).

(4) Finally, we use the proposed combination of the heuristic cost function in Equation (1) and the k-reciprocal neighborhood scheme in a modified MST-like framework, which can be regarded as a simplified version of TGIF considering topological information only: Unlike standard MST, we consider candidate edges from rooted to unrooted nodes only if they fulfill the k-reciprocal neighborhood condition. We then apply Prim’s algorithm to greedily extract the corresponding MST w.r.t. candidate edge weights as defined in Equation (1) for different parameters (see Table 1).

**Our Approach: relax TGIF.** Similar to the baseline experiments, we perform a parameter sweep for selected parameters listed in Table 1. For each parameter set and

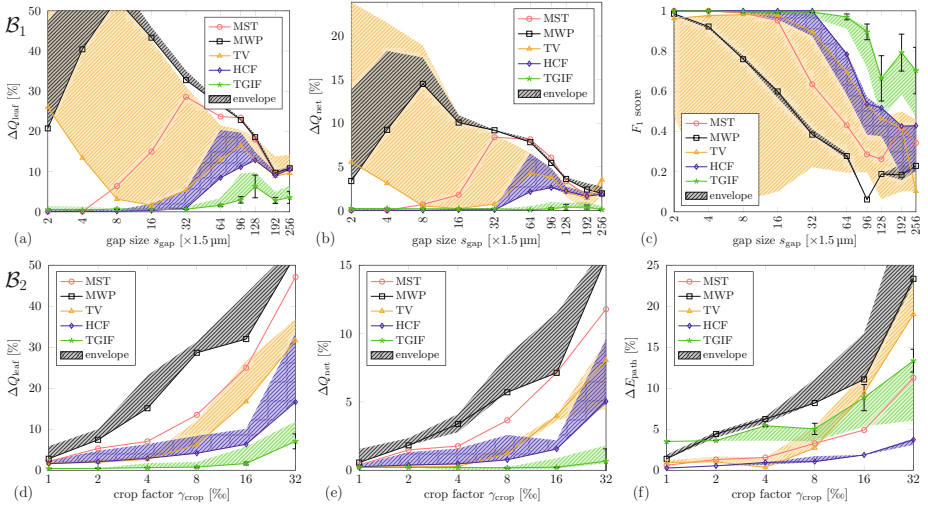


**Fig. 2.** Close-up views of (a) the ground truth (GT) network in Figure 1 and (b–f) the reconstructed networks generated by different approaches for optimal parameter values minimizing  $\Delta Q_{\text{leaf}}$ . Vessel branches are rendered as cylindrical tubes for the cropped synthetic GT network (gray), cropped gaps (red, radius scaled for better visibility), and reconstructed infill edges (blue).

each input network we run four TGIF simulations using different seeds to account for statistical variations of the reconstructed topologies. As above, the prescribed metabolic profile and angiogenesis parameters are taken from [12].

**Quantitative Analysis and Comparison.** For the comparison of the different reconstruction approaches we employ four different metrics focusing on different aspects: (1) *Topological correctness* is assessed by a precision-recall analysis. Each infill edge connecting to a static node is counted as true positive if the static target and the proximal interface to the static graph originate from the same bubble artifact (false positive otherwise). False negatives are computed as  $(\# \text{gap boundary nodes} - \# \text{true positives} - \# \text{gaps})$ . From the derived precision and recall measures, we compute the  $F_1$  score. (2) As direct comparison of the network topology may not be adequate to capture functional differences, we consider differences in the *path length* from GT nodes  $v_{\text{GT}}$  to the shortest GT root node  $\hat{v}_{\text{GT}}$  [2]:  $\Delta E_{\text{path}} = \frac{1}{\|\mathcal{V}_{\text{GT}}\|} \sum_{v_{\text{GT}} \in \mathcal{V}_{\text{GT}}} |l(P(v', \hat{v}')) - l(P(v_{\text{GT}}, \hat{v}_{\text{GT}}))|$ , where  $v'$  is the node of the reconstructed graph that is closest to  $v_{\text{GT}}$  in an Euclidean sense. As each synthetic arterial tree contains 16 roots in total, the paths may well end up at different roots ( $\hat{v}' \neq \hat{v}_{\text{GT}}$ ). The path length at unrooted nodes is set to zero. (3,4) Finally, we compare the *hemodynamic properties* of the reconstructed and GT network. To this end, we simulate the blood flow in the network using pressure boundary conditions at root and leaf nodes where the caliber-dependent pressure values are taken from literature [8]. Terminal nodes of the reconnected networks without any matching leaf node in the GT network are assigned a no-flow boundary condition. From the simulation results, we compute the relative net inflow change at all GT leaf nodes:  $\Delta Q_{\text{leaf}} = \sum_{v_{\text{GT}}} \frac{|Q_{\text{in}}(v') - Q_{\text{in}}(v_{\text{GT}})|}{Q_{\text{in}}(v_{\text{GT}})}$ . Similarly, we consider the total flow entering the reconstructed and GT network and compare their absolute relative difference  $\Delta Q_{\text{net}}$ .

Close-up views of the reconstructed networks from the different approaches are provided in Figure 2. Topological differences are visually most pronounced for long curved segments that are difficult to recover. The examples illustrate that TGIF is able to correctly fill most of the gaps and, unlike the other approaches, form bifurcations when necessary. Quantitative results w.r.t. the different metrics are shown in Figure 3 for the



**Fig. 3.** Quantitative evaluation of different reconstruction approaches for varying gap size (a–c) and cumulative artifact length (d–f). (a,d) Relative net inflow change  $\Delta Q_{\text{leaf}}$  at leaf nodes. (b,e) Relative total net inflow change  $\Delta Q_{\text{net}}$ . (c) Topological correctness of infill edges w.r.t.  $F_1$  score. (f) Relative path length error. For each problem size (fixed  $s_{\text{gap}}$  and  $\gamma_{\text{crop}}$ ), average values over all four synthetic datasets are marked for each reconstruction approach using the optimal parameter configuration (see Table 1) minimizing  $\Delta Q_{\text{leaf}}$ . Shaded areas within each subfigure outline the envelope of the average values obtained by applying these optimal parameters across  $s_{\text{gap}}$  and  $\gamma_{\text{crop}}$ , respectively. Error bars for TGI<sup>F</sup> indicate the standard deviation across the four datasets.

different reconstruction approaches and both experimental setups ( $B_1$ ,  $B_2$ ). In general, the different metrics indicate a decreasing quality of the reconstructed networks for increasing problem size. For increasing gap size ( $B_1$ ), the topological differences of the GT and reconstructed models become more pronounced as indicated by the decreasing  $F_1$  score. It is remarkable that small changes in topology can have a substantial impact on flow-related errors (see Figures 3a–3c). The figures indicate that TGI<sup>F</sup> outperforms the baseline approaches. Only for small crop factors, TGI<sup>F</sup> shows an increased path length error while the flow-related metrics remain consistently small (see Figures 3d–3f). This demonstrates the difficulty of reasonable comparison and validation of vascular networks. The parameter study reveals that, while TV proves to be highly sensitive to the parameter choice, TGI<sup>F</sup> performs consistently well across different problem sizes and topologies (functionally equivalent but structurally different networks, see above), as indicated by the narrow envelopes and small error bars in Figure 2, respectively. Finally, we find that MST outperforms MWP in our experiments. However, further experiments (not reported here) reveal that, particularly for the synthetic VSI-based networks, Murray’s minimum work principle seems to hold at the macroscale only, i.e., excluding the microvasculature ( $r \leq 5 \mu\text{m}$ ) significantly improves the performance of MWP over MST as reported in [2] for the macrostructure of murine coronary arterial trees.

## 4 Conclusions and Future Work

We proposed a new method for the reconstruction of complete 3-D arterial trees from partially incomplete image data by combining different key ideas that have been used

for the artificial generation of functionally adequate vascular models and different concepts aiming at the correction of vascular connectivity. The experiments demonstrate the effectiveness and superior performance of our approach compared to different state-of-the-art methods for vascular reconstruction. In this work, we have only considered arterial tree models rather than full-fledged vascular networks. Jointly considering arterial and venous networks raises additional problems as recently described in [13]. For instance, vascular remodeling and reconnection have to be reconsidered carefully as the hemodynamics can easily be spoiled by inserting functional shunts. Likewise, it would be interesting to exploit the image data not only for the initial estimation of the (defective) vascular topology but also to further guide and constrain the reconstruction and thus achieve better robustness against inaccuracies in, e.g., vessel direction and caliber.

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