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Interventional planning and assistance for ascending aorta dissections

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Abstract — In this paper, we present our actual works concerning interventional planning and assistance for ascending aorta dissections.

The first part is about aortic dissection retrieval in Computed Tomography Angiography (CTA) images. It is based on fast marching segmentation, mathematical morphology and digital topology frameworks.

The second part is about the registration of the outer aortic wall onto X-ray fluoroscopic angiographic images. It is based on either Digital Reconstructed Radiographs (DRRs) or parametric model projection, then on image transformation descriptors (ITDs) computing.

Our motivation is first to give 3D views of dissecting aorta in peroperative step to assess the dissection, then to augment the current angiographic view by registering aortic dissection features during the intraoperative step to assist clinicians.

Keywords aortic dissections, segmentation, 3D/2D registration, augmented reality

I. INTRODUCTION

Aortic dissections are caused by one or more aortal tissue perforation(s) because of tissue weakness and blood pressure. That consists of one or several intimal tear(s) (Fig. 1 (left)) and a clear separation between layers of the aortic media, resulting in two separate blood flow channels: the true lumen (the normal bed of blood in the aorta) and the false lumen (a channel entirely within the media which appears during an aortic dissection). The layer of detached aorta tissues between lumens is called intimal flap.

To deal with aortic dissections, physicians require preoperative 3DComputed Tomography Angiography (CTA) images (Fig. 1 (right)), then intraoperative 2D X-ray (here fluoroscopic, by a monoplane C-arm system) angiographic images (Fig. 3).

Our works consist in segmenting 3D CTA images to retrieve aortic dissection features, and to register them to X-ray images.

II. SEGMENTATION

In this section, we summarize our different works related to aortic dissection features extraction. They are schematized in Fig. 2.

Preprocessing. Let us consider an initial CT aortic dissection image : I₁. We perform a crop to accelerate image processing : I₂.

Lumens retrieval. Fast marching method (FM in the following) is an efficient implementation of level set deformable models. We initialize FM by defining a single voxel inside a lumen. Is it possible to limit the number of iterations of the evolution process to avoid the propagation to other arteries (collateral arteries) but the 3D propagation front reaches the heart : I₅. By using a morphological opening, we may cut that part of heart : we obtain the image of the two interconnected lumens I₄.

By modifying the speed function of FM, and using an intermediate labelling image (of points in the neighborhood of the intimal flap [dark grayscale]), we succeed in separating the two lumens [2] : I₅ and I₆.

Lumens distinction. By using the fact that grayscale value of blood is higher inside the true lumen, a quick operator (mean intensity inside each lumen) allowed us to distinguish between the true lumen and the false lumen, obtaining thus the false lumen image I₇ and the true lumen image I₈.

Intimal flap retrieval. A two-step procedure (a se-
Figure 2 – Aortic dissection segmentation: $I_7$ false lumen, $I_8$ true lumen, $I_9$ intimal flap, $I_{10}$ entry tears in red.

Quel of $n$ morphological dilations followed by a sequels of $n$ morphological erosions) gives the image of the intimal flap: $I_9$.

Intimal tears materializing. The first challenge of our works was to obtain the entry tears from a CT. Because they do not correspond to physical tissues or organs, such a segmentation cannot be direct. Intimal tears may be considered as 3D holes in the intimal flap. Therefore, we used a holes filling algorithm proposed in Digital Topology framework[1] (Fig. 2): the surfaces filling holes in $I_9$ are shown in red in $I_{10}$; they correspond to the location of intimal tears.

III. REGISTRATION

The main goal of our registration framework is to provide an alignment of pre- and intra-operational medical images (see Fig. 3).

Source image for registration. See the top row of Fig. 3. The reference image is a 2D X-ray fluoroscopic angiography sequence, acquired directly in the interventional room. In order to augment the aorta visibility, a filtering technique has been applied on the whole set of angiographic frames, to yield a single 2D image called Amplification image. Binaring the previous image, we obtain the Binarized amplification, which is the fixed input for the Registration algorithm.

Target image for registration. The aorta (issued from a CTA 3D image) plays the role of the (moving) target to establish the 3D/2D correspondence.

Now, we give three ways to compute a 2D model of that target:

- One way [center row of Fig. 3] is to compute a Digitally Reconstructed Radiograph (DRR) from the aorta interior; the projection parameters are computed in the registration process.
- Another alternative solution [bottom row of Fig. 3] is to define an aorta parametric model (by interactive construction[3]) and to generate its Rapid model projection.
- The parametric model reconstruction issued from the segmentation image could also be considered (* on Fig. 3).

Both Binarized DRR and Model projection images are the alternative moving inputs for the registration algorithm.

Initial 3D pose computing. A novel ITD registration technique of rapid 3D/2D alignment is applied[4]. It computes directly the spatial correspondence between projections, using image transformation descriptors (ITDs), such as moments, area...

Accurate registration. The last stage of the pipeline is a Precise 3D/2D registration technique, well initialized by the previously described algorithm.

We will consider one of the iconic methods with mul-
timodal similarity measure (correlation or entropy based).

IV. CONCLUSION

In this paper, we presented our actual works concerning aortic dissection features extraction from a CTA image, and the registration of a segmented outer aortic wall on an X-ray fluoroscopic image. We have distinguished the two lumens (Fig. 4 (a)), augmented CTA slices with these data (Fig. 4 (b)), proposed a virtual angioscopy (Fig. 4 (c)).

Our future works consist in registering the whole set of features extraction (intimal tears, intimal flap, true lumen, false lumen) on fluoroscopic angiographic images; a sketch is given in Fig. 4 (d).

REFERENCES


