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Patient-specific numerical simulation of stent-graft deployment: Validation on three clinical cases



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ABSTRACT

Endovascular repair of abdominal aortic aneurysms faces some adverse outcomes, such as kinks or endoleaks related to incomplete stent apposition, which are difficult to predict and which restrain its use although it is less invasive than open surgery. Finite element simulations could help to predict and anticipate possible complications biomechanically induced, thus enhancing practitioners' stent-graft sizing and surgery planning, and giving indications on patient eligibility to endovascular repair. The purpose of this work is therefore to develop a new numerical methodology to predict stent-graft final deployed shapes after surgery. The simulation process was applied on three clinical cases, using preoperative scans to generate patient-specific vessel models. The marketed devices deployed during the surgery, consisting of a main body and one or more iliac limbs or extensions, were modeled and their deployment inside the corresponding patient aneurysm was simulated. The numerical results were compared to the actual deployed geometry of the stent-grafts after surgery that was extracted from postoperative scans. We observed relevant matching between simulated and actual deployed stent-graft geometries, especially for proximal and distal stents outside the aneurysm sac which are particularly important for practitioners. Stent locations along the vessel centerlines in the three simulations were always within a few millimeters to actual stents locations. This good agreement between numerical results and clinical cases makes finite element simulation very promising for preoperative planning of endovascular repair.

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1. Introduction

Rupture of abdominal aortic aneurysm (AAA) is the primary cause of death of over 10,000 people in the United States each year (Kochanek et al., 2011). Prevention of AAA rupture is achieved either by open surgery or endovascular repair (EVAR). EVAR is associated with reduced postoperative mortality and morbidity compared to conventional surgery. However, the incidence of aneurysm related secondary procedures remains higher following EVAR than after open repair (Greenhalgh et al., 2010).

Endoleaks and stent-graft (SG) limb thrombosis are among the most frequent causes of secondary interventions following EVAR. Endoleaks may be associated with incomplete SG apposition on the arterial wall (Albertini et al., 2001, 2005). Graft kinking has been proved to favor stenosis and thrombosis (Carroccio et al., 2002; Cochennec et al., 2007). Highly tortuous and stiff calcified arteries are commonly associated with difficult outcomes (Albertini et al., 2006; Sternbergh et al., 2002). Therefore, mechanical behaviors of both SG and arterial wall appear to play an important role in the pathophysiology of these complications.

Within this context, finite-element analysis (FEA) could help predicting SG positioning inside patient-specific AAA, thus enabling surgeons to anticipate complications. Several teams have started to investigate this promising research field. Some of them focused on EVAR long term outcomes through fluid dynamics (Figueroa et al., 2010; Georgakarakos et al., 2014; Howell et al., 2007) and

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fluid-structure interactions to estimate the blood mechanical action onto SGs (Layman et al., 2010; Li and Kleinstreuer, 2006; Molony et al., 2010; Prasad et al., 2013). Despite their major interest, these studies present the following limitations: (i) they do not take into account SG deployment and (ii) most of them use oversimplified SG models, with a single equivalent homogenous mechanical behavior. Other groups investigated stent mechanics and their deployment in arteries (Auricchio et al., 2011; Holzapfel et al., 2005; Mortier et al., 2010). Following these pioneering studies, a number of researchers (De Bock et al., 2013; Kleinstreuer et al., 2008) have used FEA to model stentgraft structures, with both stents and graft mechanical behaviors. Our group recently achieved similar simulations (Demanget et al., 2013, 2012a) on several marketed SG limbs, which were validated against in-vitro bending tests (Demanget et al., 2012b). A step further has consisted in simulating SG deployment in aneurysm models. We developed simulations of SG deployment in idealized iliac aneurysm models (Perrin et al., 2015). De Bock et al. (2012) performed a SG deployment inside a silicone model and compared simulated and invitro SG positions. Auricchio et al. (2013) simulated the deployment of a custom-made tube aortic SG inside the corresponding patient-specific aneurysm model. Each of the above mentioned studies had limitations. The first one did not model patient-specific aneurysms and the second did not validate the simulation against in-vivo data. The third study simulated a tubular custom-made SG, much simpler than the bifurcated manufactured SGs usually used to treat AAAs.

The objective of the present study was to develop a FE methodology to simulate the deployment of marketed bifurcated SGs in patient-specific aneurysm models. In order to reduce computational costs, an original morphing technique was developed to simulate SG deployment.

2. Methods

2.1. Clinical summary

Three patients were included in this study after informed consent and approval from the Institutional Review Board. Patients, SGs and aorto-iliac characteristics are reported in Table 1. The devices were marketed by Medtronic (Santa Rosa, CA, USA) except for one right limb in clinical case #2 made by Cook Medical (Bloomington, Indiana, USA). Pre-operative and one-month post-operative computed tomography angiography (CTA) scans were available for all patients.

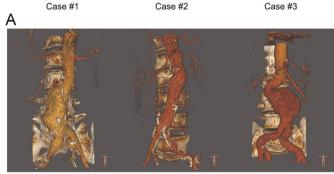
Table 1 Clinical summary of the three EVAR procedures.

	Case #1	Case #2	Case #3
Sex	Male	Male	Male
Age (y)	70	58	78
Stent-graft:			
Main body	ENBF-28-13-C-	ENBF-28-13-C-145-	ENBF-28-20-
	145-EE	EE	C-170-EE
Right limb	ENLW-16-20-C- 95-EE	ZSLE-13-56-ZT	ENLW-16-24- C-95-EE
Left limb	ENLW-16-28-C- 80-EE	ENLM-16-13-C-95- EE	None
Left extension	None	ENEW-13-13-C-80- EE	None
Aorto-iliac anatomy	Left and right	Main body rotated	60° angulated
and SG deployment features	iliac aneurysms	by 75° around the longitudinal axis	AAA neck
Aneurysm sac thrombus	Length: 60 mm Maximum thick- ness: 20 mm	Length: 100 mm Maximum thick- ness: 20 mm	None

2.2. Vessel geometries and constitutive modeling

Surgery oriented Endosize® software (Therenva, France) was used to extract aortic and iliac vessel centerlines and vascular lumen contours from pre-operative scans (Kaladji et al., 2013). Centerlines of the arteries were constituted of a set of points, spaced by 5 mm to obtain a smooth centerline interpolation, onto which were centered B-splines describing the vascular lumen contour. Each B-spline had 10 control points on the lumen surface, in each plane orthogonal to the centerline. The continuous geometry of lumen surface was generated by surface interpolation driven by the B-splines in ANSYS DesignModeler software (ANSYS, Inc., Canonsburg, PA). The arterial lumen surface was then meshed with 3-node linear shell elements (1.5 mm mean edge length) with 1.5 mm and 1.0 mm thicknesses for aortic and iliac surfaces, respectively. 3D pre-operative scans (A) and corresponding triangular meshes (B) are shown in Fig. 1.

Assuming that SG deployment induced small strains within the arterial wall, the latter was modeled as an orthotropic linear elastic material. In-plane mechanical parameters were computed by linearizing, throughout the physiological range of pressures, Holzapfel–Gasser–Ogden anisotropic hyperelastic model (Gasser et al., 2006), with parameter values obtained for above 61 year-old patient abdominal aortas by Haskett et al. (2010). For that purpose, a hyperelastic pipe (20 mm diameter, 1.5 mm thickness, 100 mm length) was subjected to *in-vivo* loading conditions (100 mmHg internal pressure and 10% longitudinal stretch (Horný et al., 2013)).



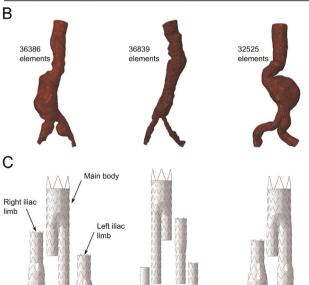


Fig. 1. Pre-operative scans (A), corresponding arterial wall meshes (B) and pre-stressed models of the various SG components used during EVAR (C) for the three clinical cases.

Then, the tangent stiffness matrix around these *in-vivo* conditions was derived by subjecting the pipe to different loading increments: pressure (5 mmHg), twist (2°) or longitudinal displacements (0.2 mm). The longitudinal, circumferential and shear elastic moduli, along with the Poisson ratio were obtained from the results of these simulations. The same value as in-plane shear modulus was chosen for the transverse shear moduli. The parameters of the hyperelastic model and the linearized elastic moduli are reported in Table 2.

2.3. Stent-graft modeling

Digitized geometries of SG main bodies and limbs were provided by the manufacturer, except the Cook Medical limb whose model had been validated in Demanget et al. (2012b), the dimensions of which were scaled to the current clinical case. Stents were meshed

Table 2Parameters of the hyperelastic Holzapfel–Gasser–Ogden model used for linearization and in-plane parameters of linearized orthotropic model.

Parameters and values of hyperelastic Holzapfel-Gasser-Ogden model:						
c_{10} (MPa)	Stress-like material parameter	100.9				
k_1 (GPa)	Stress material parameter	4.07				
k_2	Dimensionless parameter	165.55				
κ	Level of dispersion parameter	0.16				
θ (deg)	Fiber orientation angle	48.4				
K (GPa)	Bulk modulus	75.5				
In-plane parameters and values of linearized orthotropic linear elastic model:						
E_{z} (MPa)	Elastic modulus along vessel centerline	3.58				
E_{Θ} (MPa)	Elastic modulus along vessel circumferential	1.11				
	direction					
$ u_{\ThetaZ}$	Arterial wall in-plane minor Poisson ratio	0.44				
$G_{z\theta}$ (MPa)	Arterial wall in-plane shear modulus	4.0				

with linear beam elements (0.075 mm mean length). The superelastic behavior of Nitinol stents was modeled with Auricchio's model (Auricchio and Taylor, 1997) and implemented in a subroutine included in FEA software Abaqus (Simulia, Dassault Systems, Providence, RI, USA). The constitutive parameters of Nitinol, in the range of literature values, were provided by the manufacturer. Grafts were meshed with linear 4-node shell elements (0.4 mm mean edge length). Polyester fabric was modeled as an orthotropic elastic material. In-plane and bending stiffnesses characterized in a previous study of our group (Demanget et al., 2012a) were used.

During the SG manufacturing process, expanded stents diameters are oversized compared to graft diameters. For Medtronic components, a preliminary FEA was performed to tie the oversized stents to the graft. For the Cook limb, Z-stents were not oversized and modeled according to Demanget et al. (2012b). The resulting pre-stressed SG models are depicted in Fig. 1C.

2.4. Simulation methodology

A major challenge of simulating SG deployment in patient-specific models of aneurysm is to find appropriate boundary conditions for SG introduction. Our methodology, although different from the actual surgical procedure, has the potential to be used for any SG model and any aneurysm model. Also, it avoids simulating the full crimping and progressive deployment of the SG which are time expensive and may lead to numerical instabilities. A detailed description of the four steps of our simulation, which are described hereafter, is provided in Appendix A.

The main body, the iliac limbs and extensions were first compressed radially (slight crimping stage) and assembled (Fig. 2A). Then, the assembled SG was inserted inside a virtual tubular shell (Fig. 2B). From this configuration, proper displacements were prescribed onto the nodes of the virtual shell to morph its geometry onto the pre-operative geometry of the patient's aneurysm

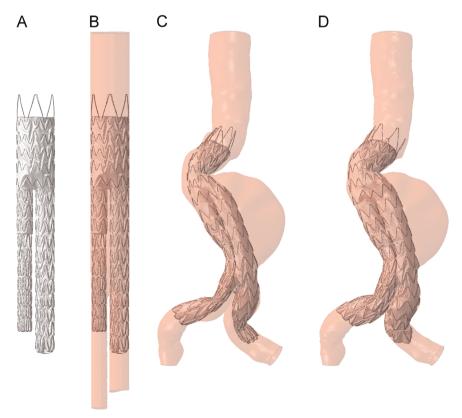


Fig. 2. Consecutive steps involving finite-element analysis: main body and limbs crimping and relative positioning (A); SG insertion and positioning in the tubular shape (B); Mesh morphing from tubular shape to actual pre-operative geometry (C); Mechanical equilibrium between SG and arteries (D).

while prescribing contact to maintain the SG inside the shell. Note that, during this step, the shell did not present any mechanical behavior and only acted as a geometrical constraint. At the end of this step, the deployment of the SG inside the pre-operative geometry of the AAA was simulated (Fig. 2C). Finally, the shell elements were ascribed the linearized AAA mechanical properties and all the boundary conditions previously assigned onto the AAA were released, for the SG to recoil and deform the vascular lumen until reaching static mechanical equilibrium (Fig. 2D). Only proximal and distal ends of the vessels remained clamped.

All simulations were carried out with the explicit FE solver of Abaqus v6.12 software. Time increments (adjusted *via* mass scaling) and time steps (Table 3) were chosen to obtain fast results while keeping the ratio of kinematic and internal energies under 10% to avoid spurious dynamic effects, as shown in Fig. 3. The FE simulations were run on 12 CPUs computers, 2.66 GHz, 24 GB RAM.

Contacts were modeled using the general penalty algorithm implemented in Abaqus, with the standard Coulomb friction law. The friction coefficient was set to 0.4, *i.e.* in the range of literature values (Vad et al., 2010).

3. Model validation methodology

3.1. Post-operative scan registration and stent segmentation

To validate the simulations, post-operative scans following EVAR procedures were used to quantify the position error of the simulated SGs. The voxel sizes of the three patient scans were respectively $0.85 \times 0.85 \times 0.625 \text{ mm}^3$, $0.76 \times 0.76 \times 0.7 \text{ mm}^3$ and $0.85 \times 0.85 \times 0.7 \text{ mm}^3$. Firstly, rigid image registration was used to align post-operative and pre-operative scans in the same coordinate system. Ten anatomical landmarks were picked on iliac bones, vertebras and vessel calcifications far from the surgical area, on each set of pre-operative and post-operative CTAs. A Matlab function based on the iterative closest point method was used for the rigid registration.

Table 3Time parameters of all steps involving FEA and corresponding computation times, for the three clinical cases. Time step is the length of the step prescribed to Abaqus. Time increment is the targeted increment, achieved using mass scaling. CPU time reflects the time a single processor would require computing the simulation. Real time reflects the real time required computing the simulation on our 12 processors setup.

	Simulation Time step (s)	on parameters Time incre- ment (s)	Computati CPU time (h)	on times Real time (h)
Case #1				
SG assembling and crimping	0.04	5E-8	199:00	16:35
Deployment in the pre- operative geometry Mechanical equilibrium	0.03	3E-8	147:00	12:15
	0.15	3E-8	845:00	70:25
Case #2				
SG assembling and crimping	0.05	4E-8	282:00	23:30
Deployment in the pre- operative geometry	0.07	4E-8	266:00	22:10
Mechanical equilibrium	0.15	4E-8	534:00	44:30
Case #3				
SG assembling and crimping	0.017	5E-8	84:00	7:00
Deployment in the pre- operative geometry	0.03	5E-8	113:00	9:25
Mechanical equilibrium	0.15	5E-8	474:00	39:20

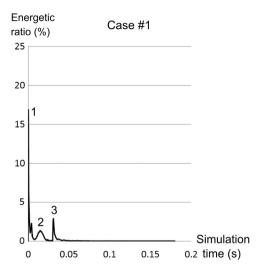


Fig. 3. Representative evolution of the ratio of the kinetic energy divided by the total energy along the deployment simulation, for case #1. 1: First contact between SG and morphed AAA. 2: Morphing step. 3: Beginning of the mechanical equilibrium simulation step.

After this procedure, the mean distance between registered preoperative points and post-operative points was 2.0 \pm 1.2 mm.

Therefrom, stent centerlines were segmented by manually picking the center of stent cross-sections on each slice of the post-operative scans (Fig. 4A). Stents located in overlap regions or in areas where vessels were highly calcified could not be properly segmented and were not considered in the validation process.

3.2. Cylinder fitting on stents

In the three registered simulation results and post-operative CTAs, a cylinder was defined for each stent using a Matlab Toutine, so that the distance from stent points to the cylinder surface was minimal. For example, Fig. 4B shows cylinders fitted on case #3 post-operative stents. In order to gauge the relevance of the simulations, radius and cylinder center position were compared between registered simulations and post-operative scans. For each simulated stent, diameter error e_D was estimated and normalized by the actual deployed stent diameter. The position error of stents center was split into a longitudinal distance error e_c along the arterial centerline and a transverse distance error e_t in the plan normal to the centerline.

4. Results

4.1. Qualitative assessment

The simulation results for the three clinical cases are presented in Fig. 5A. The superimposition of simulated and real stents, shown in Fig. 5B, gives a visual assessment of the results.

In the first clinical case, simulated and post-operative stent locations are nearly identical except for the distal stents of the right iliac limb. These stents and the two stents inside the left iliac aneurysm were not fully deployed; they are smaller in our simulation.

In the second case, simulated and actual stents are well superimposed at the proximal and distal SG ends. However, 75° longitudinal axial rotation of the main aortic body is not reproduced in the simulation.

In the third case, simulated iliac limbs are shorter and distal stents are transversally shifted, while good agreement of aortic body stents position can be observed.

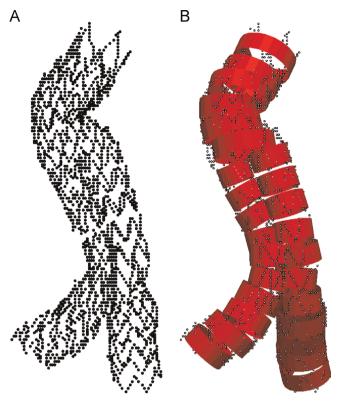


Fig. 4. Stents segmentation on the post-operative scan (A) and cylinder fitting (red) on these segmented stents (black) (B). (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

It is also worth mentioning the ability of the simulation to reproduce SG kinks in a realistic manner, as highlighted by the black arrow present in the third case in Fig. 5B.

4.2. Quantitative comparison

Values of previously defined errors are plotted in Fig. 6. Averaged error values for all stents are reported in Table 4.

Case #1 presents almost only small values of e_{cl} (mean value is 0.2 ± 1.6 mm). Stents located in the aneurysm sac and at the distal end of the right limb, have the worst position along the centerline (maximum e_{cl} value is 2.9 mm). The values of e_t are rather small and uniformly distributed on all stents (mean value is 2.7 ± 2.2 mm), the maximum transverse position error being in the aneurysm sac. Diameter agreement between simulated and real stents is best at the proximal part of the SG, while distal stents have a higher diameter error e_D , with a maximum error in the left iliac aneurysm (-35.3%).

In case #2, the distal and proximal stents are well positioned longitudinally (less than 2 mm error) whereas the stents inside the aneurysm sac have the worst longitudinal position of all cases (6.5 mm maximum error). The same observation can be made for the transverse error; e_t is small at SG ends and presents a maximum inside the aneurysm sac (e_t rises up to 14.1 mm in this area). As for case #1, most of the simulated stents have the same diameter as the actual stents (e_D ranges between -15% and 0%) and only one distal stent has a rather large diameter error (-35.6%).

In case #3, e_{cl} is between -2.5 and 2.5 mm (mean e_{cl} value is -1.3 ± 2.6 mm), except for the two most distal stents of the main body and the most distal stent of the iliac limb (where e_{cl} maximum is 5.2 mm). The transverse error is lower than in other cases $(2.3 \pm 1.3$ mm) and reaches 4.9 mm for distal stents. Mean diameter error is $-4.9 \pm 7.5\%$, with one single stent presenting a higher e_D value (-23.4%) in the aneurysm sac.

Computation times are presented in Table 3.

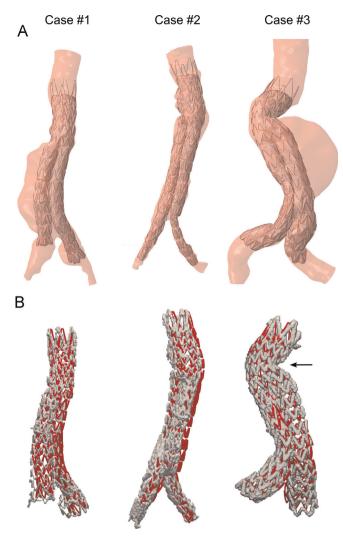


Fig. 5. Simulation results (A) and qualitative comparison of the geometry of simulated (red) and actual (white) deployed SGs (B) for the three clinical cases. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

5. Discussion

The present study reports a novel methodology to predict deployment of SGs in patient-specific aneurysm geometries. To our knowledge, this is the first time that several marketed SGs, with all their components, are numerically deployed inside patient-specific AAA models and that the results are validated against post-operative scans. Another original point of this work is the very limited manual intervention on the patient-specific part, thanks to a mesh morphing technique: the final steps of the simulation involving patient-specific data could be automated with previous SG crimping simulation, a pre-operative scan and few other parameters (SG diameters, proximal stent landing position, and overlap lengths) as only inputs.

We achieved a qualitative comparison by superimposing simulated and segmented actual stents geometries (Fig. 5). In general, this comparison was satisfactory for all cases. Most of the simulated stents were at the same location as actual stents. We could observe a slight difference in case #1 where right limb curvature and stent diameter were not exactly the same near the iliac artery bifurcation. This can be explained by incomplete stent expansion in this area. Right iliac arterial wall was not deformed enough by stent expansion and this resulted in a medial deployment of the limb, near the left

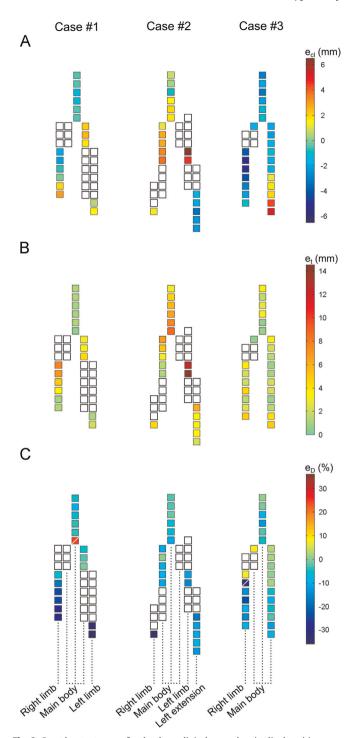


Fig. 6. Stent by stent errors for the three clinical cases: longitudinal position errors (e_{cl}) along the vessel centerline (A), transverse position errors (e_t) perpendicularly to the centerline (B) and relative diameter errors (e_D) (C). Each square is one stent; columns are representing different SG components. The error for each stent is coded by the color of the square. White squares represent stents which were not included in the statistics due to impossible segmentation on the post-operative scan. Crossed squares were not included in diameter statistics due to their elliptic shape. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

iliac artery. In case #2, main discrepancies were in the aneurysm sac, while position of stents at SG ends was well predicted. The initial rotation of the prosthesis around its longitudinal axis was not implemented in our simulations. This may explain the large errors (14.1 mm maximum e_t) at the main body stumps and proximal limb extremities. Case #3 is the simulation which showed the best

Table 4Mean, standard and maximum errors over all stents of the SG for each clinical case. The two crossed squares of Fig. 6 were not included in diameter statistics due to their elliptic shape.

	Case #1	Case #2	Case #3			
Stents position error along the Mean ± standard deviation Maximum	centerline (mm): 0.2 ± 1.6 2.9	1.1 ± 2.6 6.5	-1.3 ± 2.6 5.2			
Stents position error perpendic Mean \pm standard deviation Maximum	ularly to the center 2.7 ± 2.2	erline (mm): 5.4 ± 3.5 14.1	2.3 ± 1.3 4.9			
Stents diameter relative error (%): $ \begin{array}{cccc} \text{Mean} \pm \text{standard deviation} & -14.8 \pm 12.5 & -10.2 \pm 7.8 & -4.9 \pm 7.5 \\ \text{Maximum} & -35.3 & -35.6 & -23.4 \end{array} $						

agreement with the post-operative CT scan. Only marginal differences could be noted between simulated and actual SG shapes (especially the two distal stents).

Quantitative results were extracted to confirm the qualitative analysis. Fitting cylinders onto simulated and actual stents allowed us to compare stents center position along centerline's tangent and transverse directions, as well as stents diameter (Fig. 6 and Table 4).

In all cases, longitudinal error was less than 2.5 mm for 75% of stents. In cases #1 and #3, maximum longitudinal errors were 2.9 and 5.2 mm, respectively. These errors seem to be due to the fact that arteries were not deformed and straightened enough by the SG. In case #2, stents with the larger error values were all located in the aneurysm sac. This was partly explained by the axial rotation of the main body which was not implemented in the simulation. Another explanation was the lack of mechanical constraints around the SG in the aneurysm sac, particularly the absence of intraluminal thrombus (ILT) modeling. Finally, longitudinal error was reasonable in the perspective of clinical applications (i.e. less than 5 mm). Moreover, it is important to bear in mind that the final error is inherent to the simulation itself but also to pre- and post-operative scan registrations, stent segmentation and cylinder fitting.

The remarks are quite similar when looking at centerline transverse errors. Mean e_t value was small in case #1 as well as standard deviation (2.7 \pm 2.2 mm). High e_t values were seen only in the right iliac limb. Stents at SG ends of case #2 were well positioned radially whereas the actual deployment rotation of the main body induced higher e_t values in the aneurysm sac. In case #3, e_t confirmed the good agreement of simulation and post-operative scan.

In the three simulations, there was good diameter agreement between simulated and real stents. However, simulations seemed to slightly underestimate stent diameters as mean e_D values were all negative (-14.8% to -4.9%). Slightly overestimated AAA material properties may explain such error. In case #1, two stents were not fully deployed inside the left iliac aneurysm, as four distal stents of the right iliac limb. The effect of blood pressure (included within AAA mechanical behavior) may be one reason for the lack of stent expansion in aneurysm sacs. Local diameter underestimations (as for the right distal stent in case #2) may be related to the fact that SG balloon dilatation was not simulated.

SG longitudinal positioning was well predicted in our simulations. Implementation of this type of simulation in the pre-operative sizing process could help reducing inter-observer variability and increase accuracy of SG design, particularly in complex aortic aneurysms (Banno et al., 2014). Furthermore, assessment of SG kinks and stent apposition defects could assist in the prediction of limb thrombosis or endoleaks.

6. Limitations

Arterial walls were modeled with constant thickness and mechanical behavior linearized around *in-vivo* loading conditions. This assumption was validated by the good agreement between simulations and post-operative scans. Nevertheless, our simulations could benefit from more sophisticated modeling including thrombus (Riveros et al., 2015; Toungara and Geindreau, 2013) and calcifications. This could enhance accuracy of SG position in the aneurysm sac. Furthermore, large regional datasets gathered by other research teams (Roccabianca et al., 2014), statistical models (Vande Geest et al., 2006) or novel image processing techniques (Franquet et al., 2013) could enhance arterial wall modeling, with local patient-specific variations.

During simulations, we imposed as only boundary conditions that aortic and iliac extremities were motionless. Further developments could increase biomechanical realism by assigning elastic connectors at the ostias of internal iliac arteries or preventing any posterior displacement near the spine. Blood pressure modeling could also be improved. Fluid–structure interaction analysis would be an appropriate tool to impose pressure loads on both arterial and SG surfaces in contact with the blood. It would also allow us to simulate blood flow following SG deployment (Figueroa et al., 2010) and therefore potential endoleaks.

Finally, we did not intend to extensively reproduce all steps of the EVAR procedure in our simulation. Our goal was rather to predict final SG position with reduced computational costs. The obtained level of accuracy was acceptable from a clinical perspective. However, accuracy of SG position may be further increased by implementing intraoperative steps in the simulation, such as arterial deformation by stiff guidewire and introducer sheath, as well as SG ballooning. Pioneering work on aorto-iliac by stiff guidewire has been recently published by Kaladji et al. (2013).

7. Conclusion

EVAR is a complex surgery which requires cautious preoperative planning, from patient selection to stent-graft sizing, in order to prevent post-operative complications and secondary interventions. Within this context, numerical simulations associating patient-specific data and stent-graft modeling could provide information to choose the best-suited device in each case and anticipate complications. Towards this objective, we have proposed a novel computational methodology to predict the deployment of marketed stent-grafts inside patient-specific aneurysms. This strategy was validated on three clinical cases and numerical results showed good agreement with actual post-operative scans. These results emphasize the potential of numerical simulations to improve pre-operative planning of EVAR.

Conflict of interest

The authors have no professional, financial or commercial conflict of interest to declare.

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Appendix A. Supplementary material

Supplementary data associated with this article can be found in the online version at http://dx.doi.org/10.1016/j.jbiomech.2015.04.

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