Introduction

Along with the evolution of the aortic surgery the appearance of the thoracic endovascular aortic repair (TEVAR) provided a new therapeutic approach for the patients with acute and chronic aortic pathology. Combining the surgical with the endovascular repair broadened the vision in the treatment of the aortic arch. The surgical technique of frozen elephant trunk (FET) has developed through the years and is actually an excellent approach, providing the option for endovascular and surgical extension (1,2). However, the perioperative-stroke remains in the different case series between 2.5% and 20% (3). Additionally, the supra-aortic transposition has showed to be a reliable alternative excepting the total arch rerouting, which is related with a high rate of retrograde type A dissection (4,5). Nevertheless, there are patient groups with high risk, where neither the surgical or the hybrid repair is the first choice. In such cases total endovascular arch repair with the Bolton Relay plus double-branch endoprosthesis (Bolton Medical, Sunrise, FL, USA) could be a feasible option with excellent outcome.

However, not every patient is suitable for the approach and there are some clinical geometrical and technical details which should be considered. The geometrical and clinical details guide our path through the treatment of the thoracic aortic disease involving the aortic arch, because our primary goal is providing excellent therapeutic approach with feasible long-term outcome, reducing the aortic-related events without increasing the perioperative risks. Summarized, there are two major points, that should be considered for the use of the double branches prosthesis for the total endovascular arch repair: clinical assessment and geometrical details.

Clinical assessment

Currently, the estimation if a patient is suitable or non-suitable for classical surgery remains at the discretion of the individually treating physician as there is a lack of risk prediction models in patients with thoracic aortic pathology. However, concomitant cardiac and vascular condition and/or severe chronic obstructive lung disease
are known predictors of adverse outcome in classical surgery. Therefore, this group may qualify best for the total endovascular approach.

Regarding the usage of currently available risk scores, patients with higher EuroSCORE and ASA score levels (American Society of the Anesthesiologist) should be generally favored (6-8). Patients with connective tissue disease should be generally excluded. Life expectancy more than 2 years is recommended. Individuals with congestive heart failure in class III or IV should be excluded as well as patient with significant supraaortic atherosclerosis (stenosis of the internal carotid artery ≥70% by NASCET criteria) (8,9).

**Geometrical details**

The aortic anatomy should be evaluated with thin-sliced computed tomography of the entire aorta including the supraaortic, iliac and femoral vessels. “Shaggy aorta” due to ulcerative atherosclerosis has a very high risk of perioperative cerebrovascular embolic events and these patients might be suboptimal candidates for total endovascular arch repair. There is generally no recommendation in relation to the angulation. However, steep arches may have a weaker performance due to the risk of kinking and potential type IIIb endoleakage due to strut perforation. On the other hand, there are clearly defined needs about the diameter and length of the ascending aorta. Diameter should be 40 mm and preferably less and ascending length should be 65 mm or longer using a center-lined measurement.

The brachiocephalic trunk and the left common carotid artery (LCCA) should have regular diameters and the minimum diameter of the LCCA recommended is 7 mm. The window for the supra-aortic branches is 5 cm in length and the space from the brachiocephalic trunk offspring to the end of the LCCA has to be ≤50 mm. Currently, all devices are custom-made (Table 1).

**Table 1** Anatomical details and minimal requirements for the implantation (6)

<table>
<thead>
<tr>
<th>Geometrical details</th>
<th>Minimal requirements</th>
</tr>
</thead>
<tbody>
<tr>
<td>Length from sinotubular junction to the BT</td>
<td>≥65 mm</td>
</tr>
<tr>
<td>Diameter of the ascending aorta</td>
<td>≤40 mm</td>
</tr>
<tr>
<td>Diameter of the LCCA</td>
<td>≥7 mm</td>
</tr>
<tr>
<td>Space between BT and LCCA</td>
<td>≤50 mm</td>
</tr>
<tr>
<td>Oversizing</td>
<td>≤15%</td>
</tr>
<tr>
<td>Access vessel size</td>
<td>≥8–9 mm</td>
</tr>
</tbody>
</table>

**Prosthesis details**

The main body is delivered in a standard retrograde fashion with an outer sheath diameter of 25 French. The main body has two internal tunnels for the supra-aortic branches (LCCA—anterior tunnel, BT—posterior tunnel). The extensions are originally modified iliac limbs of the abdominal endovascular prosthesis of Bolton (Treo). The profile of the supraaortic extensions is 14 F. Oversizing more than 15% is generally not recommended (in all zones). Exception could be made by post dissection aneurysm, where the distal landing zone is sized according to the true lumen.

The self-alignment mechanism of the main body allows that the pre-curved tip adapts itself according to the curve of the aortic arch. The mounting of the supraaortic window to the outer curvature is automatically and radiopaque markers indicate the appropriate orientation. Nevertheless, exact active positioning remains a prerequisite (Figure 1).

**Procedure**

Revascularization of the left subclavian artery (LSA) is performed before the endovascular procedure for maintaining the inflow in the left vertebral artery for the cerebellar and spinal cord perfusion. The primary goal is the reduce risk of perioperative stroke and symptomatic spinal cord ischemia (6,10). If the left vertebral artery is non-dominant, overstenting of the LSA without revascularization may be performed. Neuromonitoring with motor evoked potentials and somatosensory evoked potentials combined with cerebral fluid drainage should be used if distal TEVAR extension is planned.

After systematic heparinization (100 IU/kg) the main body is delivered and the window for the supra-aortic branches is orientated correctly according to the radiopaque markers. After inducing hypotensive conditions using rapid pacing the main body is deployed. Next step is the implantation of the supra-aortic branches and begins with the posterior tunnel, which could be deployed through the right common carotid artery or the right subclavian artery using surgical cut down. The precise cannulation of the tunnel is verified.
by inflating a contrast-filled balloon. The procedure is followed likewise by the insertion of the second supra-
aortic branch for the LCCA (Figure 2). Finally, on table
angiography is performed to prove the result (Figure 3).
Mono antiplatelet therapy with 100 mg aspirin daily from
postop day 1 is recommended.

**Pros and cons**

The proximal landing zone is usually straight and, on this
account, endoleaks type IA are rare. Zone 0 per se has a
certain risk of retrograde type A dissection. However, the
incidence seems to be less than in a total arch rerouting and

*Figure 1* View of the DBP with the branches for BT and LCCA (A) side view with deployed supraaortic branches (B) inside view. DBP, double branched prosthesis; LCCA, left common carotid artery.

*Figure 2* Schematic presentation of the steps by the implantation of the aortic arch endo-prosthesis. (A) Aneurysm of the aortic arch; (B) implantation of the main body through the femoral or iliac vessels with orientation of the window for the supraaortic vessels according to the radiopaque markers. The arrows are showing the radiopaque markers; (C) the supraaortic branches are deployed after the main body. (I) The implantation of the supraaortic limbs begins with the posterior tunnel (RCCA); (II) finally, comes the deployment of the second limb for the LCCA through the anterior tunnel. The arrow shows the carotid-subclavian bypass. RSA, right subclavian artery; RCCA, right common carotid artery; LCCA, left common carotid artery; LSA, left subclavian artery.
TEVAR setting potentially due to the lack of tangential clamping of the ascending aorta (4,11). In addition, the knowledge that patients with the underlying pathology of type B aortic dissection also do have an inherently diseased ascending aorta has become accepted in the community and therefore these kinds of approaches are no longer offered to patients with this condition thereby reducing the incidence of retrograde type A aortic dissection. However, the endografts are custom made and time is needed for the manufacturing. Therefore, time per se excludes several patients from clinical implementation. Finally, there are no long-term results of the approach and the clinical experience is actually scarce.

Conclusions

Total endovascular aortic arch repair is a feasible option with an excellent aortic-related survival and very good neurological outcome. Our armamentarium in the treatment of the aortic arch has broadened with the availability of this approach. Still, there are some geometrical and clinical details, which must be taken into account when selecting patients. Nevertheless, the clinical experience with the new method is still limited and there is no data about the long-term durability of this approach. Consequently, for a potential recommendation of a broader implementation, further studies are needed.

Acknowledgements

None.

Footnote

Conflicts of Interest: Martin Czerny and Bartosz Rylski are consultants for Bolton Medical. Other authors have no conflicts of interest to declare.

References


doi: 10.21037/jovs.2018.05.27