ABSTRACT

Symptomatic unruptured abdominal aortic aneurysm (AAA) refers to a group of patients with intact AAA but who present abdominal and/or lumbar pain attributed to the aneurysm. This form of clinical presentation is potentially fatal, since its etiopathogenesis, involving acute changes in the aortic wall, including inflammation, increases the probability of impending rupture. It is clear that these patients should be referred to AAA repair. However, the timing of the intervention is controversial. Therefore, the aim of the present work was to review updated information on the diagnostic-therapeutic approach of symptomatic unruptured AAA.

Key words: Symptomatic unruptured abdominal aortic aneurysm - Inflammation - Open surgery - Endovascular aneurysm repair

INTRODUCTION

The etymology of the word aneurysm can be traced in ancient Greek language. (1,2) It derives from the word ἀνεύρυσμα (aneurysma), which means “dilation”. (2) Therefore, “aortic aneurysm” refers to an abnormal dilation of the aorta that compromises one or more wall segments. In this sense, an increase in diameter greater than 50% of the aortic diameter at the level of the diaphragm is accepted as abnormal. However, this definition is not always applicable, since often the limit between the healthy and pathological aorta is not precise. (3,4) Consequently, there is a general consensus in considering abdominal aortic aneurysm (AAA) to be a dilation of the aorta greater than or equal to 30 mm. (4-7) Multiple studies have been developed to evaluate the results of elective AAA repair, as well as those associated with emergency repair of ruptured aneurysms. (8-15) However, there is a third type of potentially fatal and scarcely studied clinical presentation of AAA: symptomatic unruptured AAA. (16) Therefore, the aim of the present work was to review updated information on the diagnostic-therapeutic approach of symptomatic unruptured AAA.

METHODS

Searches were carried out in PubMed/Medline, EMBASE, and Cochrane Clinical Trials electronic databases to identify clinical studies that evaluated the diagnostic-therapeutic approach of symptomatic unruptured AAA, using the terms “abdominal aortic aneurysm”, “symptomatic aneurysm”, “unruptured aneurysm”, “open surgery”, and “endovascular aneurysm repair”.

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The selection of articles was made according to the following criteria: a) publications issued from 1990 to the present; b) observational studies, clinical trials, systematic reviews and recommendations of scientific societies; c) human studies; and d) articles referring to management of symptomatic unruptured AAA. Studies were excluded if the full text was not accessible.

Primary outcomes of this article were: a) to synthesize knowledge about epidemiology, prognosis and diagnostic-therapeutic approach of symptomatic unruptured AAA; and b) to develop an algorithm towards its diagnostic-therapeutic management.

Definition and epidemiology
Symptomatic unruptured AAA refers to a group of patients with intact AAA but who present abdominal and/or lumbar pain attributed to the aneurysm. Another symptom includes tenderness to palpation overlying the AAA in the abdomen, back, or flank. (17) The presence of pain is due to multiple causes, including acute changes in the aortic wall, as we will describe later. In large aneurysms, pain may be caused by compression of adjacent structures. Symptomatic unruptured AAA patients generally do not have arterial hypotension because the aortic wall is intact, with no evidence of retroperitoneal hemorrhage. (7) The reported incidence of this type of clinical presentation is between 3% and 15%. (18,19)

Pathophysiology
The pathophysiology of symptomatic unruptured AAA involves acute changes in the aneurysmal wall, including increased wall stress and intra-thrombus expansion or new bleeding, raising the probability of impending aneurysm rupture. (20) Evidence demonstrates the role of aneurysmal wall inflammation in the etiopathogenesis of this type of clinical presentation. In this sense, increased 18-fluorodeoxyglucose metabolism by positron emission tomography (PET/CT) in patients with symptomatic unruptured AAA compared with asymptomatic ones correlated with increased inflammatory infiltrate density in aneurysmal wall biopsies. (21)

Prognosis and repair outcomes
The current perioperative mortality rate in symptomatic unruptured AAA patients is below that previously reported in the literature. In this regard, De Martino et al. showed a global in-hospital mortality rate of 1.7% (2.2% in operated patients and 0% in patients undergoing endovascular repair). (22) Similarly, Chandra et al. published a contemporary (2005-2014) global perioperative mortality of 5.9%, involving 8% of surgical patients and 5% of those undergoing endovascular procedures. (17) Regarding postoperative complications and long-term survival, De Martino et al. reported that the rate of postoperative complications, including kidney failure, acute myocardial infarction, arrhythmias and respiratory insufficiency was 35% in patients with symptomatic unruptured AAA versus 20% and 63% in those with asymptomatic and ruptured AAA, respectively. In the case of 1 and 4-year survival, this was 83% and 58%, respectively, in patients with symptomatic unruptured AAA, while in asymptomatic ones it was 89% and 73%, and in those with ruptured AAA, 49% and 35%, respectively. (22) In conclusion, in-hospital mortality of symptomatic unruptured AAA repair is similar or discreetly superior to that observed for the elective intervention. However, the rates of postoperative complications and long-term survival are intermediate compared with asymptomatic and ruptured AAA repair. (23-27)

Diagnostic-therapeutic approach
When a symptomatic AAA is suspected, a large caliber venous access should be placed, invasive blood pressure monitoring should be performed, and diagnostic confirmation and immediate treatment should be accomplished at the center where the patient is located. If the conditions to carry out the diagnosis and treatment in the medical center are not met, immediate referral to a high complexity hospital should be activated. (7) Most patients will present with hemodynamic stability since, by definition, there is no rupture of the aortic wall. (17) A high degree of clinical suspicion is essential given the poor prognosis of ruptured and symptomatic unruptured AAA. In a meta-analysis that included studies published after 1990, ruptured AAA misdiagnosis was seen in 32% of cases. The most common erroneous differential diagnoses were ureteric colic and myocardial infarction. (28,29) Figure 1 postulates an algorithm for the diagnostic-therapeutic approach of symptomatic AAA. All individuals admitted to an emergency center with abdominal and/or lumbar pain with a known diagnosis of AAA and who are stable from a hemodynamic point of view, must quickly undergo an abdominal-pelvic computed tomography angiography (CTA). (5) Clinicians may use an abdominal ultrasound to help make the diagnosis whenever it is available. (30) While ultrasound can identify the presence of AAA and intraabdominal fluid, it is less effective at detecting signs of aortic rupture. (31) Ultrasound is therefore not considered a confirmatory test for rupture; however, it represents a useful tool in identifying AAA. Those patients with clinical suspicion and presence of AAA on ultrasound could be considered high-risk for symptomatic AAA, and this may allow for expedited referral CTA. On the other hand, in patients with low clinical risk of symptomatic AAA, the absence of AAA on ultrasound may be sufficient not to carry out further tests, especially if there is another more probable cause that explains the symptoms. (32) If symptomatic AAA is confirmed, a retroperitoneal hemorrhage or the absence of aortic rupture with persistent pain despite optimal medical treatment, including analgesia and control of vital signs, indicates mandatory emergency repair. If, on the contrary, no tomographic findings of aneurysmal rupture are observed, the timing of intervention is controversial and remains a challenge. (8) Numerous retrospective case series have published higher morbidity and mortality rates in emergency AAA repair compared with urgent intervention (18-26% vs. 4-5%), (33-35) In addition, no deaths associated with aneurysm rupture have been reported in patients with symptomatic unruptured AAA, in whom the intervention was postponed and performed semi-electively. (18) Concerning this aspect, different series which have evaluated the causes of death in this group of patients have shown that, in most cases, these are secondary to myocardial infarction, respiratory insufficiency, kidney failure, multiorgan failure and sepsis. (34,36) In our opinion, considering that the morbidity and mortality of patients with symptomatic unruptured AAA undergoing emergency surgery is related to cardio-re-nal-pulmonary dysfunction, it is reasonable to prioritize the optimization of these systems prior to AAA repair. In this sense, we consider that these objectives could be achieved in a short time span of 12-24 hours, provided the medical center has the necessary resources. It is essential that the patient be admitted to a critical care unit, with suitable vital sign monitoring. From a cardiovascular viewpoint, there is evidence on the usefulness of focused cardiac ultrasonogra-
phy (FoCUS) for critical patient assessment, as well as for presurgical evaluation. (37) An echocardiogram performed in the emergency room will provide essential information for hemodynamic management, including left ventricular ejection fraction, presence of significant valve diseases and the degree of patient’s preload, through the transmitral filling pattern, E/E’ relationship, pulmonary systolic pressure, and inferior vena cava diameter, and inspiratory collapsibility. (37) Frequently, elderly patients present with reduced preload prior to surgery with a negative impact on cardiac output, and this can be optimized with an adequate volume load. Some cases may require more invasive monitoring with a Swan Ganz catheter, allowing therapeutic guidance as a function of the hemodynamic state of the patient. Moreover, achieving adequate preload will reduce the risk of postoperative kidney failure. From a respiratory standpoint, reversible obstructive pulmonary disease can improve through a short treatment with intravenous steroids and aerosol bronchodilators. (34)

Therefore, in this subgroup of patients with symptomatic unruptured AAA it is crucial to identify individuals at high preoperative risk who may benefit from a fast optimization of their clinical status (cardiological, pulmonary and/or renal) prior to an urgent intervention. (17,34) With the objective of answering this problem, different publications have evaluated the usefulness of scores and certain biochemical markers to predict in-hospital mortality in patients with symptomatic unruptured AAA. Antonello et al. reported that the Glasgow Aneurysm Score is a good predictor of perioperative morbidity and mortality after urgent open surgery of symptomatic unruptured AAA. The authors suggest that patients with score ≤90 can safely undergo an emergency open repair. However, patients with score ≥90 should undergo a thorough assessment and optimization of the preoperative clinical sta-

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**Fig. 1. Algorithm for the diagnostic-therapeutic approach of symptomatic AAA**

**Modified Sullivan’s algorithm (34)**

AAA: abdominal aortic aneurysm; CTA: computed tomography angiography; NLR: neutrophil-to-lymphocyte ratio; PLR: platelet-to-lymphocyte ratio

1. Critical Care Unit admission
2. Vital signs monitoring
3. Optimization of cardio-renal-pulmonary function

- Acute onset of abdominal/lumbar pain
- Previous diagnosis of AAA or ultrasonographic evidence of AAA in the emergency room
  - Yes
  - No

- Hemodynamic stability
- Hemodynamic instability

- Abdominal-pelvic CTA
- Absence of retroperitoneal hemorrhage
- Retroperitoneal hemorrhage

- Inadequate clinical status, NLR ≥ 6.4 and PLR ≥ 185
- Adequate clinical status, <6.4 and PLR <185

- Emergency AAA repair
- Urgent AAA repair once the patient has been clinically optimized (12-24 hours after admission)
tus. (38) On the other hand, considering the role of inflammation in the etiopathogenesis of this disease, Garagoli et al. recently evaluated the usefulness of inflammatory biochemical markers for the prediction of in-hospital mortality in patients submitted to surgical and endovascular repair of symptomatic unruptured AAA. The authors concluded that patients with neutrophil-to-lymphocyte ratio ≥ 6.4 and/or platelet-to-lymphocyte ratio ≥ 185 are at high risk and could benefit from a surveyed waiting conduct prior to optimization of the presurgical clinical status or, even consider an endovascular repair. (36,39) These biomarkers are widely available, have low cost and, in addition, have the advantage of representing the inflammatory state of the patient at the time of hospital admission. This is different from the Glasgow Aneurysm Score that uses clinical variables referred to the patient’s prior medical history and considers shock as the only variable of the clinical status at the time of presentation, which we regard as insufficient since, as previously mentioned, these individuals generally present with hemodynamic stability. (36,39)

However, during this preoperative evaluation, it is fundamental that the treating physician is alert to signs and symptoms leading to a mandatory emergency intervention. The development of hypotension, tachycardia, oliguria or metabolic acidosis requires an immediate repair. (7)

Decision on the type of symptomatic unruptured AAA repair: open surgery versus endovascular intervention

Evidence shows that endovascular intervention reduces morbidity and mortality after symptomatic unruptured AAA repair. In this sense, this type of repair offers some potential advantages compared with conventional surgical treatment in this clinical setting: less physiological aggression to the organism, less need for deep anesthesia, less blood loss, minimizes hypothermia and reduces intervention time. (7) The analysis of the ENGAGE Registry demonstrated lack of difference in the elective endovascular repair of asymptomatic AAA versus symptomatic unruptured AAA. (19) A systematic review identified 23 observational studies evaluating 7040 symptomatic unruptured and ruptured AAA repairs (with surgical and endovascular intervention) and showed that patients undergoing endovascular intervention presented a lower rate of in-hospital mortality compared with open surgery (odds ratio 0.624, 95% CI 0.518-0.752; p <0.0001), as well as reduced length of hospital stay, lower bleeding and decreased intervention time. (40)

However, to perform endovascular AAA repair it is essential to fulfill the required anatomical criteria in addition to an institutional program for the emergency endovascular intervention. In this context, the role of CTA is crucial to define those patients fit for endovascular repair. (5,8) Moreover, we consider that the transfer to a high-complexity vascular center is appropriate for hemodynamically stable patients and anatomically apt for endovascular repair, especially if the surgical risk is high.

CONCLUSIONS

Symptomatic unruptured AAA constitutes a clinical form of potentially life-threatening AAA, with an intermediate prognosis between asymptomatic patients and those with ruptured aneurysms. Although the timing of intervention is still controversial, we consider it is essential to stratify individual preoperative risk, with the object of identifying high-risk patients who would benefit from the optimization of their clinical status prior to the intervention. Endovascular repair is feasible in this clinical scenario, mainly in patients at high surgical risk, provided the required anatomical criteria are fulfilled and the institution has a program for emergency endovascular intervention.

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Conflicts of interest

None declared.

(See authors’ conflict of interests forms on the web.

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