ABSTRACT – Purpose: The subclavian arteries can vary on their origin, course or length. One of the most common anatomical variations is the right subclavian artery originating as the last branch of the aortic arch. This artery is known as a retroesophageal right subclavian artery or “luxory artery”. The right retroesophageal subclavian artery usually is described as not producing symptoms, being most discoveries coincidental. Nevertheless, it may be the site of formation of atherosclerotic plaque, inflammatory lesions or aneurysm. Case Report: The present study describes a case of right retroesophageal subclavian artery and discusses the findings according to their clinical and surgical implications. Conclusion: The anatomic and morphologic variations of the aortic arch and its branches are significant for diagnostic and surgical procedures in the thorax and neck. If a right retroesophageal subclavian artery is diagnosed during aortic arch repair, corrective surgery should be considered. Intensive care patients should be screened before long term placement of nasogastric tube, in order to avoid fistulization and fatal hemorrhage.

INTRODUCTION

The subclavian arteries can vary on their origin, course or length. Right retroesophageal subclavian artery (RRESA) is a frequent defect of the embryological aortic arches and its incidence is estimated at between 0.5 and 1.8%. This variation is due to interruption of the fourth right aortic arch between the notches for the common carotid artery and subclavian artery with the left fourth arch remains intact. A regression of the right subclavian artery proximal portion occurs and the retroesophageal aortic arch persists.

The RRESA is usually described as not producing symptoms, being most discoveries coincidental. Nevertheless, it may be the site of formation of atherosclerotic plaque, inflammatory lesions or aneurysm. The seriousness of RRESA aneurysm is associated with the high risk of clot-related events, tracheal or venous compression and rupture.

The anatomic and morphologic variations of the aortic arch and its branches are significant for diagnostic and surgical procedures in the thorax and neck. The present study describes a case of RRESA and discusses the findings according to their clinical and surgical implications.

CASE REPORT

During a routine dissection at the School of Medicine of Triângulo Mineiro Human Anatomy Laboratory, an unusual origin and course of the carotid arteries and the right subclavian artery was observed. The body was that of a 54-year-old white female. Particular care was taken in dissecting the regions of the neck and thorax to preserve the aortic arch and its branches. With the aid of an electronic digital caliper (range of 0-300 mm, resolution 0.01 mm, Gehaka, SP, Brazil), the diameters and total length of the common carotid arteries were measured from their point of origin to their terminal branching. The diameters of the subclavian arteries at their point of origin were also measured. A common trunk of the right and left common carotid artery (bicarotid trunk), the left subclavian artery and the RRESA were derived from the aortic arch in that order (Figure 1). The right recurrent laryngeal nerve was not formed, but the nerves to the larynx and esophagus branched directly from the vagus as it descended in the neck. The left recurrent laryngeal nerve was not formed, but the nerves to the larynx and esophagus branched directly from the vagus as it descended in the neck. The RRESA originates distal and posterior from the aortic arch. V indicates the left vagus nerve and the arrowheads indicate the left recurrent laryngeal nerve. RCCA = right common carotid artery, LCCA = left common carotid artery, ES = esophagus and TR = trachea.

DISCUSSION

Since the incidence of a RRESA is reported to range from 0.2% to 2.5%, this anomaly is not markedly rare and our percentage is in accordance to that reported in the literature. The RRESA is more frequent in women and mongoloid children and is also associated with Chromosome 22q11 deletion. Our casuistic is small (only 3 cases, 1 female and 2 male) but we found that one of our male cadavers had a mental disorder not well characterized. Since the frequency of a RRESA is high in at least two genetic disorders mentioned above, and most of these diseases course with mental disorders, this could be an explanation why we had more male than female cadavers with the variation since it is described to be more frequent in women. Although described as usually asymptomatic, the RRESA can cause dysphagia lusoria as a clinical finding, due to the compression of the esophagus between the right common carotid artery and the trachea anterior to it and the right subclavian artery posterior to it. Nevertheless, acute ischemia of the right upper limb due to thrombosis of an aberrant subclavian artery was described by Akers Jr et al. and Boas et al. Akers Jr et al. reported a case of nonocclusive and presumably nonatherosclerotic thrombosis in a 35-year-old woman that led to an extensive distal embolization requiring amputation of the right hand. Boas et al. reported a case of acute ischemia of the right upper extremity due to an extensive thrombosis of a RRESA in a 79-year-old woman that was successfully treated by extensive thrombectomy of the arteries in the upper extremity and right subclavian-to-carotid artery transposition. In both cases, the authors mention the delayed diagnosis of the underlying etiology of the thrombosis. This is a clear example of when knowledge of an anatomical variation is helpful in clinical practice. The presence of a RRESA should be considered especially when ischemia of the upper limb is severe and occlusion is located at the shoulder level.

Another rare but clinically important complication of the RRESA is the arterioesophageal fistula, an usually fatal event. The presence of a RRESA is a risk factor for rupture into the esophagus because the close anatomical relationship between the aberrant artery and the upper respiratory and digestive tract. The mechanism underlying the development of a fistula involves the induction of limited necrosis of the digestive and arterial walls by pulsatile compression of the esophageal wall between the RRESA and a rigid intubation catheter. Feugier et al. revisited the literature and found 11 cases of fistula between a nonaneurismal retroesophageal subclavian artery and the esophagus. Hematemesis was precipitated by esogastric intubation in all cases. Fistulization
was associated with predisposing factors in three reports and hemorrhage was fatal in seven cases. Feugier et al.² tested that clinical diagnosis of this complication is difficult because of its extreme rarity. In the absence of any evidence suggesting the possibility of an anomaly of the aortic arch system, there are no specific manifestations. They also conclude that the extreme unfavorable prognosis of this rare complication combined with the relatively high incidence of the RRESA in the general population underlines the need of prevention. If a RRESA is diagnosed during aortic arch repair, corrective surgery should be considered. Intensive care patients should be screened before long term placement of nasogastric tube.

REFERENCES