Vascular Ring; Spectrum of Pathologies and Scenarios for Diagnosis and Management

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Abstract: Background: Vascular rings represent approximately 1% of all congenital cardiac anomalies, with Edward’s classification being the first to outline them into a complete or partial vascular ring. A Complete ring is a combination of patent vessels, atretic vascular segments or ligamentous structures. Methods: We reviewed patients’ records from our Pediatric Cardiac Surgery Unit, over 10 years. We encountered 63 patients from January 2009 to January 2019. Our patients were 57 (90.5%) complete vascular rings, 5 (7.9%) pulmonary artery slings, and 1 (1.6%) patient with innominate artery compression. Results: The age of our Patients ranged from 2 to 57 months (mean 16.25±1.3 m), weight ranges from 3k to 26k (mean 9.87±0.8kg). Males were 38 (60.3%) and male: female ratio was 1.5:1. In our series various types of vascular rings were encountered; Double Aortic Arch was the right dominant arch variant in 25 (53.2%) and left arch dominant in 7 (14.9%) while the co-dominant variant was 15 (31.9%). Fifty-seven cases of vascular rings operated through left posterolateral thoracotomy. Five cases of pulmonary artery sling (7.9%) operated by median sternotomy and one case (1.6%) of innominate artery compression. No reported intraoperative mortality in our study. Conclusion: Vascular ring is a rare anomaly that needs suspicion for diagnosis. The operative strategy relies on extensive dissection and division of the non-dominant arch and division of the ligamentum, freeing all fibrous bands between the arch and esophagus and trachea. For augmented results; we need collaboration from pediatricians and pulmonologists who must be aware of such anomalies.

Keywords: Anomaly, Arch, Sling, Ring, Compression

1. Background

Under the term vascular rings; a constellation of anomalies related to defective development of the branchial arches. They present as an abnormal position, doubling, interrupted anomalous origin of the aortic arch or its branches [1]. They encircle the aero digestive tracks within the mediastinum; thus presentation is wheezy chest, recurrent pneumonia, stridor, troubled feeding or choking, and failure to thrive. This quite uncommon subcategory of cardiac anomalies accounts for 1% or less with equal incidence in both sexes. They may present alone or are associated with other major anomalies of the heart like Ventricular septal defect, Fallot’s tetralogy, or trans-positioning of great vessels [1].

2. Methods

2.1. Study Design

It is a retrospective observational longitudinal study on vascular rings management at pediatric cardiac surgery unit. We reviewed our patients’ records over a period of 5 years. Mansoura University Children Hospital is a high volume, referral institution sees from 8-12 of these patients per year. We encountered 63 patients subjected to surgery from January 2015 to January 2020. Our patients were 57 (90.5%) complete vascular rings, 5 (7.9%) pulmonary artery slings, and 1 patient (1.6%) with innominate artery compression. About one-quarter of these patients (23.8%) have a chromosome abnormality. The ethical committee had approved both the records review process and the study plan.
The classification adopted and used generally classifies vascular rings into 2 main types:

1. Complete Rings. [1, 2]
   - Double Aortic Arch; (Right arch dominant. -Left arch dominant. Balanced). Right arch/Left ligamentum
   - Double Aortic Arch; (Mirror image branching. -Mirror image branching.)

2. Incomplete Rings
   - Innominate artery compression syndrome.
   - Pulmonary artery sling.
   - Left arch/aberrant Right subclavian artery.

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**Figure 1.** Embryonic 6 aortic arches and various rings [2]. (a) Schematic diagram depicting development of the embryonic 6 pairs of aortic arches and branches. Gray areas represent segments that normally involute; (b) A normal left aortic arch which results from regression of the right arch. The first branch arising from the arch is the brachiocephalic artery, followed by the left common carotid and left subclavian arteries; (c) Hypothetical double aortic arch system with bilateral aortic arches and bilateral ductus arteriosi; (d) Right aortic arch with mirror image branching. This anomaly results from regression of the left arch between the LSC artery and the descending aorta. The first branch arising from the arch is the left brachiocephalic artery, followed by the RCC and then the RSC arteries; (e) Right aortic arch with aberrant left subclavian artery arising from a retroesophageal diverticulum of Kommerell. This anomaly results from regression of the left arch between the LCC and LSC arteries (gray), usually with persistence of the left sixth arch as ligamentum arteriosum; (f) Left aortic arch with aberrant right subclavian artery without diverticulum of Kommerell. This anomaly results from regression of the right arch between the RCC and RSC arteries, including the right ductus arteriosus (gray).

**Figure 2.** a MSCTA (frontal view, left image) done for the representative patient in our study population showing RAA with Left aberrant subclavian artery with tracheal compression, b (back view, right image).
2.2. Data Collection

Table 1. Showing types of vascular rings in our study.

| Type of vascular ring | number | percentage |
|-----------------------|--------|------------|
| DAA                   | 47     | 74.6%      |
| Aberrant RT innominate| 1      | 1.6%       |
| RAA                   | 10     | 15.9%      |
| Pul. Art. Sling        | 5      | 7.9%       |

Pertinent data extracted from their records either preoperative, intraoperative, and postoperative such as demographic data, presenting symptoms, findings in preoperative chest x-ray (CXR), and computed tomography (CT) and intra-operative correlation, surgical complications, and eventually the outcome (Table 1).

2.3. Statistical Analysis

Data analysis was carried out via using SPSS (Statistical Package for Social Science) version 21. Qualitative data were represented in the form of mean ± standard deviation. Continuous variable data were expressed as the mean ± standard deviation. Statistical comparisons were made through Student’s t-test for continuous variables and chi-square test for categorical variables. Results were considered statistically significant if P-value is less than 0.05.

2.4. Radiological Diagnosis

Plain chest X-ray postero-anterior and lateral views were done in 100% as routine in the outpatient clinic. Upon suspecting or finding a right sided arch; patients were further investigated. No Barium swallow was used, due to inconvenience; especially in young infant or preschool children. Echo-Doppler examination (88.9%) which was very helpful in diagnosing and defining which type of vascular ring. Bronchoscopy was used in 25 cases (39.7%), done by pulmonologists before referral. Cardiac catheterization in only 2 patients (3.2%). Multislice CT with 3D reconstruction was used in 59 patients (93.7%). It is considered an excellent, non-invasive diagnostic modality. See table 3.

Table 2. Showing presenting symptoms in relation to different types of vascular rings.

| Diagnostic modalities | Total (n=63)| Percentage |
|-----------------------|-------------|------------|
| Chest X-ray           | 63          | 100%       |
| Echo                  | 56          | 88.9%      |
| Bronchoscope          | 25          | 39.7%      |
| Angiography           | 2           | 3.2%       |
| MSCT                  | 59          | 93.65%     |

3. Results

Patients’ age in our study ranged from 2 months to 57 months (mean 16.25±1.3 m), weight ranges from 3kg to 26kg (mean 9.87±0.8 kg). Males were 38 cases (60.3%) and male: female ratio was 1.5:1. In table 1 various types of vascular rings were encountered. In our series, Double Aortic Arch (DAA) was right dominant arch variant in 25 (53.2%) and left arch dominant in 15 (31.9%). Symptoms presented in our series were mostly due to tracheal compression in form of stridor, bronchial asthma, feeding problem, and failure to thrive shown in table 2. We depended on a variety of diagnostic methods in our series, mainly CXR in 100% see table 3. CXR findings like an absent left aortic knuckle, right-sided aortic arch, tracheal compression or indentation. Echocardiography with tissue Doppler assessment was used not only for confirming diagnosis but also helped in detecting 4 cases with associated cardiac anomalies (1 Fallot’s tetralogy (1.6%), 1 Patent ductus arteriosus (PDA) (1.6%), 2 patent foramen ovale (PFO) (3.2%).

Fifty-seven cases of vascular rings (90.5%) operated by left posterolateral thoracotomy. Five cases of pulmonary artery sling (7.9%) operated by median sternotomy and one case of innominate artery compression (1.6%), no reported intraoperative complications nor intraoperative mortality in our study.

No injury happened to either phrenic or recurrent laryngeal nerve. One case was complicated with pneumothorax (1.6%) and was treated by an intercostals tube. No bleeding in our cases as a complication. Three cases (4.7%) were re-ventilated. Chylothorax happened in 4 cases (6.3%) and managed by total parental nutrition and fat intake restriction alone in one case (25%) and in 3 cases (75%), somatostatin was added but eventually, we went back to theatre for thoracic duct ligation in 2 cases (50%).

We have only one case of mortality (1.6%), pulmonary artery sling with diffuse narrowing of lower tracheal and this case died in the intensive care unit (ICU) 12 h early postoperatively. In our study, total postoperative stay ranged from (3-23 days).

Correlations through Q- square and student t-test revealed no significant association between the type of complication and type of vascular ring and age or weight, or gender, P value>0.05. Also, no significant association between ICU and ward stay and age or gender or presenting symptoms like stridor or asthma, p value>0.05.

There is a significant association between ward stay and types of vascular rings, p-value<0.05. Also, there is a significant association between ICU and ward stay and Post-Operative complications, p-value<0.05.
4. Discussion

During fetal life, any anomaly associated with the development of the 6 pairs of aortic arches can lead to a vascular ring. Gross in 1945, introduced that term where a vascular anomaly encircles the aerodigestive tracts either completely or partially after he successfully relieved tracheal obstruction in a double aortic arch patient. Thus, they are classified as complete or partial; but both share the same symptoms and the same treatment algorithm. The double aortic arch (DAA) is by far the most common variant, seen in 57 cases (74.6%) and this is similar to other reports from similar countries where the incidence of congenital heart diseases are near equal; they reported 68% and 56.3% incidence of DAA in 2 different reports [1, 3]. The right-sided aorta with aberrant subclavian and left ligamentum is our second common type; representing 15.9% which is lower than most reports, where RAA represents 45% or 26.8% [4, 5]. The third common type in our study was pulmonary artery sling accounting for 7.9% and it’s considered high if compared to Ronald’s report, it constitute 3.7% [5].

Usually, the presenting symptoms of the incomplete vascular ring are severe and annoying; so patients present earlier. The severity of the symptoms is directly related to the degree of compression exerted by the rings. Because of this, complete rings present earlier versus incomplete rings. This is similar to other reported surgical series [6]. On the other hand, it could be totally asymptomatic and discovered during an investigation for other conditions.

The presence of right-sided aortic arch is a diagnostic clue in plain CXR; also, the lateral indentation of the tracheal air column. We have done CXR in 100% of our cases. Bronchoscopy was done in 39.7%, findings were external compression, narrowing, and transmitted pulsation; but we no longer recommend except in severe dyspnea or stridor. But we think noninvasive radiologic imaging is enough to help and bronchoscope may cause laryngeal edema [4].

We used echocardiography principally due to its accuracy, simplicity and because of being non-invasive method, but in the presence of the highly specific and sensitive diagnostic MSCT, lately, we did not use echocardiography except if we suspect associated cardiac anomalies, exactly in 56 out of 63 cases (88.9%). This nearly coincides with Shah and his colleagues, who used echocardiography in 95% in their study [7].

We had 4 cases with associated anomalies, Fallot tetralogy (1.6%), PDA (1.6%), PFO (3.2%). Compared to Khalfan’s study in 2006, the associated cardiac anomalies were found in 14 patients (17%), ventricular septal defect in 10 patients (12%), atrial septal defect in 4 patients (5%), patent ductus arteriosus in 3 patients (4%), and tetralogy of Fallot in 3 patients (4%) [8].

In our study; cardiac catheterization was needed in 3.2% as an accurate method to delineate the anatomy but this was no longer used because of being an invasive technique and the need for contrast and replaced by Echocardiography and/or CT chest. Cardiac catheterization was used largely as in Khalfan’s series [8] that reported 33 (41%) of 81 cases, also in another series of cardiac catheterization was used in 21 (47.7%) from 44 cases [9].

Multislice CT is a noninvasive modality for diagnosis. It illustrates the arch dominance and tracheal compression. We used it in 59 cases, and our results match the results of many reports in which all cases were diagnosed by contrast-enhanced MSCT [10, 11]. On contrary, in Raghavan’s study, they used Magnetic resonant Imaging (MRI), but as we notice, this was a relatively old study [3]. However, we recommend MSCT for all our cases for better delineation and good decision for operation. The only disadvantage is radiation exposure. Our preference was based on the accessibility to that new imaging modality and the preference of the radiologist and surgeon interpreting the images as it provides an excellent correlation with intraoperative findings.

Stridor is the commonest presenting symptom among cases of complete vascular ring; mostly these cases were misdiagnosed as foreign body inhalation and referred for diagnostic bronchoscopy or sent for ENT evaluation. In our series, 26 of 63 cases (41.3%) presented with stridor with 24 from the DAA group. Our results are comparable to other series with similar results [3, 7].

In our series, aerodigestive symptoms like dysphagia for solid foods or feeding troubles was seen in (12.7%). Other studies reported higher percentage 27% and 31.3% [7, 12].

Four of our patients had associated cardiac anomalies (6.3%). Other series reported a higher incidence of cardiac anomalies as in the series but in our series, there were no associated non-cardiac anomalies compared to the same report, they documented respiratory anomalies in 13 patients (40.6%) [12].

Our principle of operating as soon as possible in any case with a vascular ring was due to early repair allows normal trachea and esophagus growth. Similar to Yoon’s report in 2012, [12] we approached through a limited left posterolateral thoracotomy through the 3rd intercostal space in (92.1%), similar to others who reported (91.8%) and (92%) [4, 8].

Five sternotomies were used for cases with pulmonary artery sling. That allowed easy dissection and a wide non-compromised field for re-implantation. In our series, no Video-assisted thoracoscopic surgery (VATS) was used in contrast to Rahul’s report in 2007 who operated 39 cases using VATS (60.9%) of 64 cases [7].

We tried our best to extubate as early as possible, on the operating table, or after transfer to the surgical intensive care unit to avoid upper airway edema. With pulmonary toilet, nasotracheal suction and chest percussion following extubation. Usually, better results are harvested and minimal postoperative atelectasis or pneumonia. We always inform and educate their parents that symptoms could persist up to six months before disappearing completely. In our 63 cases, postoperative stay ranges from (3 to 23 days), the extended stay was attributed to the development of chylothorax, and that coincides with a recent report where the mean hospital stay was (18.8±9.3) days [13].

Mortality in our series was one case with pulmonary artery
sling. The infant had diffuse narrowing of the lower trachea extending to the left main bronchus. We started to operate off-pump, then we had to go on by-pass emergently, it was complicated by difficult weaning from the ventilator as the stenosed tracheal segment mostly caused air trapping, CO2 narcosis, and death.

5. Conclusion

A vascular ring is a rare congenital cardiovascular anomaly that needs a high index of clinical suspicion for diagnosis. Operative strategy to minimize improper or incomplete management and prevent recurrence mainly relies on extensive dissection, delineation of the different components of the anomaly, and division of non-dominant arch and division of the ligamentum, freeing all fibrous bands between the arch and esophagus and trachea. For better results; we need collaboration from many specialists who will receive these patients before us as pediatricians and pulmonologists who must be aware of such anomaly with its multiple variants.

Limitation of the Study

This is a retrospective study that has the inherent deficiencies of this category of studies. Besides that, it is a single-center study, the number of cases is relatively small over a single decade period of time, with different variants of the same pathology. Data of follow up was needed but not documented in our files.

List of Abbreviations

- VSD: Ventricular septal defect
- CXR: Chest x-ray
- CT: computed tomography
- DAA: Double Aortic Arch
- PDA: Patent ductus arteriosus
- PFO: Patent foramen ovale
- ICU: Intensive care unit
- MRI: Magnetic resonant Imaging

Declarations

Ethics Approval and Consent to Participate

The study was approved by the Institutional Review Board (IRB) according to the code of ethics of the World Medical Association (Declaration of Helsinki) for studies involving humans, though the Committee’s reference number was not applicable. The need for informed consent was waived due to the retrospective nature of the study.

Consent for Publication

Not applicable.

Availability of Data and Material

Data are available on request.

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Competing Interests

No conflict of interest and nothing to disclose.

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