

Unilateral internal jugular vein phlebectasia in an adult: Management and one year follow-up

SAGE Open Medical Case Reports
Volume 7: 1–4
© The Author(s) 2019
Article reuse guidelines:
sagepub.com/journals-permissions
DOI: 10.1177/2050313X19836351
journals.sagepub.com/home/sco



Mazyad Alenezi¹, Abeer Alaglan² , Abdulhakeem Almutairi¹, Sultan Alanazy¹ and Osama Al Wutayd³

Abstract

Phlebectasia describes an anomalous, fusiform dilatation of a vein. In the neck region, the internal and external jugular veins are mostly affected. To our knowledge, this is the first case in Saudi Arabia of internal jugular phlebectasia affecting an adult female. We describe a 61-year-old female with complaints of a neck swelling she noticed 4 years ago. Initially, the swelling increased in size and reached a stable level. It was asymptomatic and only enlarged during Valsalva maneuver. Flexible nasolaryngoscopy and computerized tomography scan showed unremarkable examination. Follow-up after 1 year with US Doppler showed no progression. Internal jugular phlebectasia is a rare disorder which is often diagnosed during childhood. More often than not, it does not cause any significant morbidity. Since it is a benign condition, observation is advised with regular monitoring. For asymptomatic lesions, surgical intervention is recommended if cosmetic or psychological concerns are present.

Keywords

Phlebectasia, internal jugular vein, head and neck, otolaryngology, vascular anomalies

Date received: 27 November 2018; accepted: 5 February 2019

Introduction

Phlebectasia is an aberrant, fusiform dilatation of a vein.¹ It can affect any neck vein, mainly, internal and external jugular and, less commonly, anterior jugular and superficial communicus veins.² Two-thirds of the cases arise from the internal jugular vein.³ To our knowledge, this is the first case in Saudi Arabia of internal jugular phlebectasia affecting an adult female.

Case presentation

A 61-year-old female presented with complaints of a neck swelling she noticed 4 years ago. Initially, the swelling increased in size and reached a stable level. The patient also reported that the swelling enlarged with coughing and straining. She denied stridor, dysphagia, dysphonia, neck pain, and having other neck masses. On examination, there was a mass located in the right lower region of the patient's neck with no overlying skin changes. The mass increased in size with Valsalva (Figure 1). On palpation, the mass was compressible with transmitted pulse. Auscultation showed no bruit. Flexible nasolaryngoscopy showed unremarkable examination. Computerized

tomography (CT) scan of head and neck with contrast was done (Figures 2–4). No treatment was indicated. Thus, the patient was scheduled for follow-up and observation. On patient follow-up after 1 year with US Doppler, the internal jugular vein was patent within size compared to previous CT image and no thrombus formation (Figure 5).

Discussion

Internal jugular phlebectasia is a rare disorder which can be either unilateral or bilateral, and it is diagnosed often during

¹Department of Otolaryngology, Head and Neck Surgery, College of Medicine, Qassim University, Buraidah, Saudi Arabia

²Qassim University, Buraidah, Saudi Arabia

³Department of Family and Community Medicine, Unaizah College of Medicine, Qassim University, Unaizah, Saudi Arabia

Corresponding Author:

Mazyad Alenezi, Department of Otolaryngology, Head and Neck Surgery, College of Medicine, Qassim University, P.O. Box 6655, Buraidah 51452, Qassim, Saudi Arabia.

Email: mazyad@qumed.edu.sa





Figure 1. Right-sided neck swelling during Valsalva maneuver.

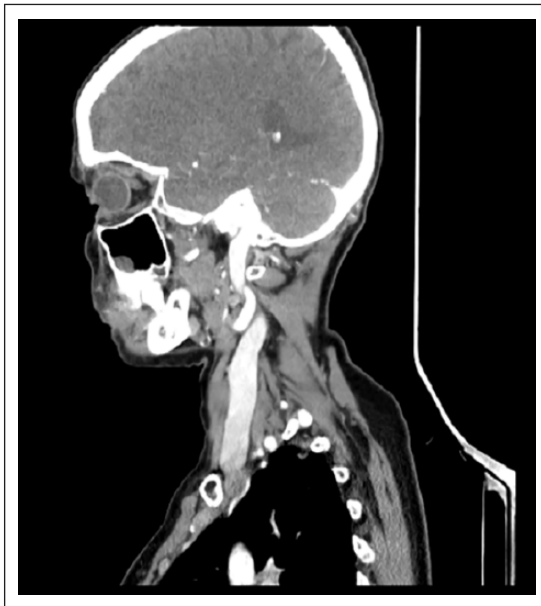


Figure 2. Sagittal contrast-enhanced CT image of the neck showing a well-defined large saccular internal jugular vein.

childhood.⁴ Thus, adulthood onset of internal jugular phlebectasia when compared to childhood onset is extremely rare.^{2,5} A couple of cases were reported to have internal jugular phlebectasia presenting in adulthood; however, the neck swelling was evident since birth for both cases.^{3,6} Due to the rarity of this condition, no likely etiology has been found. However, it is hypothesized that a primary lack of elasticity of the venous wall can be a reason explaining mainly congenital cases. In addition, elevated pressure in the internal jugular vein is

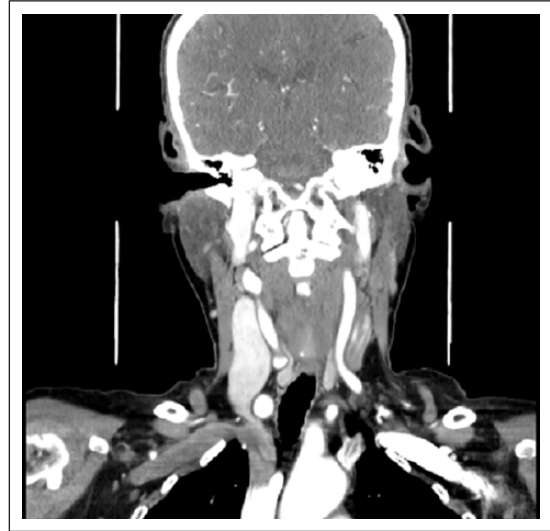


Figure 3. Contrast-enhanced coronal CT image is showing the extent of the dilatation of the right internal jugular vein.



Figure 4. Contrast-enhanced axial CT image showing dilated right internal jugular vein with heterogeneous contrast filling.

another hypothesis that can lead to phlebectasia.^{7,8} Trauma was the etiology in one case who presented with an internal jugular phlebectasia following a motorcycle traffic accident with a compound fracture of the right zygomatic bone.⁸ Unlike our patient, in adults, internal jugular vein phlebectasia is most common on the left side of the neck, while in children, it was commonly found on the right side.^{2,9} It typically presents with a benign swelling over the lateral side of the neck on the affected side, seen on effort. More often than not, it does not cause any significant morbidity which was the presentation of our patient.¹⁰ The chief complaint of patients is mainly due to cosmetic concerns.⁸ Other infrequent presentations reported were associated with pain and chest heaviness.^{2,5} Dyspnea and

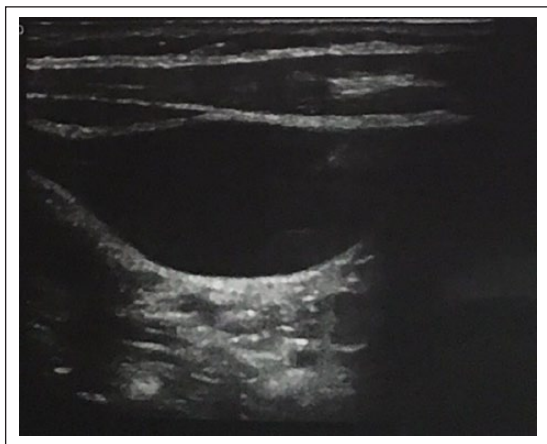


Figure 5. Doppler ultrasound of the right internal jugular vein on 1-year follow-up showing patent vein with the same diameter compared to previous CT and no thrombus formation.

dysphonia seemed to be the presenting complaint in patients having malignancies combined with internal jugular phlebectasia.¹¹ Generally speaking, symptomatic presentation or developing complications such as Horner's syndrome and thrombosis is rare.¹² Valsalva maneuver is the most valuable clinical sign used to diagnose the dilatation of internal jugular vein.¹⁰ Internal jugular phlebectasia is first noticed on physical examination, subsequently color Doppler imaging confirms the diagnosis and is said to be the gold standard.^{12,13} Furthermore, CT scans are utilized to rule out any causative structural lesions.⁸ The differential diagnosis of a progressive swelling in the neck on straining includes tumors or cysts of the upper mediastinum, external laryngeal diverticula or laryngoceles, inflation of the apex of the lung, as well as cystic hygromas.^{5,6} Since internal jugular phlebectasia is a benign condition, conservative observation is advised with regular monitoring. However, if the patient decides on conservative management, it should be emphasized that observing the lesion for any changes and recording the natural course is essential. Also, the patient must be protected from all potential injuries to avoid hemorrhage and infection.¹⁴ Surgical intervention for internal jugular phlebectasia is generally reserved for symptomatic patients together with the asymptomatic who present with cosmetic or psychologic concerns.^{12,13,15} The conventional surgical approaches include ligation of the internal jugular vein via a transcutaneous cervical approach or transaxillary approach as well as longitudinal constriction suture venoplasty and encapsulation or partial resection of the phlebectasia.¹⁶ Both procedures have been reported to be safe and successful in eliminating the phlebectasia.^{8,15} In recent years, constriction suture venoplasty and encapsulation has been the preferable choice since it did not require occlusion.¹⁵ Surgical resection is also indicated in the setting of thrombosis within the phlebectic segment, which is an uncommon complication.^{17,18} As such complications are rare, they do not necessitate anticoagulation therapy.^{19,20} However, since there is grave concern of embolization of the thrombus to the lungs, it is an indication for surgery.¹⁸

Conclusion

Internal jugular phlebectasia presenting in adults is a very rare phenomenon as it is a disorder often seen during childhood. It typically presents with a benign swelling over the lateral side of the neck, exacerbated by Valsalva maneuver. The main complaint is predominantly cosmetic concerns. The gold standard for diagnosis is color Doppler imaging which can be aided by CT to rule out any causative structural lesions. Because internal jugular phlebectasia is a benign condition, conservative observation is advised. Surgical intervention for internal jugular phlebectasia is commonly reserved for symptomatic patients together with the asymptomatic who present with cosmetic or psychologic concerns. The conventional surgical approaches include ligation of the internal jugular vein as well as longitudinal constriction suture venoplasty and encapsulation or partial resection of the phlebectasia.

Declaration of conflicting interests

The author(s) declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

Ethics approval

Our institution does not require ethical approval for reporting individual cases or case series.

Funding

The author(s) received no financial support for the research, authorship, and/or publication of this article.

Informed consent

Written informed consent was obtained from the patient(s) for publication of this case report and any accompanying images. A copy of the written consent is available for review by the Editor-in-Chief of this journal.

ORCID iD

Abeer Alaglan  <https://orcid.org/0000-0003-3919-4976>

References

- Gerwig WH Jr. Internal jugular phlebectasia. *Ann Surg* 1952; 135(1): 130–133, <http://www.ncbi.nlm.nih.gov/pubmed/14895158> (accessed 2 February 2018).
- Krstačić A, Župetić I, Soldo SB, et al. Jugular phlebectasia in adult—an overlooked cause of cervical pain. *Neurol Sci* 2017; 38(9): 1703–1704, <http://www.ncbi.nlm.nih.gov/pubmed/28474146> (accessed 2 February 2018).
- Gerek M, AkçLam T, Ümittalas D, et al. Internal jugular phlebectasia surrounded by mature adipose tissue. *Otolaryng Head Neck Surg* 2016; 128: 761–763, <http://journals.sagepub.com/doi/abs/10.1016/S0194-59980223272-6?journalCode=otoj> (accessed 2 February 2018).
- Pul N and Pul M. External jugular phlebectasia in children. *Eur J Pediatr* 1995; 154(4): 275–276, <http://www.ncbi.nlm.nih.gov/pubmed/7607276> (accessed 2 February 2018).

5. Ogbole GI, Irabor AE, Adeoye PO, et al. Internal jugular phlebectasia in an African adult. *BMJ Case Rep* 2010; 2010: bcr0220102724, <http://www.ncbi.nlm.nih.gov/pubmed/22778187> (accessed 2 February 2018).
6. Huang C-C and Chen H-C. Internal jugular vein phlebectasia. *Vasc Med* 2013; 18(6): 372–373, <http://journals.sagepub.com/doi/10.1177/1358863X13497785> (accessed 2 February 2018).
7. Paleri V and Gopalakrishnan S. Jugular phlebectasia: theory of pathogenesis and review of literature. *Int J Pediatr Otorhinolaryngol* 2001; 57(2): 155–159, <http://www.ncbi.nlm.nih.gov/pubmed/11165654> (accessed 2 February 2018).
8. Rha EY, Choi IK and Byeon JH. Internal jugular phlebectasia in a patient with facial trauma. *Arch Plast Surg* 2013; 40(4): 456–458, <http://www.ncbi.nlm.nih.gov/pubmed/23898449> (accessed 2 February 2018).
9. Thulasiraman V, Ramesh Pandian T, Cheralathan S, et al. Internal jugular phlebectasia as an incidental finding in cervical spine surgery. *Indian J Orthop* 2010; 44(4): 471–473, <http://www.ncbi.nlm.nih.gov/pubmed/20924495> (accessed 2 February 2018).
10. Ferreira L and Haguette É. Bilateral internal jugular phlebectasia. *Int Arch Otorhinolaryngol* 2007; 11(2): 220–223, http://arquivosdeorl.org.br/conteudo/acervo_eng.asp?Id=429 (accessed 2 February 2018).
11. Micozkadioglu SD and Erkan AN. Internal jugular vein anomaly: a lateral branch of the internal jugular vein in the neck. *Egypt J Ear Nose Throat Allied Sci* 2011; 12(1): 77–79, <https://www.sciencedirect.com/science/article/pii/S2090074011000144> (accessed 2 February 2018).
12. Zohar Y, Ben-Tovim R and Talmi YP. Phlebectasia of the jugular system. *J Cranio Maxillofac Surg* 1989; 17(2): 96–98, <https://www.sciencedirect.com/science/article/pii/S1010518289800538> (accessed 2 February 2018).
13. Haney JC, Shortell CK, McCann RL, et al. Congenital jugular vein phlebectasia: a case report and review of the literature. *Ann Vasc Surg* 2008; 22(5): 681–683, <http://www.ncbi.nlm.nih.gov/pubmed/18462919> (accessed 2 February 2018).
14. Bindal SK, Vasisth GOP and Chibber P. Phlebectasia of internal jugular vein. *J Surg Tech Case Rep* 2012; 4(2): 103–105, <http://www.ncbi.nlm.nih.gov/pubmed/23741586> (accessed 2 February 2018).
15. Jianhong L, Xuewu J and Tingze H. Surgical treatment of jugular vein phlebectasia in children. *Am J Surg* 2006; 192(3): 286–290, <http://www.ncbi.nlm.nih.gov/pubmed/16920419> (accessed 3 November 2018).
16. Hung T and Campbell AI. Surgical repair of left internal jugular phlebectasia. *J Vasc Surg* 2008; 47(6): 1337–1338, <http://www.ncbi.nlm.nih.gov/pubmed/18514849> (accessed 2 February 2018).
17. Chakraborty S, Dey PK, Roy A, et al. Internal jugular vein phlebectasia presenting with hoarseness of voice. *Case Rep Vasc Med* 2013; 2013: 386961, <http://www.ncbi.nlm.nih.gov/pubmed/24369523> (accessed 3 November 2018).
18. Matsunaga K and Kishi K. Phlebectasia of the external jugular vein with thrombosis: report of a case. *Surg Today* 2014; 44(6): 1180–1183, <http://www.ncbi.nlm.nih.gov/pubmed/23589057> (accessed 26 January 2019).
19. Calligaro KD, Ahmad S, Dandora R, et al. Venous aneurysms: surgical indications and review of the literature. *Surgery* 1995; 117(1): 1–6, <http://www.ncbi.nlm.nih.gov/pubmed/7809821> (accessed 26 January 2019).
20. Phookan S, Strickland PT, Hanna B, et al. Internal jugular venous pseudoaneurysm in a patient with heart failure and severe tricuspid regurgitation. *Case Rep Vasc Med* 2017; 2017: 3592459, <https://www.hindawi.com/journals/crivam/2017/3592459/> (accessed 26 January 2019).