

# World Journal of *Clinical Cases*

*World J Clin Cases* 2021 February 26; 9(6): 1247-1498



## Contents

Thrice Monthly Volume 9 Number 6 February 26, 2021

### EDITORIAL

- 1247 Interactive platform for peer review: A proposal to improve the current peer review system  
*Emile SH*

### MINIREVIEWS

- 1251 Animal models of cathartic colon  
*Meng YY, Li QD, Feng Y, Liu J, Wang EK, Zhong L, Sun QL, Yuan JY*

### ORIGINAL ARTICLE

#### Case Control Study

- 1259 New indicators in evaluation of hemolysis, elevated liver enzymes, and low platelet syndrome: A case-control study  
*Kang SY, Wang Y, Zhou LP, Zhang H*

#### Retrospective Study

- 1271 Analysis of hospitalization costs related to fall injuries in elderly patients  
*Su FY, Fu ML, Zhao QH, Huang HH, Luo D, Xiao MZ*
- 1284 Effect of alprostadil in the treatment of intensive care unit patients with acute renal injury  
*Jia Y, Liu LL, Su JL, Meng XH, Wang WX, Tian C*

#### Clinical Trials Study

- 1293 Etomidate *vs* propofol in coronary heart disease patients undergoing major noncardiac surgery: A randomized clinical trial  
*Dai ZL, Cai XT, Gao WL, Lin M, Lin J, Jiang YX, Jiang X*

#### Observational Study

- 1304 Healthy individuals *vs* patients with bipolar or unipolar depression in gray matter volume  
*Zhang YN, Li H, Shen ZW, Xu C, Huang YJ, Wu RH*
- 1318 Impact of metabolism-related mutations on the heart rate of gastric cancer patients after peritoneal lavage  
*Yuan Y, Yao S, Luo GH, Zhang XY*

### CASE REPORT

- 1329 Efficacy of afatinib in a patient with rare EGFR (G724S/R776H) mutations and amplification in lung adenocarcinoma: A case report  
*He SY, Lin QF, Chen J, Yu GP, Zhang JL, Shen D*

- 1336** Esophageal superficial adenosquamous carcinoma resected by endoscopic submucosal dissection: A rare case report  
*Liu GY, Zhang JX, Rong L, Nian WD, Nian BX, Tian Y*
- 1343** Do medullary thyroid carcinoma patients with high calcitonin require bilateral neck lymph node clearance? A case report  
*Gan FJ, Zhou T, Wu S, Xu MX, Sun SH*
- 1353** Femoral epithelioid hemangioendothelioma detected with magnetic resonance imaging and positron emission tomography/computed tomography: A case report  
*Zhao HG, Zhang KW, Hou S, Dai YY, Xu SB*
- 1359** Noninvasive tools based on immune biomarkers for the diagnosis of central nervous system graft-*vs*-host disease: Two case reports and a review of the literature  
*Lyu HR, He XY, Hao HJ, Lu WY, Jin X, Zhao YJ, Zhao MF*
- 1367** Periodontally accelerated osteogenic orthodontics with platelet-rich fibrin in an adult patient with periodontal disease: A case report and review of literature  
*Xu M, Sun XY, Xu JG*
- 1379** Subtalar joint pigmented villonodular synovitis misdiagnosed at the first visit: A case report  
*Zhao WQ, Zhao B, Li WS, Assan I*
- 1386** Wilson disease — the impact of hyperimmunity on disease activity: A case report  
*Stremmel W, Longerich T, Liere R, Vacata V, van Helden J, Weiskirchen R*
- 1394** Unexplained elevation of erythrocyte sedimentation rate in a patient recovering from COVID-19: A case report  
*Pu SL, Zhang XY, Liu DS, Ye BN, Li JQ*
- 1402** Thoracic pyogenic infectious spondylitis presented as pneumothorax: A case report  
*Cho MK, Lee BJ, Chang JH, Kim YM*
- 1408** Unilateral pulmonary hemorrhage caused by negative pressure pulmonary edema: A case report  
*Park HJ, Park SH, Woo UT, Cho SY, Jeon WJ, Shin WJ*
- 1416** Osseous Rosai-Dorfman disease of tibia in children: A case report  
*Vithran DTA, Wang JZ, Xiang F, Wen J, Xiao S, Tang WZ, Chen Q*
- 1424** Abdominopelvic leiomyoma with large ascites: A case report and review of the literature  
*Wang YW, Fan Q, Qian ZX, Wang JJ, Li YH, Wang YD*
- 1433** Unusual presentation of granulomatosis with polyangiitis causing periaortitis and consequent subclavian steal syndrome: A case report  
*Cho U, Kim SK, Ko JM, Yoo J*
- 1439** Postoperative discal pseudocyst and its similarities to discal cyst: A case report  
*Fu CF, Tian ZS, Yao LY, Yao JH, Jin YZ, Liu Y, Wang YY*

- 1446** Treatment of oral lichen planus by surgical excision and acellular dermal matrix grafting: Eleven case reports and review of literature  
*Fu ZZ, Chen LQ, Xu YX, Yue J, Ding Q, Xiao WL*
- 1455** Nonalcoholic fatty liver disease as a risk factor for cytomegalovirus hepatitis in an immunocompetent patient: A case report  
*Khiatah B, Nasrollah L, Covington S, Carlson D*
- 1461** Early reoccurrence of traumatic posterior atlantoaxial dislocation without fracture: A case report  
*Sun YH, Wang L, Ren JT, Wang SX, Jiao ZD, Fang J*
- 1469** Intrahepatic cholangiocarcinoma is more complex than we thought: A case report  
*Zeng JT, Zhang JF, Wang Y, Qing Z, Luo ZH, Zhang YL, Zhang Y, Luo XZ*
- 1475** Congenital hepatic fibrosis in a young boy with congenital hypothyroidism: A case report  
*Xiao FF, Wang YZ, Dong F, Li XL, Zhang T*
- 1483** Polidocanol sclerotherapy for multiple gastrointestinal hemangiomas: A case report  
*Yao H, Xie YX, Guo JY, Wu HC, Xie R, Shi GQ*
- 1490** Gastrointestinal stromal tumor with multisegmental spinal metastases as first presentation: A case report and review of the literature  
*Kong Y, Ma XW, Zhang QQ, Zhao Y, Feng HL*

**ABOUT COVER**

Editorial Board Member of *World Journal of Clinical Cases*, Dr. Quach is an Associate Professor of Gastroenterology at the University of Medicine and Pharmacy at Hochiminh City, Viet Nam, where he received his MD in 1997 and his PhD in 2011. Dr. Quach has published more than 100 reviews and original papers in local and international journals. He has received several awards, including Outstanding Presentation at the Biannual Scientific Congress of Vietnamese Nationwide Medical Schools, Medal of Creativeness from the Vietnamese Central Youth League, etc. Currently, he serves as a Vice President of the Vietnam Association of Gastroenterology and Secretary General of the Vietnam Federation for Digestive Endoscopy. (L-Editor: Filipodia)

**AIMS AND SCOPE**

The primary aim of *World Journal of Clinical Cases* (*WJCC*, *World J Clin Cases*) is to provide scholars and readers from various fields of clinical medicine with a platform to publish high-quality clinical research articles and communicate their research findings online.

*WJCC* mainly publishes articles reporting research results and findings obtained in the field of clinical medicine and covering a wide range of topics, including case control studies, retrospective cohort studies, retrospective studies, clinical trials studies, observational studies, prospective studies, randomized controlled trials, randomized clinical trials, systematic reviews, meta-analysis, and case reports.

**INDEXING/ABSTRACTING**

The *WJCC* is now indexed in Science Citation Index Expanded (also known as SciSearch®), Journal Citation Reports/Science Edition, Scopus, PubMed, and PubMed Central. The 2020 Edition of Journal Citation Reports® cites the 2019 impact factor (IF) for *WJCC* as 1.013; IF without journal self cites: 0.991; Ranking: 120 among 165 journals in medicine, general and internal; and Quartile category: Q3. The *WJCC*'s CiteScore for 2019 is 0.3 and Scopus CiteScore rank 2019: General Medicine is 394/529.

**RESPONSIBLE EDITORS FOR THIS ISSUE**

Production Editor: Ji-Hong Lin; Production Department Director: Xiang Li; Editorial Office Director: Jin-Lai Wang.

**NAME OF JOURNAL**

*World Journal of Clinical Cases*

**ISSN**

ISSN 2307-8960 (online)

**LAUNCH DATE**

April 16, 2013

**FREQUENCY**

Thrice Monthly

**EDITORS-IN-CHIEF**

Dennis A Bloomfield, Sandro Vento, Bao-Gan Peng

**EDITORIAL BOARD MEMBERS**

<https://www.wjgnet.com/2307-8960/editorialboard.htm>

**PUBLICATION DATE**

February 26, 2021

**COPYRIGHT**

© 2021 Baishideng Publishing Group Inc

**INSTRUCTIONS TO AUTHORS**

<https://www.wjgnet.com/bpg/gerinfo/204>

**GUIDELINES FOR ETHICS DOCUMENTS**

<https://www.wjgnet.com/bpg/GerInfo/287>

**GUIDELINES FOR NON-NATIVE SPEAKERS OF ENGLISH**

<https://www.wjgnet.com/bpg/gerinfo/240>

**PUBLICATION ETHICS**

<https://www.wjgnet.com/bpg/GerInfo/288>

**PUBLICATION MISCONDUCT**

<https://www.wjgnet.com/bpg/gerinfo/208>

**ARTICLE PROCESSING CHARGE**

<https://www.wjgnet.com/bpg/gerinfo/242>

**STEPS FOR SUBMITTING MANUSCRIPTS**

<https://www.wjgnet.com/bpg/GerInfo/239>

**ONLINE SUBMISSION**

<https://www.f6publishing.com>

# Unusual presentation of granulomatosis with polyangiitis causing periaortitis and consequent subclavian steal syndrome: A case report

Uiju Cho, Sung-Kyung Kim, Jeong Min Ko, Jinyoung Yoo

**ORCID number:** Uiju Cho 0000-0002-6229-8418; Sung-Kyung Kim 0000-0002-2742-6410; Jeong Min Ko 0000-0002-8996-5786; Jinyoung Yoo 0000-0002-5053-1489.

**Author contributions:** Cho U and Yoo J performed the pathologic examination and diagnosed the patient's case, reviewed the literature and contributed to manuscript drafting; Kim SK was the patient's physician and contributed to manuscript drafting; Ko J performed the radiologic analysis of the patient's case and contributed to manuscript drafting; all authors issued final approval for the version to be submitted.

**Informed consent statement:**

Written informed consent was obtained from the patient.

**Conflict-of-interest statement:** The authors declare that there is no conflict of interest.

**CARE Checklist (2016) statement:**

The authors have read the CARE Checklist (2016), and the manuscript was prepared and revised according to the CARE Checklist (2016).

**Open-Access:** This article is an open-access article that was selected by an in-house editor and

**Uiju Cho**, Department of Pathology, St. Vincent's Hospital, College of Medicine, The Catholic University of Korea, Seoul 06591, South Korea

**Sung-Kyung Kim**, Department of Internal Medicine, St. Vincent's Hospital, College of Medicine, The Catholic University of Korea, Seoul 06591, South Korea

**Jeong Min Ko**, Department of Radiology, St. Vincent's Hospital, College of Medicine, The Catholic University of Korea, Seoul 06591, South Korea

**Jinyoung Yoo**, Department of Hospital Pathology, St. Vincent's Hospital, College of Medicine, The Catholic University of Korea, Seoul 06591, South Korea

**Corresponding author:** Jinyoung Yoo, MD, PhD, Doctor, Professor, Department of Hospital Pathology, St. Vincent's Hospital, College of Medicine, The Catholic University of Korea, Jidong 93, Paldal-gu, Seoul 06591, South Korea. [jinyyoo@catholic.ac.kr](mailto:jinyyoo@catholic.ac.kr)

## Abstract

### BACKGROUND

Granulomatosis with polyangiitis (GPA) is a rare autoimmune disease that involves small-to-medium-sized vessels and forms necrotizing vasculitis with granulomatous inflammation. The formation of a large vessel lesion in GPA patients has been scarcely reported, and it can cause confusion in the diagnosis.

### CASE SUMMARY

A 27-year-old man presented with mild left-sided pleuritic chest pain that started one year prior. An imaging study revealed up to 2.5 cm-sized two irregular nodular consolidation nodule in the left lower lobe. Both nodules showed central necrosis. Also, there was a periaortic mass occluding the branching porting of the subclavian artery. He had positive anti-neutrophil cytoplasmic antibodies (ANCA), but myeloperoxidase-ANCA and proteinase 3-ANCA were negative. The patient also developed symptoms of subclavian vein syndrome during the follow-up. Wedge resection of the lung revealed necrotizing vasculitis, destructive parenchymal abscess and surrounding granuloma, and therefore diagnosed of GPA. The patient started on methotrexate and steroid therapy with a relief of symptomatic.

### CONCLUSION

fully peer-reviewed by external reviewers. It is distributed in accordance with the Creative Commons Attribution NonCommercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>

**Manuscript source:** Unsolicited manuscript

**Specialty type:** Medicine, research and experimental

**Country/Territory of origin:** South Korea

**Peer-review report's scientific quality classification**

Grade A (Excellent): 0  
Grade B (Very good): B  
Grade C (Good): 0  
Grade D (Fair): 0  
Grade E (Poor): 0

**Received:** October 7, 2020

**Peer-review started:** October 7, 2020

**First decision:** December 21, 2020

**Revised:** December 30, 2020

**Accepted:** January 8, 2021

**Article in press:** January 8, 2021

**Published online:** February 26, 2021

**P-Reviewer:** Alshewered ASH

**S-Editor:** Zhang L

**L-Editor:** A

**P-Editor:** Ma YJ



Here, we present an unusual manifestation of GPA with periaortitis and consequent subclavian steal syndrome, which has never been previously described. This case alerts us that we should include GPA in the differential diagnosis of large vessel vasculitis as well as subclavian steal syndrome.

**Key Words:** Granulomatosis with polyangiitis; Wegener granulomatosis; Systemic vasculitis; Subclavian steal syndrome; Periaortitis; Case report

©The Author(s) 2021. Published by Baishideng Publishing Group Inc. All rights reserved.

**Core Tip:** Granulomatosis with polyangiitis (GPA) is a systemic process characterized by necrotizing vasculitis and granulomatous inflammation. The large vessel involvement by GPA is not only rare but also causes major diagnostic difficulty. We present a rare case of GPA, showing both lung and periaortic lesion in a male patient. Periaortic lesion of the patient caused vascular occlusion and consequent subclavian vein syndrome. On pathologic examination, the lung nodules had typical histologic features of GPA. This case reminds us that the GPA should not be excluded from the diagnosis because of a large vascular lesion present in an otherwise suspicious setting.

**Citation:** Cho U, Kim SK, Ko JM, Yoo J. Unusual presentation of granulomatosis with polyangiitis causing periaortitis and consequent subclavian steal syndrome: A case report. *World J Clin Cases* 2021; 9(6): 1433-1438

**URL:** <https://www.wjgnet.com/2307-8960/full/v9/i6/1433.htm>

**DOI:** <https://dx.doi.org/10.12998/wjcc.v9.i6.1433>

## INTRODUCTION

Granulomatosis with polyangiitis (GPA), formerly called Wegener granulomatosis, is a systemic process characterized by necrotizing vasculitis and granulomatous reaction involving small- to medium-sized vessels. It is rare for the GPA to present with lesions in the large blood vessels. Moreover, GPA as a cause of periaortitis and consequent subclavian steal syndrome has never been described. Here, we present a GPA case showing such rare findings that evoke physicians of a new aspect of the disease.

## CASE PRESENTATION

### Chief complaints

A 27-year-old man presented with mild left-sided pleuritic chest pain.

### History of present illness

The patient's symptom started one year prior, and it was aggravated by breathing and laughing. He also had occasional substernal area discomfort.

### History of past illness

He had no past medical history except for allergic rhinitis.

### Personal and family history

He had no family history. He had a smoking history of 12 pack-years, and he was a social drinker.

### Physical examination

The patient's lungs were clear on auscultation, and the other physical examination findings, including skin, neurologic and otorhinolaryngologic examinations, were within normal limits.



### Laboratory examinations

Laboratory studies revealed an elevated level of C-reactive protein but normal levels of creatinine and blood urea nitrogen. White blood cell count was within the normal limit. Antinuclear antibodies were positive, with a low titer of 1:40. Enzyme-linked immunosorbent assay for anti-neutrophil cytoplasmic antibodies (ANCA) was positive, but myeloperoxidase-ANCA and proteinase 3-ANCA were negative. Other autoimmune antibodies were not found. The urine analysis results were within normal limits.

### Imaging examinations

Chest computed tomography revealed a peribronchial, 2.5 cm-sized irregular nodular consolidation nodule in the left lower lobe (Figure 1A). Another solid nodule was located adjacent to the main nodule. Both nodules showed central necrosis. Vascular wall thickening from the distal portion of the aortic arch to the proximal descending aorta was also noted (Figure 1B and C). This periaortic lesion occluded the branching portion of the left subclavian artery. The bronchial arteries had no occlusions.

### Further diagnostic work-up

Transbronchial lung biopsy showed nonspecific interstitial inflammation and fibrosis. Tests for tuberculosis, including the polymerase chain reaction and culture, were all negative. An open lung biopsy was recommended, but the patient refused. During follow-up, the patient complained of exercise-induced arm fatigue and paresthesia. The blood pressure of the left brachial artery was 105/65 mmHg, which was more than 15 mmHg lower than that of the right brachial artery blood pressure (129/74 mmHg). Continuous Doppler ultrasonography showed reversed flow in the left vertebral artery (Figure 1D). Brain magnetic resonance imaging and angiography also revealed occlusion of the left proximal subclavian artery along with retrograde filling of the distant subclavian artery. Subclavian steal syndrome was diagnosed based on the patient's symptoms and imaging studies. The patient underwent lung wedge resection.

On gross examination, the nodules in the lung parenchymal consolidation showed irregular edges and pale-brown cut surfaces (Figure 2A). Histologically, there was bronchocentric chronic inflammation, extensive geographic parenchymal necrosis, granulomatous inflammation with vasculitis. The areas of necrosis featured microabscesses and large basophilic zones of geographical necrosis with serpiginous borders. Epithelioid macrophages were palisading around the borders of the necrosis, and a few multinucleated giant cells were also present. Vasculitis involved the bronchial arteries and small vessels. Lymphocytes infiltrated the vascular wall and destroyed the elastic laminae (Figure 2B-F). Special staining and culture techniques showed no fungal or bacterial infection, and all the tests for Mycobacteria, including Ziehl-Neelsen stain, culture and polymerase chain reaction, were also negative.

---

## FINAL DIAGNOSIS

---

The pathology findings were consistent with granulomatosis with polyangiitis.

---

## TREATMENT

---

The patient was started on methotrexate and steroid therapy.

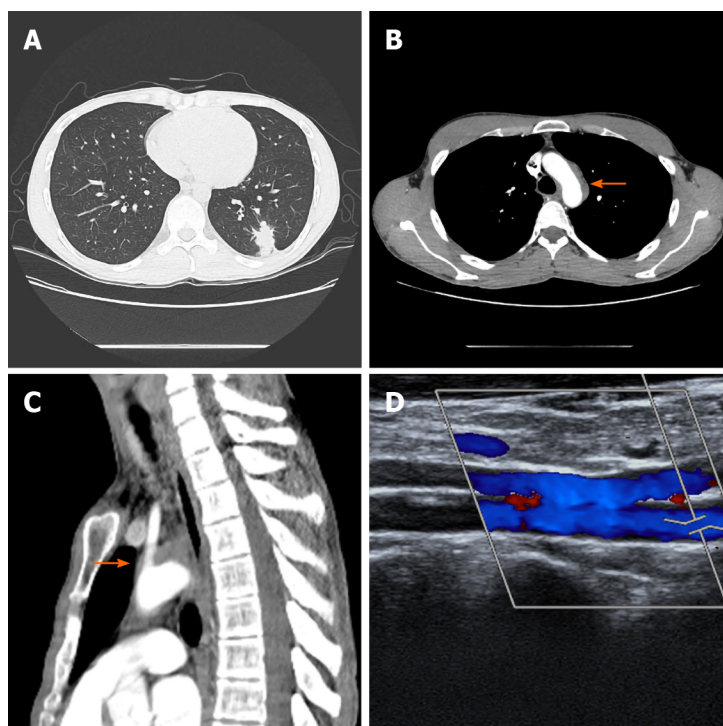
---

## OUTCOME AND FOLLOW-UP

---

During 11 mo follow up, the patient did well with a relief of chest pain. Fatigue and paresthesia on the left arm did not worsen but remained. Twelve months after the lung wedge resection, the patient developed headache and aphasia and visited the hospital emergency room. Computed tomography angiography and magnetic resonance imaging revealed middle cerebral artery territory infarct of the left brain. The left proximal carotid coronary artery, proximal subclavian artery, and left inferior division of the middle cerebral artery M2 segment were occluded. Treatments of the acute brain infarct, *e.g.*, aspirin, clopidogrel, rosuvastatin, low-molecular-weight heparin, were





**Figure 1 Initial computed tomography scans.** A: Chest computed tomography scan with contrast enhancement showing a nodule abutting the pleura in the left lower lobe of the lung; B and C: Computed tomography scan showing a heterogeneously enhancing soft tissue dense lesion partially occluding the aortic arch; and D: Doppler ultrasonography for the evaluation of subclavian steal syndrome demonstrated a reverse flow in the left vertebral artery.

added to the methotrexate and steroid therapy. The patient was discharged after the symptomatic relief and had been following up eventless.

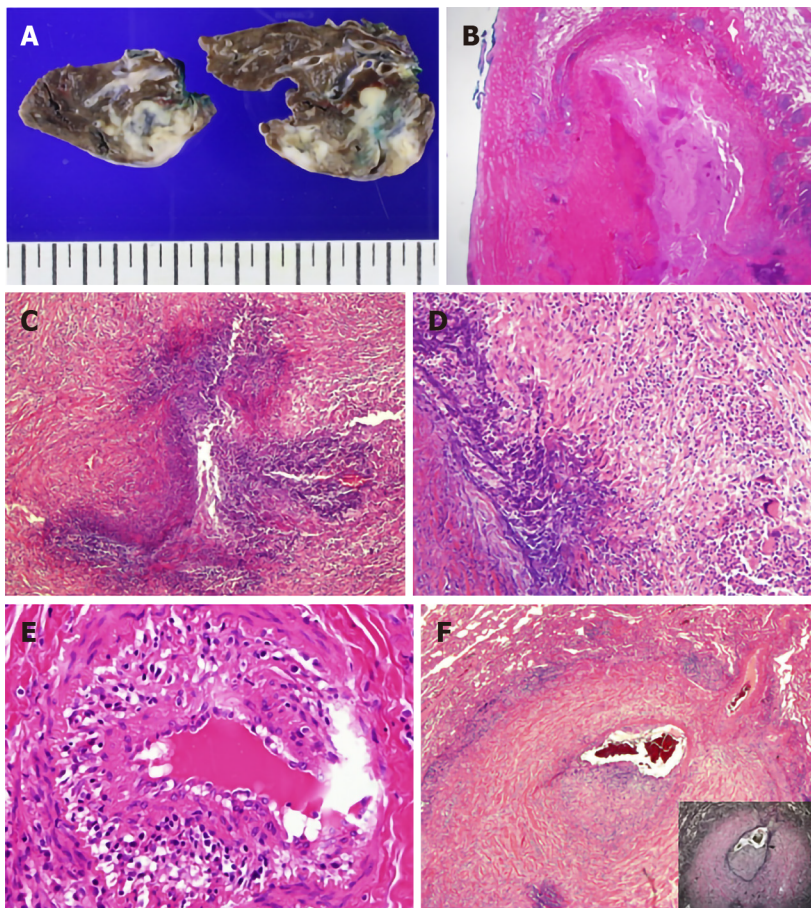
## DISCUSSION

Many pulmonary conditions can cause vasculitis, but most are secondary conditions, such as necrotizing granulomatous infection. Granulomatosis with polyangiitis<sup>[1]</sup>, microscopic polyangiitis, and eosinophilic granulomatous polyangiitis are the few primary idiopathic vasculitis diseases that affect the lung.

Because of the patient's subclavian steal syndrome, the major differential diagnosis, in this case, was Takayasu arteritis (TA). Subclavian steal syndrome is a syndrome with symptoms like exercise-induced arm pain, fatigue, and paresthesia caused by flow reversal in the vertebral artery ipsilateral to stenosis of the subclavian artery<sup>[2]</sup>. Atherosclerosis is the most common cause of subclavian steal syndrome. Other conditions, such as TA, compression of the subclavian artery in the thoracic outlet, and aortic or heart surgery complications, may also lead to subclavian steal syndrome<sup>[3]</sup>.

Lung aberrations in TA are stenosis of the main pulmonary artery branch associated with the parenchymal ischemic change<sup>[4]</sup>. Vasculitis with geographic necrotizing granuloma is not a feature of TA<sup>[5]</sup>. Besides, concentric wall thickening of the large vessel on imaging is the typical finding in TA<sup>[6]</sup>, but in our case, the aortic lesion was not concentric. Thus, the patient's overall pathologic and radiologic findings suggested against this form of vasculitis.

Involvement of large vessels by GPA is a rare phenomenon, and less than 40 cases have been reported in the literature to date<sup>[7-9]</sup>. Although the most frequent site of involvement is an abdominal aorta, there have been some case reports of GPA presenting in the thoracic aorta as periaortitis or aortic aneurism<sup>[8]</sup>. Histologic examination of the large vessel lesions demonstrated necrotizing granulomatous inflammation of the vessel wall itself or periaortic tissue extending to the aortic wall<sup>[8,10]</sup>. The pathogenic mechanism of the large vessel involvement by GPA is yet to be elucidated. ANCA may play a role in this process by inducing vasculitis in the vasa vasorum<sup>[9,11]</sup>. Large vessel involvement may also be explained by polyangiitis overlap syndrome or a novel clinical overlap syndrome<sup>[12]</sup>. However, histologic and clinical observations in the previous studies suggest that large vessel involvement belongs to



**Figure 2 Pathology findings in the lung.** A: Gross sections of the wedged resected lung show pale-brown or ivory colored, irregularly shaped nodules. Areas of necrosis are grossly identifiable; B: The lung displayed extensive geographic necrosis areas with the total destruction of the normal lung parenchyma. Pleural plaque-like fibrosis was also present [Hematoxylin and eosin (HE) stain,  $\times 12.5$ ]; C: Smaller serpiginous abscesses were seen (HE stain,  $\times 100$ ); D: Granulomatous inflammation with a few multinucleated giant cells surrounded the necrotic area (HE stain,  $\times 200$ ); E: Some of the small vessels had many inflammatory cells infiltrating into the vessel wall (vasculitis) (HE stain,  $\times 400$ ); and F: Arterioles were markedly affected by vasculitis. The vessel wall was destroyed, and the lumen was partially occluded by infiltrating inflammatory cells and fibrosis. Cicatricial fibrosis surrounded the vessel wall (HE stain, inset: elastic stain,  $\times 40$ ).

the spectrum of ANCA-associated vasculitis rather than overlap with other large vessel vasculitides<sup>[9,11,13]</sup>.

## CONCLUSION

Interestingly, our patient had a GPA with associated periaortitis, which led to the consequent subclavian steal syndrome and brain infarct, which has never been previously described. Such unusual presentations can result in delayed diagnosis since GPA is known to involve not large vessels but small- to medium-sized vessels. This is an instructive case showing that GPA should be included in the differential diagnosis of large vessel vasculitis as well as subclavian steal syndrome.

## REFERENCES

- 1 **Lazim QJ**, Atrah SSG, Mutlag KJ, Alhilfi HSQ, Fahad AM, Alshewered AS. Granulomatosis (Wegener's granulomatosis) with polyangiitis presented as pulmonary manifestation: a case report. *Respirol Case Rep* 2020; **8**: e00674 [PMID: 33082957 DOI: 10.1002/rcr2.674]
- 2 **Fisher CM**. A new vascular syndrome-"the subclavian steal. *N Engl J Med* 1961; **265**: 912-913 [DOI: 10.1056/NEJM196111022651812]
- 3 **Osiro S**, Zurada A, Gielecki J, Shoja MM, Tubbs RS, Loukas M. A review of subclavian steal syndrome with clinical correlation. *Med Sci Monit* 2012; **18**: RA57-RA63 [PMID: 22534720 DOI: 10.12659/msm.882721]
- 4 **He Y**, Lv N, Dang A, Cheng N. Pulmonary Artery Involvement in Patients with Takayasu Arteritis. *J*

- Rheumatol* 2020; **47**: 264-272 [PMID: [31092716](#) DOI: [10.3899/jrheum.190045](#)]
- 5 **Nakajima N.** Takayasu arteritis: consideration of pulmonary involvement. *Ann Vasc Dis* 2008; **1**: 7-10 [PMID: [23555332](#) DOI: [10.3400/avd.AVDedit00107](#)]
- 6 **Zhu FP**, Luo S, Wang ZJ, Jin ZY, Zhang LJ, Lu GM. Takayasu arteritis: imaging spectrum at multidetector CT angiography. *Br J Radiol* 2012; **85**: e1282-e1292 [PMID: [23175494](#) DOI: [10.1259/bjr/25536451](#)]
- 7 **Milkowska-Dymanowska J**, Laskowska P, Rzuczkowski M, Białas AJ, Piotrowski WJ, Górski P. Untypical Manifestations of Granulomatosis with Polyangiitis—A Review of the Literature. *SN Compr Clin Med* 2019; **1**: 616-626 [DOI: [10.1007/s42399-019-00083-4](#)]
- 8 **Ozaki T**, Maeshima K, Kiyonaga Y, Torigoe M, Imada C, Hamasaki H, Haranaka M, Ishii K, Shibata H. Large-vessel involvement in granulomatosis with polyangiitis successfully treated with rituximab: A case report and literature review. *Mod Rheumatol* 2017; **27**: 699-704 [PMID: [25736357](#) DOI: [10.3109/14397595.2015.1021950](#)]
- 9 **Chirinos JA**, Tamariz LJ, Lopes G, Del Carpio F, Zhang X, Milikowski C, Lichtstein DM. Large vessel involvement in ANCA-associated vasculitides: report of a case and review of the literature. *Clin Rheumatol* 2004; **23**: 152-159 [PMID: [15045631](#) DOI: [10.1007/s10067-003-0816-0](#)]
- 10 **Blockmans D**, Baeyens H, Van Loon R, Lauwers G, Bobbaers H. Periaortitis and aortic dissection due to Wegener's granulomatosis. *Clin Rheumatol* 2000; **19**: 161-164 [PMID: [10791632](#) DOI: [10.1007/s100670050038](#)]
- 11 **Carels T**, Verbeken E, Blockmans D. p-ANCA-associated periaortitis with histological proof of Wegener's granulomatosis: case report. *Clin Rheumatol* 2005; **24**: 83-86 [PMID: [15565392](#) DOI: [10.1007/s10067-004-0998-0](#)]
- 12 **Ohta H**, Shirai S, Nasu K, Tei M, Kambara H, Ono T, Shintaku M. Ga-67 uptake in the aorta in Wegener's granulomatosis: overlap with Takayasu's arteritis? *Clin Nucl Med* 1998; **23**: 859-860 [PMID: [9858310](#) DOI: [10.1097/00003072-199812000-00023](#)]
- 13 **Zoma AA.** Clinical overlap in proximal artery stenosis: two disorders or one? *Lancet* 2016; **387**: 2350 [PMID: [27302275](#) DOI: [10.1016/S0140-6736\(16\)30524-4](#)]



Published by **Baishideng Publishing Group Inc**  
7041 Koll Center Parkway, Suite 160, Pleasanton, CA 94566, USA

**Telephone:** +1-925-3991568

**E-mail:** [bpgoffice@wjgnet.com](mailto:bpgoffice@wjgnet.com)

**Help Desk:** <https://www.f6publishing.com/helpdesk>

<https://www.wjgnet.com>

