

criteria are still used today and they only include symptoms from the temporal artery. There are not established criteria for the involvement of other large vessels. Prospective studies using CTA show a rather common participation of the aorta though, either in the form of thickened wall or presented as an aneurysm, at a percentage of 45%-65% [3,4]. Studies using Fludeoxyglucose-Positron Emission Tomography (FDG-PET) have detected involvement of the aorta in >50% of patients with temporal arteritis [5]. Symptoms of the aortic involvement are not always present. Most often the patient complains about lower limb claudication. At physical examination vascular bruits and aortic regurgitation murmur can be noted. Pain in the chest or the abdomen can indicate a latent aneurysm of the aorta. Other vessels which are less commonly affected are the brachiocephalic trunk, the carotids, subclavian and axillary arteries, and even rarer there is involvement of the splanchnic, renal, and femoral arteries [4].

In general, patients with temporal arteritis don't seem to have increased risk of death, when compared with the general population mortality rates [6]. When GCA is affecting large vessels, though, and especially when aorta is involved, it is associated with increased mortality [7]. The development of an aortic aneurysm is the most serious complication, which can result in rupture and death. Another serious complication is the permanent loss of vision that occurs when ophthalmic and retinal arteries are affected. In milder cases, transient loss of vision, also known as amaurosis fugax can occur. Critical limb ischemia has also been reported but it is rare. There are few reports of acute myocardial infarction due to inflammation of the coronary arteries in the context of GCA [8,9].

Even fewer are the reports of pericardial effusion that can correlate with GCA [10]. Pericarditis is the inflammation of the pericardial sac, caused by either infectious or immunologic disorders. Some researchers have proposed that GCA occurs in a spectrum of clinical manifestations, ranging from isolated cranial involvement, cranial and large vessel involvement and large vessel involvement without cranial participation [11]. Regarding the case of pericardial effusion, the pathophysiology is unknown. To the best of authors' knowledge, only a few case reports have described this condition and it is not always associated with aortic manifestations. It is only logical to assume, that vasculitis of the coronary arteries is present. Other possible pathophysiological mechanisms include inflammatory cytokines, immunological reactions, and immune complexes deposition in the pericardium, causing inflammatory interstitial lesion with or without granulomas [12]. The exact prevalence of this implication is unknown but Zenone et al. [13] reported a prevalence of 3,5% of pericardial effusion in a retrospective study of 114 patients with GCA. Tiosano et al. [14] have found that

pericarditis is independently associated with GCA, especially in patients younger than 70-year old [14].

It is well known that Polymyalgia Rheumatica is a disease that can present in the context of giant cell arteritis and it comes with constitutional symptoms of low-grade fever and fatigue and achiness and stiffness on the shoulders and the pelvic girdle. The incidence of pericardial effusion in patients with polymyalgia rheumatica and a negative temporal artery biopsy is unknown, but reports of such cases exist [15].

Conclusion

In patients with biopsy diagnosed GCA, it is not a common practice to examine the heart or the aorta. Examining the patient for pericardial rub sound and aortic bruits should be regular in these patients. Given that the implications of these conditions might be very severe, if untreated, we suggest early checking with CT angiography or PET CT scan and heart ECHO. Contrariwise, in a patient with pericarditis, and a negative testing for common causes, before attributing this condition to unknown or idiopathic origins, the physician should consider checking for GCA in the differential diagnosis.

What is new?

Pericardial effusion is rare but serious in the context of Giant Cell Arteritis. The authors want to point out the significance of early checking with CT angiography or PET CT scan and heart ECHO when Giant Cell Arteritis is diagnosed to rule out pericardial effusion.

List of Abbreviations

GCA Giant cell arteritis

Consent for publication

Written informed consent was obtained from the patient.

Ethical approval

Ethical approval is not required at our institution for publishing an anonymous case report.

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References

1. Kermani TA, Schäfer VS, Crowson CS, Hunder GG, Gabriel SE, Matteson EL, et al. Increase in age at onset of giant cell arteritis: a population-based study. *Ann Rheum Dis.* 2010;69:780–781. <https://doi.org/10.1136/ard.2009.111005>
2. Hunder GG, Bloch DA, Michel BA, Stevens MB, Arend WP, Calabrese LH, et al. The American college of Rheumatology 1990 criteria for the classification of Giant Cell Arteritis. *Arthritis Rheum.* 1990;33. <https://doi.org/10.1002/art.1780330810>

3. Agard C, Barrier JH, Dupas B, Ponge T, Mahr A, Fradet G, et al. Aortic involvement in recent-onset giant cell (temporal) arteritis: a case-control prospective study using helical aortic computed tomodensitometric scan. *Arthritis Rheum.* 2008;59:670–6. <https://doi.org/10.1002/art.23577>
4. Prieto-González S, Arguis P, García-Martínez A, Espígol-Frigolé G, Tavera-Bahillo I, Butjosa M, et al. Large vessel involvement in biopsy-proven giant cell arteritis: prospective study in 40 newly diagnosed patients using CT angiography. *Ann Rheum Dis.* 2012;71:1170–6. <https://doi.org/10.1136/annrheumdis-2011-200865>
5. Meller J., Strutz F, Siefker U, Scheel A, Sahlmann CO, Lehmann K, et al. Early diagnosis and follow-up of aortitis with [18F]FDG PET and MRI. *Eur J Nucl Med Mol Imaging.* 2003;30:730–6. <https://doi.org/10.1007/s00259-003-1144-y>
6. Lensen KD, Voskuyl AE, Comans EF, van der Laken CJ, Smulders YM. Extracranial giant cell arteritis: a narrative review. *Netherlands J Med.* 2013;74:182–92.
7. Kermani TA, Warrington KJ, Crowson CS, Ytterberg SR, Hunder GG, Gabriel SE, et al. Large-vessel involvement in giant cell arteritis: a population-based cohort study of the incidence-trends and prognosis. *Ann Rheum Dis.* 2012;72(12):1989–94. <https://doi.org/10.1136/annrheumdis-2012-202408>
8. Hupp SL, Nelson GA, Zimmerman LE. Generalized giant-cell arteritis with coronary artery involvement and myocardial infarction. *Arch Ophthalmol.* 1990;108(10):1385–7. <https://doi.org/10.1001/archophth.1990.01070120031015>
9. Godoy P, Araújo Sde A, Paulino E Jr, Lana-Peixoto MA. Coronary giant cell arteritis and acute myocardial infarction. *Arquivos Brasileiros de Cardiologia.* 2007.88(4):84–7. <https://doi.org/10.1590/S0066-782X2007000400027>
10. Tasliyurt T, Sivgin H, Bekar L, Sahin S, Uzun Kaya S, Koseoglu RD, et al. A rare cause of pericardial effusion: giant cell arteritis. *Case Rep Rheum.* 2014. <https://doi.org/10.1155/2014/424295>
11. Matthew J. Koster, Eric L. Matteson and Kenneth J. Warrington. Large-vessel giant cell arteritis: diagnosis, monitoring and management. *Rheumatology (Oxford).* 2018;57:ii32ii42. <https://doi.org/10.1093/rheumatology/kex424>
12. Guindon A, Rossi P, Bagneres D, Aissi K, Demoux AL, Bonin-Guillaume S, et al. [Pericarditis: a giant cell arteritis manifestation]. *Revue de Médecine Interne.* 2007;28(5):326–31. <https://doi.org/10.1016/j.revmed.2007.01.023>
13. Zenone T, Puget M. Pericardial effusion and giant cell arteritis. *Rheumatol Int.* 2014;34(10):1465–9. <https://doi.org/10.1007/s00296-014-2958-6>
14. Tiosano S, Adler Y, Azrielant S, Yavne Y, Gendelman O, Ben-Ami Shor D, et al. Pericarditis among giant cell arteritis patients: from myth to reality. *Clin Cardiol.* 2018.41:623–7. <https://doi.org/10.1002/clc.22927>
15. Calvo E, Becerra E, López-Longo FJ, Cabrera FJ, Carreño L, Paravisini A, et al. Pericardial tamponade in a patient with polymyalgia rheumatica. *Clin Exp Rheumatol.* 2009;27(1):83.

Summary of the case

1	Patient (gender, age)	Female, 72
2	Final diagnosis	Pericardial effusion in the context of Giant Cell Arteritis
3	Symptoms	Fever, Headache
4	Medications	Methylprednisolone, calcium, vitamin D, proton pump inhibitors and biphosphonate agents
5	Clinical procedure	Chest CT scan
6	Specialty	Internal Medicine