CASE REPORT

Immunoglobulin G4-related constrictive pericarditis and the importance of a thorough workup: a case report

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Abstract

Background: Constrictive pericarditis remains a problematic diagnosis and a thorough investigation is critical. Among possible aetiologies, immunoglobulin-G4 (IgG4)-related pericardial disease is an unusual cause of pericardial constriction. We report a challenging diagnostic case of pericardial constriction due to IgG4-related disease.

Case presentation: A 68-year old male with a history of inferior myocardial infarction with right ventricle (RV) involvement was thrice-hospitalized due to marked ascites and peripheral oedema. Systemic congestion was initially attributed to RV dysfunction due to previous infarction. Yet, at the final admission, a re-assessment echocardiogram followed by cardiac computed tomography, magnetic resonance and right heart catheterization raised a possible diagnosis of constrictive pericarditis with a finding of abnormal pulmonary venous return. Patient therefore underwent pericardiectomy and surgical correction of pulmonary venous return. Pericardium histology revealed an IgG4-related pericardial constriction. Patient was later discharged on corticosteroids with marked symptomatic improvement.

Conclusion: IgG4-related disease remains a rare cause of pericardium constriction while also presenting a challenging diagnosis in everyday clinical practice. This case exemplifies the difficulties faced by clinicians when reviewing a possible case of constrictive pericarditis, while highlighting the importance of a multimodality assessment.

Keywords: Case report, Constrictive pericarditis, Immunoglobulin G4-related disease, Right heart failure

Background

ImmunoglobulinG4 (IgG4)-related disease is a systemic syndrome with multiorgan lymphoplasmacytic infiltration. Yet, the frequency of IgG4-related pericarditis remains unknown. Given its non-specific presentation, the correct diagnosis may be delayed, especially in those with other causes of right heart failure (RHF). We present a challenging diagnostic case of constrictive pericarditis due to IgG4-related disease in accordance with the CARE reporting checklist.

Case presentation

A 68-year old male with a previous inferior myocardial infarction (MI) with right ventricle (RV) involvement was thrice-admitted due to abdominal distension and peripheral oedema. He denied fever, dyspnoea or known hepatic or renal disease. Vital signs were stable and physical examination identified marked ascites and peripheral oedema. Blood tests revealed an NT-proBNP of 1525 pg/mL (reference: < 125 pg/mL), creatinine 1.25 mg/dL (reference: < 0.8–1.3 mg/dL), alanine aminotransferase 30 U/L.
possible diagnosis of constrictive pericarditis with an inci-
pressure-waveform showing a “dip and plateau” pattern. A
Doppler imaging of 10 cm/s). Cardiac computed tomog-
16 mm, peak tricuspid annular systolic velocity by tissue-
dysfunction (tricuspid annular plane excursion of
also showed dilated inferior vena cava and borderline RV
mitral inflow E velocity was identified. Echocardiogram
infiltration (Fig. 2A–D).

As such, IgG4 immunochemistry study was performed,
over, focal fibrosis with a storiform pattern was observed.
Dense lymphoplasmacytic infiltration with numerous plasmocytes. More-
trates and focal fibrosis with storiform pattern are key
histological diagnosis. Dense lymphoplasmacytic infiltr-
and focal fibrosis with storiform pattern was observed.
As such, IgG4 immunochemistry study was performed,
which confirmed an extensive IgG4-positive plasma-cell
infiltration (Fig. 2A–D).

A diagnosis of IgG4-related constrictive pericarditis with concomitant anomalous pulmonary venous return was therefore established. The patient was started on corticosteroids and discharged with marked symptom improvement.

Discussion and conclusions
Multimodality imaging is often required in diagnosing constrictive pericarditis. Several concurring RHF aetiologies were presented during this patient's workup, namely: (a) previous MI with RV involvement; (b) cardiac CT with pericardial thickening and anomal-
pulmonary venous return also contributing to RV volume overload and restrictive physiology; (c) CMR revealing diffuse pericardial LGE and a Qp/Qs of 1.4. However, no imaging modality could unequivocally identify constrictive physiology, possibly given the low fluid status caused by previous intravenous diuretic use. Moreover, while several RHC signs suggestive of constrictive pericarditis were present in our case, we were not able to assess the systolic area index, an additional sensitive marker for ventricular interdependence in constrictive pericarditis [1].

Diagnosing constrictive pericarditis is, therefore, challenging, particularly in those with other possible causes for RHF. Besides diagnosis, management could also be controversial. While findings of pericardial LGE at CMR could have led us to consider a trial of anti-
flammatory therapy before surgery [2], concomitant hemodynamically relevant left-to-right shunt possibly contributing to RV overload justified Heart-Teams’ decision to offer simultaneous pericardiectomy and shunt correction, also allowing us to reach the final histological diagnosis. Dense lymphoplasmacytic infiltrates and focal fibrosis with storiform pattern are key histopathological features of IgG4-related disease [3]. When present, IgG4 immunostaining is recommended to establish a definitive diagnosis.

IgG4-related disease is an autoimmune condition characterized by tissue IgG4 positive plasma cell infiltration [4–6]. Although cardiovascular involvement has been reported, constrictive pericarditis remains an unusual manifestation and often underrecognized presentation [6–8]. No guidelines exist regarding optimal treatment. Management usually relies on pericardiectomy and immunosuppressive therapies [7, 8].

Constrictive pericarditis remains a rare form of IgG4-related disease. A high-suspicition index, multimodality assessment and thorough investigation are critical in establishing the correct diagnosis.
Fig. 1 (See legend on previous page.)
Abbreviations
CMR: Cardiac magnetic resonance; CT: Computed tomography; IgG4: Immunoglobulin-G4; LGE: Late-gadolinium enhancement; MI: Myocardial infarction; RHC: Right heart catheterization; RHF: Right heart failure; RV: Right ventricle.

Supplementary Information
The online version contains supplementary material available at https://doi.org/10.1186/s12872-022-02468-1.

Additional file 1. Cardiac magnetic resonance (short axis, free-breathing cine) showing ventricular septum "d-shape".

Acknowledgements
None to report.

Authors’ contributions
SM wrote the first draft of this article; MC, PF, CA, CF, DM and BS incorporated feedback in subsequent drafts and revisions; MC, PF, CA, CF, DM, BS, MM and JPN contributed to revisions and reviewed the final draft; SM composed the figure and submitted the final version of this article, on behalf of all the authors. All authors read and approved the final manuscript.

Funding
None to report.

Availability of data and materials
Not applicable.

Declarations

Ethics approval and consent to participate
The patient has verbally consented to the publication of his medical case in a Medical Journal. The corresponding author (SM) also obtained written consent from a family member. This investigation was conducted in accordance with the World Medical Association Declaration of Helsinki (seventh revision, Fortaleza, 2013) and the Declaration of Istanbul (2008). Furthermore, the authors declare that the figures within this article do not allow the identification of the patient. Dates were not specified to comply with patient confidentiality. This case report was exempt from ethics’ board approval.

Consent for publication
The patient has verbally and written consented to the publication of his medical case in a Medical Journal. The corresponding author (SM) obtained the informed consent.

Competing interests
The authors declare that they have no competing interests.

Fig. 2  A (Hematoxiline-eosin, 100X): Pericardium with diffuse fibrous thickening, with focal areas of intersecting fascicles (storiform pattern—asterisk), and fibrinous areas at the surface (arrow). B (Hematoxiline-eosin, 400X): Patchy lymphoplasmacytic infiltration, and areas with Russell bodies (arrow)—atypical plasma cells with eosinophilic, homogeneous immunoglobulin-containing inclusions in cytoplasm. C (CD138 immunostaining, 400X): CD138 membranous immunostaining demonstrating plasma cells forming clusters. D (IgG4 immunostaining, 400X): cytoplasmatic immunostaining demonstrating an increased absolute count of IgG4-positive cells (> 20 per high-power field) with a marked storiform fibrosis pattern (arrow).
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Received: 4 September 2021   Accepted: 21 January 2022
Published online: 04 February 2022

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