



Concurrent acute coronary and Takotsubo syndrome: two in one

M. Wagener^{1,2} · R. Twerenbold^{1,3} · Michael J. Zellweger¹ · Philip Haaf¹

Received: 1 February 2023 / Accepted: 6 February 2023 / Published online: 13 March 2023
© The Author(s) 2023

A 69-year-old woman with metabolic syndrome was referred for emergency coronary angiography with chest pain, signs of left ventricular decompensation, elevated high-sensitivity cardiac troponin T (3686 ng/l, norm < 14 ng/l) and inferolateral ST-segment elevations. Apart from the acute chest pain itself no further emotional or physical stress factors could be identified.

Coronary angiography revealed a 1-vessel coronary artery disease with a distally occluded right posterolateral branch (Panel 1a, 1b, Supplement Video S1) and elevated left ventricular end-diastolic pressure of 34 mmHg. Considering the tortuosity and small vessel diameter, the patient was treated conservatively with dual antiplatelet therapy. Echocardiography showed severely depressed ejection fraction with akinesia of the apical two thirds of the myocardium with basal hyperkinesia. Cardiovascular magnetic resonance (CMR) imaging showed evidence of a focal acute transmural inferolateral midventricular myocardial infarction (Panel 2a, 2b) as well as a myocardial oedema of the akinetic apical two thirds of the myocardium (Panel 3a, 3b and 4a-d, Supplement Video S2). The findings of the patient were interpreted as concurrent Takotsubo syndrome (TTS) and acute myocardial infarction.

Two months later, left-ventricular global systolic function completely normalized, myocardial oedema as well as

apical ballooning resolved (Supplement Panel S1a-b, S2a-d, Supplement Video S3). Furthermore, wall thinning with dyskinesia developed of the transmurally infarcted inferolateral midventricular segment (Supplement Panel S3a-b) (Fig. 1).

Discussion

Acute coronary syndrome (ACS) is still widely regarded as an exclusion criterion of TTS and discrimination between them can sometimes be challenging. Although, concomitant coronary artery disease in TTS is known to be present in around 10–29% of cases, patients with concurrent TTS and acute obstructive CAD are often misdiagnosed as classical ACS [1] since presence of ACS is still widely regarded as an exclusion criterion for TTS.

A multimodality imaging approach including CMR should be considered when coronary angiography findings do not match with ventriculography or echocardiography. The presence of myocardial infarction should not exclude TTS per se but ACS may occasionally trigger - rather than exclude TTS, as recently outlined in the latest consensus paper on TTS [2].

✉ Philip Haaf
philip.haaf@usb.ch

¹ Department of Cardiology and Cardiovascular Research
Institute Basel (CRIB), University Hospital Basel, University
of Basel, Basel, Switzerland

² Galway University Hospitals, Galway, Ireland

³ University Center of Cardiovascular Science, University
Heart and Vascular Center Hamburg, Clinic for Cardiology,
University Medical Center Hamburg-Eppendorf, Hamburg,
Germany

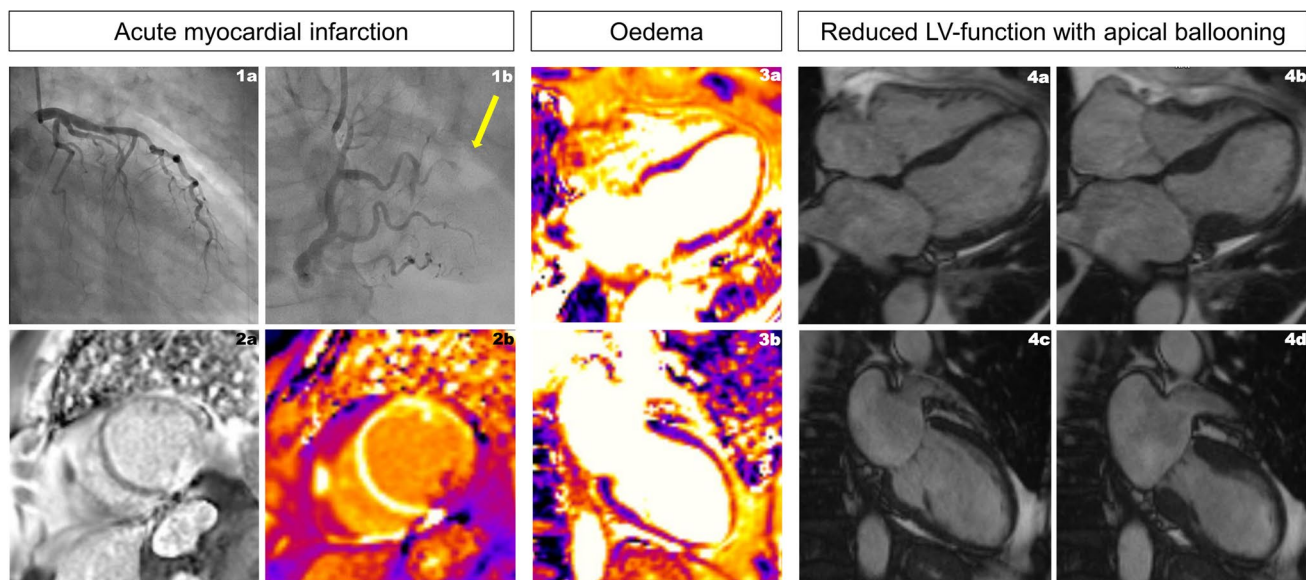


Fig. 1 1a – RAO caudal view of the left anterior descending and circumflex artery; 1b – RAO cranial view of the right coronary artery with occluded distal right posterolateral branch (yellow arrow); 2a/b – Cardiovascular Magnetic Resonance (CMR) imaging with evidence of a focal acute transmural inferolateral midventricular myocardial infarction (2a: Late Gadolinium Enhancement; 2b: T1 post-contrast

Map); 3a and 3b – CMR T2 map showing myocardial oedema of the apical two thirds of the myocardium; 4a–d –four chamber view cine images in diastole (4a) and apical ballooning in systole (4b); two chamber view cine images in diastole (4c) with apical ballooning in systole (4d)

Supplementary Information The online version contains supplementary material available at <https://doi.org/10.1007/s10554-023-02813-1>.

Author contributions Drafting of the manuscript: MW, PH. Critical revision of the manuscript for important intellectual content: MW, RT, MJZ, PH. Final approval of the Manuscript: MW, RT, MJZ, PH.

Funding Open access funding provided by University of Basel. The authors declare that no funds, grants, or other support were received during the preparation of this manuscript.

Declarations

Competing interests The authors declare no competing interests.

Consent for publication The authors confirm that written consent for submission and publication of this case report including images and associated text has been obtained from the patient in line with COPE guidance.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes

were made. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit <http://creativecommons.org/licenses/by/4.0/>.

References

1. Templin C et al (2015) Clinical features and outcomes of takotsubo (stress) cardiomyopathy. *N. Engl. J. Med* 373(10):929–938. <https://doi.org/10.1056/NEJMoa1406761>
2. Ghadri J-R et al (2018) International Expert Consensus Document on Takotsubo Syndrome (Part I): clinical characteristics, Diagnostic Criteria, and pathophysiology. *Eur Heart J* 39:2032–2046. <https://doi.org/10.1093/eurheartj/ehy076>

Publisher's Note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.