

Myeloproliferative invasion of arterial walls: Premortem diagnosis by temporal artery biopsy

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A 72-year old woman with a previous history of polycythemia vera presented with acute chest pain. The full blood count showed leukocytosis (WBC $43 \times 10^9/L$) with 90.1% neutrophils, microcytic anemia (Hb 103 g/L), thrombocytosis ($528 \times 10^9/L$) and no blasts. CT scan revealed a spontaneous intramural hematoma of the descending aorta (Figure 1) and an intra-abdominal hematoma. Active bleeding of the right inter-

nal iliac artery caused the latter and was treated with coil embolization. Bone marrow biopsy showed evolution to a myeloproliferative-dysplastic syndrome with excessive neutrophil counts, not further classifiable. Because of the hypothesis of invasion of the vascular walls by



FIGURE 1 CT scan showing an intramural hematoma of the descending aorta

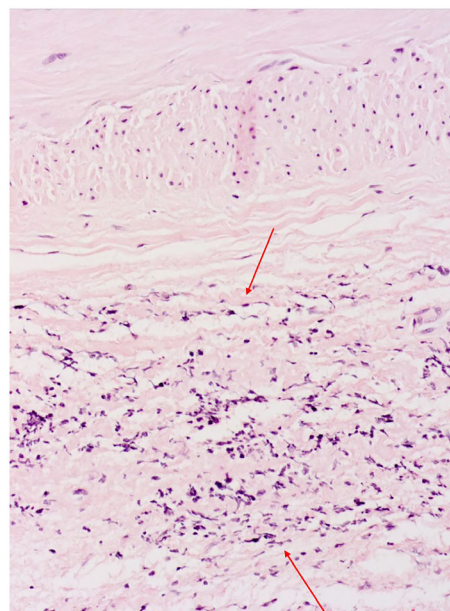


FIGURE 2 A section through a temporal artery segment (Hematoxylin eosin stain, magnification $\times 200$) shows an increased cellularity in the tunica adventitia. The tunica intima is slightly hyperplastic, but free of inflammation. There are no signs of giant cell arteritis, as no inflammation is present in the tunica media, and no giant cells or fragmentation of the elastica interna or externa are seen

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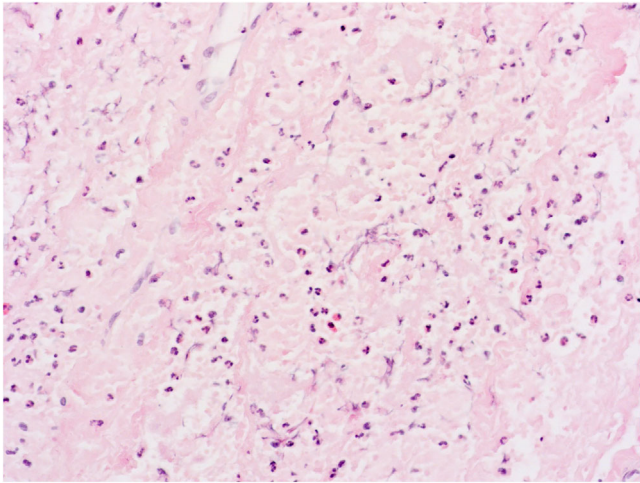


FIGURE 3 A section through a temporal artery segment (Hematoxylin eosin stain, magnification x 400). Detail of the tunica adventitia shows increased neutrophil infiltration

neutrophils, we performed a biopsy of the temporal artery. Although the latter was clinically normal, the biopsy showed a dense neutrophilic infiltrate in the adventitia without inflammation or giant cells in the tunica media, not diagnostic of giant cell arteritis (Figure 2 and 3). These findings support the hypothesis of arterial wall damage and intramural hemorrhage due to infiltration of malignant neutrophils, as has been reported in some post mortem studies [1,2]. Our case demon-

strates that temporal artery biopsy can be used to safely diagnose this complication of myeloproliferative disease.

CONFLICT OF INTEREST

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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