Vascular Ehlers-Danlos syndrome with distinct histopathologic features

Hee Sang Hwang¹, Jin Woo Song², Se Jin Jang³

Departments of ¹Pathology and ²Internal Medicine, Asan Medical Center, University of Ulsan College of Medicine, Seoul, Korea

Ehlers-Danlos syndrome is a connective tissue disorder that presents with heterogeneous manifestations depending on the involved genes. Vascular Ehlers-Danlos syndrome (vEDS), also known as Ehlers-Danlos syndrome type IV, is caused by a heterozygous mutation in the COL3A1 gene that encodes the pro-α1 chains of type III collagen, which attenuates the structural integrity of type III collagen-enriched tissues, such as skin, lung, liver, intestine, and blood vessel. vEDS can result in fatal complications such as arterial or intestinal rupture, but its clinical diagnosis is often challenging. Here, we present a case of vEDS that was diagnosed based on histopathologic and confirmatory genetic examinations.

CASE REPORT

A 19-year-old male was admitted to our hospital for recurrent episodes of spontaneous pneumothorax (6 times), which started about 8 months ago. His father had a history of recurrent pneumothorax and also suffered from spontaneous intra-abdominal bleeding due to aneurysmal rupture of the common hepatic artery 7 years ago. The patient had undergone wedge resection of the lung at another hospital, after which he experienced a secondary pneumothorax, but no specific pathologic diagnosis was made. Thoracic computed tomography at our hospital revealed intra-alveolar accumulation of red blood cells and hemosiderin-laden macrophages (Fig. 1B, C), indicating recurrent episodes of intra-alveolar hemorrhage. The most diagnostic histologic finding was the presence of several round fibro-osseous nodules scattered in a hemorrhagic background of lung parenchyma (Fig. 1D, E). Disruption of the vascular wall was not identified, even in the examination under van Gieson elastic staining. Due to suspicion of vEDS based on the patient’s familial history and the histopathologic findings, COL3A1 gene sequencing was performed. The sequencing test revealed a heterozygous germline mutation in the spliceosome acceptor site in intron 23 of the COL3A1 gene (COL3A1 c. 1662 + 1G > A [p. Gly537_Pro554del]) (Fig. 1F), a known pathogenic mutation in the ClinVar database (https://www.ncbi.nlm.nih.gov/clinvar/variation/101269/). Hence, the patient was diagnosed with vEDS.

DISCUSSION

vEDS comprises about 4%–5% of Ehlers-Danlos syndrome cases and has a high rate of fatal complications related to cardiovascular and gastrointestinal systems, such as cerebrovascular hemorrhage or intestinal rupture. In addition, pregnancy can be fatal in patients with vEDS, who might develop ruptures in the uterus or uterine vessels [1]. Therefore, prompt diagnosis is the mainstay of medical management of vEDS to lower the rate of such fatal complications [2]. However, the rarity of vEDS makes its diagnosis challenging, frequently leading to delayed diagnosis. The pathologic diagnosis of vEDS is challenging because the characteristic histopathological findings of vEDS are not widely

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recognized. Abrupt caliber changes in the muscularis propria with diverticular formation have been identified in colon resection specimens of vEDS patients [3]. Kawabata et al. [4] reported that fragile lung parenchyma with spontaneous laceration, acute hemorrhage and/or hematoma, and fibro-osseous nodules, which are hypothesized to be a consequence of the organization of hematomas, are frequently identified in surgical lung biopsy specimens of vEDS patients, findings which serve as clues for pathologic diagnosis. In line with these findings, a previous report described a case of vEDS that was diagnosed based on identification of fibro-osseous nodules and vascular disruption in the lung parenchyma, suggesting the potential of histopathologic features in clinical diagnosis of vEDS [5]. Therefore, the distinct histopathologic features of vEDS should be considered when assessing surgical lung biopsy specimens from patients with recurrent spontaneous pneumothorax and/or hemorrhagic lung parenchymal cysts.

In conclusion, identifying characteristic histopathologic features in surgical lung biopsy samples, such as parenchymal fragility, spontaneous laceration, parenchymal hemorrhage, and fibro-osseous nodules, can contribute to early clinical diagnosis and proper management of vEDS.

**Ethics Statement**
This case study was approved by the Institutional Review Board of Asan Medical Center (2020-1889), which waived the requirement for informed consent.

**Availability of Data and Material**
Data sharing not applicable to this article as no datasets were generated or analysed during the study.

**Code Availability**
Not applicable.

**ORCID**
Hee Sang Hwang https://orcid.org/0000-0001-9486-9214
Jin Woo Song https://orcid.org/0000-0001-5121-3522
Se Jin Jang https://orcid.org/0000-0001-8239-4362

**Author Contributions**
Conceptualization: SJJ, JWS, HSH. Data curation: HSH. Formal analysis: HSH, SJJ. Investigation: HSH. Methodology: HSH, JWS. Resources: SJJ, JWS. Supervision: SJJ. Writing—original draft: HSH. Writing—review & editing: HSH, SJJ. Approval of final manuscript: all authors.
Conflicts of Interest
The authors declare that they have no potential conflicts of interest.

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