

Pneumorrhachis: an under-recognized entity correlates with severity of spontaneous pneumomediastinum

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Spontaneous pneumomediastinum (SPM) is an uncommon condition that predominantly affects young males. Due to its rarity, the optimal management still remains uncertain without consensus.

We read with great interest the article titled “*Clinical Experience of Spontaneous Pneumomediastinum: Diagnosis and Treatment*” by Kim *et al.* (1), who reported one of the largest series of SPM in literature in the last decade (2-6). The authors demonstrate the result of no difference in symptoms, clinical course and SPM, which was conservatively managed regardless of the degree of severity via their proposed radiologic classification.

In the recent 2 years at our institution, a preliminary investigation for SPM was initiated retrospectively based on the authors’ classification. Surprisingly, we found that several cases of severe type of SPM exhibit additional epidural air, which is also known as pneumorrhachis (Figure 1A,B). Spontaneous pneumorrhachis is a rare condition and occasionally correlates with SPM. It is hypothesized to originate from air leak spreading through the posterior mediastinum into the epidural space via the

cervical fascia planes or neural foramen (7). Owing to its scarcity in nature, there is no established incidence of pneumorrhachis associated with SPM except for only one published study revealing 4 out of 42 cases (9.5%) of SPM as well as extensive subcutaneous emphysema, suggesting that in SPM, pneumorrhachis is uncommon but not extraordinary (8).

With the continuing advancement in imaging studies and frequent adoption of chest computed tomography in cases of SPM, pneumorrhachis may be increasingly detected. In spite of its benign and self-limiting entity, we would like to reclassify the degree of severity in SPM, encompassing the most severe type with concomitant pneumorrhachis and/or other extrathoracic air leak such as pneumoperitoneum and pneumoretroperitoneum (Figure 1C,D).

We hope this modified classification will be beneficial for future studies in a large scale or with prospective intent, enabling a validated outcome analysis based on degree of severity and drawing a more reliable and convincing conclusion in terms of diagnostic and therapeutic purposes for SPM.

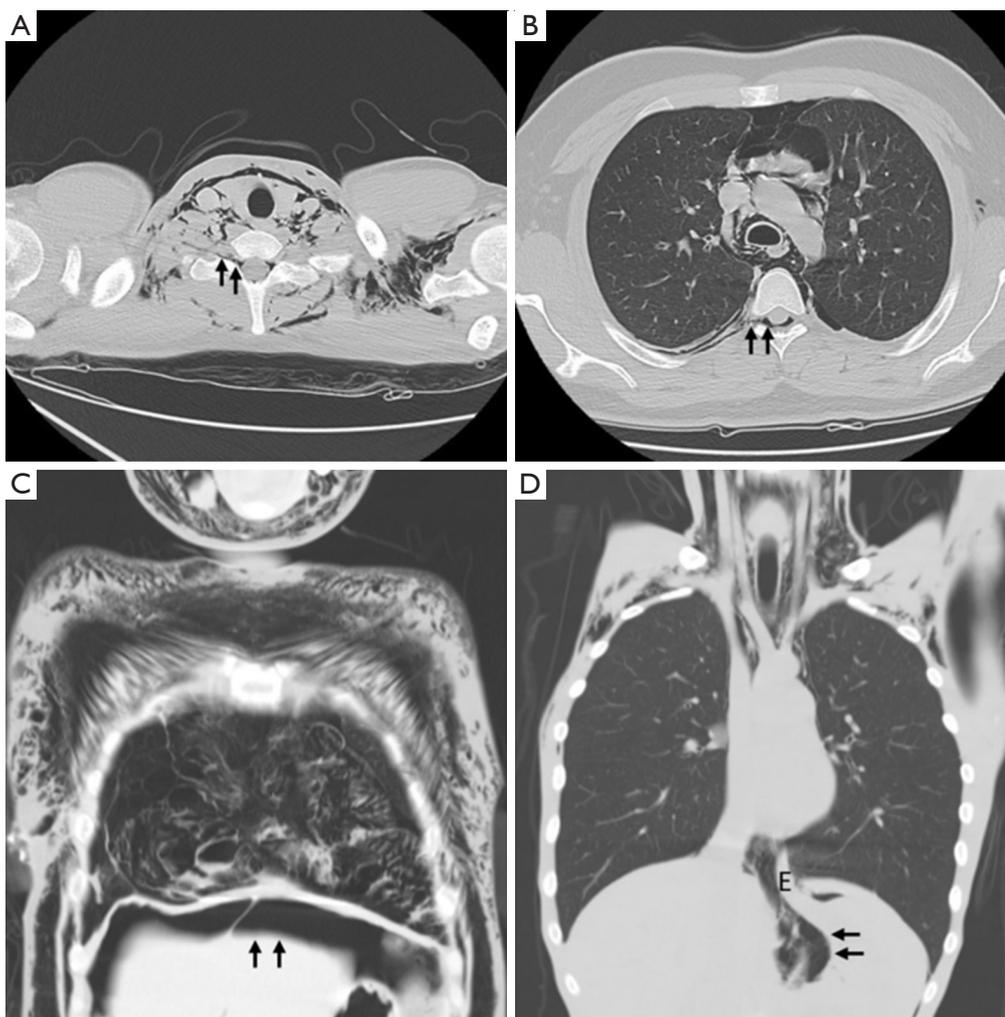


Figure 1 Spontaneous pneumomediastinum with concomitant extrathoracic air leaks. (A) Patient 1 disclosed neck and chest wall subcutaneous emphysema, pneumomediastinum, and pneumorrhachis originated from deep cervical fascial planes (arrows); (B) patient 2 revealed pneumomediastinum with pneumorrhachis, which resulted from adjacent neural foramen (arrows); (C) patient 3 showed massive subcutaneous and mediastinal emphysema with concomitant pneumoperitoneum (arrows); (D) patient 4 demonstrated pneumomediastinum and air dissection along periesophageal fascial planes to cause pneumoretroperitoneum (arrows). E, esophagus.

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Footnote

Conflicts of Interest: The authors have no conflicts of interest to declare.

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