Klippel-Feil Syndrome Accompanied by Pulmonary Artery Sling

Yoshiyuki Kawano, Akira Tamura, Yusei Abe and Junichi Kadota

Key words: Klippel-Feil syndrome, pulmonary artery sling

(DOI: 10.2169/internalmedicine.47.0727)

A 27-year-old man was diagnosed as Klippel-Feil syndrome accompanied by a ruptured aneurysm of the non-coronary sinus of Valsalva and underwent a patch closure for the ruptured aneurysm (1). Multislice computed tomography performed before the operation showed that the left pulmonary artery originated from the posterior aspect of the right pulmonary artery and passed rightward around the trachea and between the lower portion of the trachea and esophagus (Picture 1). Since he did not have any symptoms caused by compression of the trachea and esophagus by the left pulmonary artery, an operation for this congenital anomaly was not performed. To the best of our knowledge, this is the first case report describing a patient with Klippel-Feil syndrome accompanied by pulmonary artery sling.

References