Abstract:

Pheochromocytoma is a type of neuroendocrine tumors with catecholamine secretion. They have adrenal medullary origin. The classic symptoms of tumor are headache, palpitation and diaphoresis. Surgical resection of tumor renders symptoms relief. Occasionally some uncommon presentations of pheochromocytoma have been found in the literature. Here, we presented a case of a 42year old female who suddenly collapsed mimicking acute coronary syndrome leading to the diagnosis of a pheochromocytoma.