Meigs syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Meigs syndrome. ORPHA:314451

Meigs syndrome is a rare neoplastic disease characterized by the clinical triad of benign ovarian tumor (typically, ovarian fibroma or fibroma-like tumor), hydrothorax and ascites, which resolve after tumor resection. Patients usually present with dyspnea, pelvic mass with or without a tender, distended abdomen and/or weight loss.