Pulmonary arterial hypertension associated with portal hypertension

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Pulmonary arterial hypertension associated with portal hypertension. ORPHA:275813

Pulmonary arterial hypertension associated with portal hypertension (PAH-PH) is a form of pulmonary arterial hypertension (PAH), characterized by an elevated pulmonary arterial resistance leading to right heart failure observed as a complication of portal hypertension.