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# Eisenmenger syndrome

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base.*

*Eisenmenger syndrome. ORPHA:97214*

Eisenmenger syndrome (ES) is a form of pulmonary arterial hypertension (PAH) associated with unoperated congenital heart disease and is characterized by congenital heart malformations with reversed or bi-directional shunting through an intra-cardiac or intervascular (usually aorto-pulmonary) communication with the development of PAH.