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# Neuhauser anomaly

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Neuhauser anomaly. ORPHA:99078*

Neuhauser anomaly is a rare cardiovascular morphological anomaly due to maldevelopment of embryonal aorta resulting in right aortic arch and left ligamentum arteriosum characterized by tracheoesophageal compression symptoms (stridor, dyspnea, dysphagia, apnoeic episodes, recurrent respiratory infections).