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# Laubry-Pezzi syndrome

INSERM

## Source

INSERM. (1999). *Orphanet: an online rare disease and orphan drug data base*. [Laubry-Pezzi syndrome](#). ORPHA:99094

Laubry-Pezzi syndrome is a rare, non-syndromic, congenital heart malformation characterized by the prolapse of an aortic valve cusp into a subjacent ventricular septal defect due to Venturi effect, resulting in aortic regurgitation. Patients typically present with symptoms of progressive aortic valve insufficiency, such as shortness of breath, heart palpitations, chest pain and exercise intolerance.