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Pulmonary Arteriovenous Malformation: Case ReportAndré Luiz Mascarenhas Silva1, Florentino Andrade De Melo Júnior1, Ana Paula Sampaio De Mattos1, Saulo Sacramento Meira1 1School Of Medicine, State University Of Southwest Bahia, Salvador-BA, Brazil. Abstract This Is A Clinical Case Study Involving The Respiratory System Of A Patient In The State Of Bahia. The Aim Of This Paper Is To Report The Jan 4th, 2024Macrocephaly- Capillary Malformation SyndromeMacrocephaly- Capillary Malformation Syndrome ... Neurological Complications (obstructive Hydrocephalus, Increased Intracranial Pressure, Cerebellar Tonsillar Ectopia Or Chiari Malformation; Epilepsy In People With HMEG) May Warrant Neurosurgical Intervention. Regular Monitoring Is Recommended (MRI Of The Brain In The First 8 Years Of Life, Renal Apr 1th, 2024Proctalgia Secondary To Rectal Arteriovenous Malformation ...Microcatheter (Medtronic, Dublin, Ireland) Superselec-tively Into A Superior Rectal Artery Branch. This Proced-ure Was Performed Under General Anaesthesia. Excellent Symptomatic Response Was Reported, With Improved Quality Of Life. Proctalgia Recurred Three Years Later, Req Apr 1th, 2024. A Rare Case Of Bilateral Jugular Venous MalformationSuperficial Veins Of Head And Neck. J Anat Soc India. 2005; 49:61-62. 5. Hollinshead WH. Anatomy For Surgeons, 3rd Ed, Harper And Row, Philadelphia. 1982. 6. Deslangiers B, Vaysse P, Combes JM. Contribution To The Study Of The Tributaries And The Termination Of The External Jugular Vei Feb 4th, 2024Type I Split Spinal Cord Malformation: Literature Review ...Of Bone In The Middle Of The Thecal Sac (Figure 2b). This Island Of Bone Met In The Midline And Went Intradurally, Splitting The Spinal Cord Into Two Halves. We Then Brought In The Microscope For Visualization, Illumination And Magnification. We Opened The Dura Feb 3th, 2024Hepatic Arteriovenous Malformation Of Cortical And Vascular Development In One ...Marcelo Valente, Kette D. Valente, Sofia S. M. Sugayama, Chong Ae Kim Summary: Vascular And Cortical Anomalies Have Been Found In A Family With Parietal Foramina Type 2 (PFM2), Which Is Determined By The ALX4 Gene. It Is Believed That ALX4 Has A Bone-restricted Expression. We Report A Case Of PFM With Age-related Size Variation In A 4-year-old ... Apr 3th, 2024

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