Anorectal Embryology and Anatomical Structure

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DESCRIPTION

The anorectal region is derived from the hindgut, which is the posterior part of the embryonic gut tube. The development of the anorectal region involves complex interactions between the endoderm, mesoderm, and ectoderm. During early embryonic development, the hindgut extends caudally and forms a cloacal membrane, which separates the hindgut from the surface ectoderm. The cloacal membrane consists of two parts: the urorectal septum and the anal membrane. The urorectal septum divides the cloaca into two parts: the urogenital sinus and the anorectal canal.

The anorectal canal initially communicates with the exterior through the anal membrane. The anal membrane is formed by the ectoderm and the underlying cloacal endoderm. At around the 8th week of gestation, apoptosis (programmed cell death) occurs in the center of the anal membrane, forming an opening called the anal pit. The anal pit then elongates to form the anal canal. The levator ani muscle, which is important for maintaining fecal continence, develops from the mesoderm in the pelvic region. The puborectalis muscle, a part of the levator ani, originates from the ventral mesoderm and attaches to the anorectal junction. The external anal sphincter muscle develops from the surface ectoderm. Malformations of the anorectal region are common and can range from mild to severe. These malformations can occur due to genetic mutations, environmental factors, or a combination of both. Some examples of anorectal malformations include anal atresia (absence of the anal opening), imperforate anus, and anorectal agenesis.

The anorectal region is a complex area that includes the anus and rectum, and is responsible for the elimination of solid waste from the body. An understanding of the embryology and anatomical structure of this region is important for diagnosing and treating various anorectal disorders. Embryologically, the anorectal region arises from the endoderm, ectoderm, and mesoderm germ layers. The rectum and upper anal canal arise from the endoderm, while the lower anal canal arises from the ectoderm. The rectum and upper anal canal arise from the body. An understanding of the embryology and anatomy of the anorectal region is essential for healthcare professionals who diagnose and treat patients with anorectal disorders.

The internal anal sphincter is a smooth muscle that is under involuntary control, while the external anal sphincter is a skeletal muscle that is under voluntary control. The blood supply to the anorectal region is provided by the inferior rectal artery, which is a branch of the internal pudendal artery, and the superior rectal vein, which drains into the inferior mesenteric vein. The venous drainage of the anorectal region is through the inferior rectal veins, which drain into the internal iliac vein, and the superior rectal vein, which drains into the inferior mesenteric vein. The anorectal region is a complex area with a diverse embryological and anatomical structure. Understanding the anatomy and physiology of this region is important for the diagnosis and treatment of various anorectal disorders.

The embryonic development of the anorectal region involves the fusion of the urorectal septum and the cloacal membrane, which forms the anal canal and separates it from the urogenital sinus. The anal canal is surrounded by internal and external anal sphincters, which control the flow of feces and prevent incontinence. The anatomy of the anorectal region is further complicated by the presence of the levator ani muscle, the pelvic floor, and the pudendal nerve. The anatomic variations in these structures can lead to various clinical conditions, including fecal incontinence, anal fissure, and haemorrhoids.

Overall, understanding the embryology and anatomy of the anorectal region is essential for healthcare professionals who diagnose and treat patients with anorectal disorders.