HYPOSPADIA AND ASSOCIATED CONGENITAL GENITAL DEFECTS IN A JAMNAPARI KID (CAPRA HIRCUS): A CASE STUDY

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Abstract: Hypospadias generally occur as a result of imperfect closure or fusion of the urethral grooves during phallus elongation. The purpose of this case study is to report rare congenital urinary tract anomaly, hypospadias and other associated congenital genital tract anomalies in a jamnpari goat kid.

Keywords: Hypospadias, urethral grooves, congenital, genital tract, anomalies.

Introduction

Congenital defects of the urinary tract are not common in farm animals (Dennis and Leipold, 1979). Hypospadias is a rare congenital malformation of the urethra in domestic mammals. Hypospadias generally occur as a result of imperfect closure or complete lack of fusion of the urethral grooves during phallus elongation (Kahn et al., 2005; Radostits et al., 2007). In a survey of the occurrence of congenital anomalies in goats, the occurrence of congenital hypospadias was 0.066% (Al-Ani et al. 1998). Hypospadias is accompanied by hypoplasia of the corpus cavernosum urethra, causing the urethra to open anywhere along its length at one or more locations (Alam et al., 2005). The hypospadias is thus classified on the basis of anatomic localization such as glandular, penile, scrotal, perineal, or anal (Ader and Hobson, 1978; Kahn et al., 2005). The purpose of this study is to report a rare congenital urinary tract anomaly, hypospadias and other associated congenital genital tract anomalies in a kid.

Case history and Observation

A five day old Jamnapari kid (Fig. No.1) was presented to TVCC with the complaints of mild depression, stranguria and dysuria through an opening at abnormal location. Clinical
examination revealed a slit-like opening of the urethra on the under-surface of the shaft of the penis just anterior to bifid scrotum each containing a normal sized testis (Fig. No.2). Associated genital defects included underdeveloped cord like penis and absence of prepuceal orifice. Anus was present and defecation was normal.

**Treatment and discussion**

An attempt was made to locate the urethral opening by means of an infant catheter but in vain. During the course of observation, kid has urinated in a projectile manner from an opening at the perineum. Hence, surgical correction was not attempted as the animal was urinating normally and also there was no discomfort to the kid. Owner was advised to bring the animal for follow-up or if at all any complication arises at later stage.

Formation of the external male genitalia is a complex developmental process involving genetic programming, cell differentiation, hormonal signaling, enzyme activity and tissue remodeling (Baskin, 2002). Disturbance in the process of development might lead to disruption of the fusion of the urogenital folds at different sites along the urogenital tract. Environmental toxicants and xenoestrogens acting during fetal life have been partly implicated in an increasing incidence of hypospadias, as well as other reproductive tract abnormalities (Baskin *et al.*, 2001). The aetiology of hypospadias is not well understood; it seems to be multifactorial and may be related to genetic, endocrinological, and environmental factors (Silver, 2000). One possible explanation for increase in the incidence of hypospadias may be environmental contaminations. Farm animals are constantly exposed to estrogenic compounds which are known for their ability to disrupt reproduction, the so called oestrogenic endocrine disruptors (Kim *et al.*, 2004). Surgical correction was not advised in the present case because it was associated with other congenital abnormalities like of underdeveloped penis and agenesis of prepuceal orifice.

Recently, association between environmental oestrogen-like compounds and hypospadias has been suggested (Sharpe and Skakkebaek, 1993). Karyotype and gonadal histological features can be used to characterize this disorder as an abnormality of sexual differentiation.

There is no conflict of interest that could be perceived as prejudicing the impartiality of the case reported.

**References**


Fig.1: Jamnapari kid with multiple congenital anomalies of the urogenital tract.

Fig.2: Jamnapari kid showing hypospadias, cleft scrotum and agenesis of prepucial orifice.