

CASE REPORT

A RARE CASE OF BIFID PENIS WITH IMPERFORATE ANUS

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ABSTRACT: BACKGROUND: Diphallus is rare anomaly & accompanying anomalies vary from bifid scrotum, bladder exstrophy, imperforate anus and colo-rectal anomaly such as duplication, and other associated anomalies. We report a case of 2 day old newborn who had bifid penis, bifid scrotum with imperforate anus.

KEYWORDS: Diphallus, Imperforate Anus.

INTRODUCTION: Bifid penis is a rare anomaly and accompanying anomalies vary from imperforate anus, colorectal anomaly, bladder extrophy and others. Incidence is 1 in every 5 million live births.

CASE PRESENTATION: A 2 day old male newborn was brought us because of distended abdomen, abnormal genitalia & imperforate anus. Thus antenatal history was uneventful, and investigations were negative for other associated anomalies postnatally.

The birth history was normal. It is a consanguineous marriage. Examination of genitalia revealed a well formed double penis, functioning urethra, bifid scrotum with each compartment containing a testicle. Perineal examination showed imperforate anus.

The values of blood analysis were within normal limits. Postoperatively abdominal ultrasonography showed bilateral normal kidneys. The day after he underwent colostomy, we performed an intravenous pyelography which showed normal kidneys and urethra. It was diagnosed as high lining recto urethral fistula.

DISCUSSION: Diphallus or duplication of the penis is rare anomaly that occurs 1 in every 5 million live births.¹ They may be associated with different anomalies such as anorectal malformation and lower genitourinary abnormalities.² Azmy has reported complete duplication of the hindgut which had two anal openings and a lower urinary tract with diphallus.³

The first case was reported in 16094. Schneider classified diphallus in three groups; diphallus of glans alone, bifid diphallus, and complete diphallus. Vilanova and Raventos have added a fourth category pseudodiphallus.^{1,4} The meatus may be normal, epispadic or hypospadic and the scrotum may be normal or bifid⁵ Treatment of diphallus usually includes excision of the duplicated penile structure and its urethra.^{1,6} Associated congenital anomalies are present in majority of the cases.^{1,3,7}

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Fig: Bifid Penis and Bifid Scrotum

CONCLUSION: The babies with diphallus though have a rare incidence have to be examined carefully because of the high incidence of other systemic anomalies. Treatment includes excision and repair of associated anomalies.

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