

## Hypospadias - Paediatric Full Clinical Guideline - Burton Only

Reference no.: WC/NP/44N/Aug 20/v006

### 1. Introduction

Hypospadias is a congenital malformation of the male genitalia characterised by incomplete development of the urethra in which the opening is located on the ventrum rather than at the tip of the penis. Severity depends on the location of the urethral opening. In most cases, the opening is located on the glans. Moderate or severe forms of hypospadias occur when the opening is situated more proximally on the penile shaft, at the base of the penis or, in rare cases, in the scrotum or perineum.

### 2. Aim and Purpose

To inform staff, within all areas, in the management of neonates and infants with hypospadias.

### 3. Classification

Hypospadias can be classified according to location (glandular, coronal, subcoronal, distal penile, mid shaft, proximal penile, peno-scrotal, scrotal or perineal) or according to severity i.e. mild (glandular or coronal), moderate (sub-coronal or penile) and severe (peno-scrotal, scrotal or perineal).

### 4. Management

The aetiology of hypospadias remains unknown with environmental exposure in the form of endocrine disruptors being the most likely explanation. Appendix 1 is a general guideline on the management of newborns noticed to have hypospadias on the Postnatal Wards.

In cases of undescended testes together with hypospadias (proximal defect), chromosome analysis should be requested and on call consultant to be informed. An urgent referral to Paediatric Urologist may be required.

Distal defects (e.g. hooded prepuce) can be referred to Urologist after reassessment in the clinic at 2-3 months of age.

Advise parents to avoid circumcision on religious grounds until reviewed by Urologist.

5. **Suggested key review criteria for monitoring and/or audit**

Appropriateness of management/follow up according to the severity of hypospadias

Associated congenital/chromosomal anomalies

Timing of referral to Paediatric Urologist

Delayed diagnosis (beyond neonatal age group)

6. **References**

Abdullah NA, Pearce MS, Parker L, Wilkinson JR, Jaffray B and McNally RJQ. Birth prevalence of cryptorchidism and hypospadias in northern England, 1993–2000 Arch Dis Child. 2007;92:576-579

Ahmed SF, Dobbie R, Finlayson AR, Gilbert J, Youngson G, Chalmers J & Stone D. Prevalence of hypospadias and other genital anomalies among singleton births, 1988–1997, in Scotland Arch Dis Child Fetal Neonatal Ed. 2004;89:149-151

Leonard J. Paulozzi, J. David Erickson, and Richard J. Jackson  
Hypospadias Trends in Two US Surveillance Systems Pediatrics.  
1997;100:831-834

Baskin LS, Ebberts MB. Hypospadias: anatomy, etiology, and technique. *J Pediatr Surg* 2006;41 (3) :463–72

<b>Title:</b> Management of Hypospadias			<b>Policy No:</b> WC/NP/44N <b>Version No:</b> 6		
<b>Document Type:</b>		<b>Clinical / Non Clinical:</b>		<b>Effective from:</b>	
Divisional Guideline		Clinical		Aug 2020	
<b>Responsibility:</b>		<b>Essential Reading for:</b>		<b>Information for:</b>	
Dr M Ahmed <b>Consultant Paediatrician</b>		All Paediatric Medical Staff		All Paediatric Nursing Staff All Midwives	
<b>Original Issue Date:</b>		<b>Date of Last Review:</b>		<b>Next Review Date:</b>	
January 2005		Aug 2020		Aug 2023 Extended March 2024	
<b>Reason for amendment:</b>					
Routine review and update					
<b>Linked Trust Policies:</b>		<b>Consulted:</b>		<b>Stored:</b>	
58: Policy Framework		All Paediatric Medical Staff All Paediatric Nursing Staff All Senior Nurse Managers All Midwives		Division of Women & Children's Guideline Intranet Server	
<b>Approved by:</b>		<b>Approved by Women's and Children's Division 27<sup>th</sup> August 2020</b>			

## Appendix 1: Management of Hypospadias

