P3 Hydronephrosis

National Cancer Institute

Source

National Cancer Institute. P3 Hydronephrosis. NCI Thesaurus. Code C123185.

Postnatal Hydronephrosis with the following clinical findings: 1) calyceal dilation and the ureter are the same as those in UTD P2, 2) the renal parenchymal is thinned, has increased echogenicity and/or has decreased corticomedullary differentiation, or 3) the bladder is abnormal (wall thickening, ureterocele, posterior urethral dilation). Cases in which there are parenchymal abnormalities but the APRPD is less than 15 mm, are classified as UTD P3. (Adapted from: Hiep T. Nguyen, Carol B. Benson, Bryann Bromley, Jeffrey B. Campbell, Jeanne Chow, Beverly Coleman, Christopher Cooper, Jude Crino, Kassa Darge, C.D. Anthony Herndon, Anthony O. Odibo, Michael J.G. Somers, Deborah R. Stein; Multidisciplinary consensus on the classification of prenatal and postnatal urinary tract dilation (UTD classification system); Pediatric Urology; December 2014 Volume 10, Issue 6, Pages 982-998)