ARPKD is a rare, monogenic ciliopathy characterized by the formation of bilateral renal cysts and congenital hepatic fibrosis.–The urinary bladder was empty with Foleys catheter in situ, no ascites was seen.last BG: pH: 7.45 PC02: 31.1 HCO3: 22 BE: –11 * Serum Lytes: NA: 133 K: 3.9 Cl: 109 * UA: Leukocytes: 20–23, gravity: 1.015 * BUN: 6 Cr: 0.18 * AST: 96 ALT: 34 – She also had abdominal distention, abdominal U/S was done and showed bilateral medullary nephrocalcinosis and no hepatic hemangiomas.Prenatally, oligohydramnios without any enlargement and/or increased echogenicity of .kidneys was observed