Hirschsprung disease

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hirschsprung disease. ORPHA:388

Hirschsprung disease (HSCR) is a congenital intestinal motility disorder that is characterized by signs of intestinal obstruction due to the presence of an aganglionic segment of variable extent in the terminal part of the colon.