"WAGR syndrome/11p deletion syndrome is a rare genetic syndrome in which there is a predisposition to several conditions, including certain malignancies, distinctive eye abnormalities, and/or intellectual disability. ""WAGR"" is an acronym for the characteristic abnormalities associated with the syndrome. The acronym stands for (W)ilms' Tumor, the most common form of kidney cancer in children; (A)niridia, partial or complete absence of the colored region of the eye(s) (iris or irides); (G) Genitourinary abnormalities, such as undescended testicles or hypospadias in males, or internal genital or urinary anomalies in females; and Mental (R)etardation (intellectual disability). A combination of two or more of these conditions is usually present in most individuals with WAGR syndrome/11p deletion syndrome. The clinical picture varies, depending upon the combination of associated abnormalities. WAGR syndrome/11p deletion syndrome is a rare genetic disorder that is thought to affect males more frequently than females. Because some affected individuals have external genitalia that strongly resembles that of the opposite sex, incorrect sex identification may occur initially"