GENE	PHENOTYPE	PHENOTYPE MIM NUMBER	INHERITANCE	CPT CODE
ACTL6A	BAF-related disorder	N/A	de novo	81479
ARID1A	Coffin-Siris Syndrome 2	614607	AD	81479
ARID1B	Coffin-Siris Syndrome 1	135900	AD	81479
ARID2	Coffin-Siris Syndrome 6	617808	AD	81479
DPF2	Coffin-Siris Syndrome 7	618027	AD	81479
PHF6	Borjeson-Forssman-Lehmann syndrome (BFLS)	301900	XLR	81479
SMARCA2	Nicolaides-Baraitser syndrome	601358	AD	81479
SMARCA4	Coffin-Siris Syndrome 4	614609	AD	81479
SMARCB1	Coffin-Siris Syndrome 3	614608	AD	81479
SMARCE1	Coffin-Siris Syndrome 5	616938	AD	81479
SOX11	Mental retardation, autosomal dominant 27	615866	AD	81479