DISEASE:
KBG syndrome

NAME:	KBG syndrome
DESCRIPTION:	A rare congenital malformation syndrome characterized by a typical facial dysmorphism, macrodontia of the permanent upper central incisors, short stature, skeletal anomalies, developmental delay and behavioral abnormalities.
ORPHACODE:	2332
SYNONYMS:	Short stature-facial and skeletal anomalies-intellectual disability-macrodontia syndrome
XREF(S):	Orphanet MeSH ICD-10 OMIM
ANALYTE(S):	ANKRD11
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RELATED CONTENT

Related Genetic Tests

- Short Stature (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Laboratories

- Centre de Génétique Médicale UCL
- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)

Related Analytes

- ankyrin repeat domain containing 11

Related Gene Panels

- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes)) - UCL
- Short Stature (46 genes) - IPG