CGT Basic v3.1.1

Gene	Disease
CFTR	Cystic fibrosis
FMR1	Fragile X syndrome
G6PD	Hemolytic anemia, G6PD deficient (favism)
GJB2	Deafness, autosomal recessive, type 1A
HBA1	Thalassemia, alpha-
HBA2	Thalassemia, alpha-
HBB	HBB-related hemoglobinopathy
SMN1	Spinal muscular atrophy