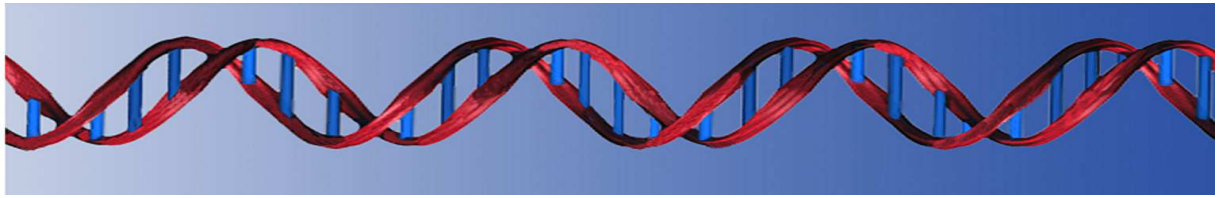

Molekulargenetik

Erbkrankheit, Genotyp-Variante	Gen, Genort, Chromosom
Retinitis Pigmentosa (autosomal dominant)	RHO, RP11, RP1, RDS
Retinitis Pigmentosa (X-Chromosomal)	RP3
Rett-like-Syndrom	CDKL5
Rett-Syndrom	MECP2
Rhesusinkompatibilität	RhCE, RhD
Romano- Ward Syndrom	KCNQ1, KCNH2, KCNE1, KCNE2, SCN5A, ANK2, KCNJ2, CACNA1C