meta_1461
meta_1587
meta_901
BNP,NP-proBNP,VEGF,ADMA,
TBX1 (22q11 deletion syndrome)
ELN (Williams-Beuren syndrome)
EVC2 and EVC (Wolf-Hirschhorn syndrome)
STRA6 (15q24 recurrent microdeletion syndrome)
FBN1 (Marfan syndrome)
MID1 (Opitz G/BBB syndrome)
RAD21 (Cornelia de Lange syndrome 4)
SETBP1 (Schinzel-Giedion midface retraction).
In addition, the genes contributing to non-syndromic CHD included
CRELD1
RAF1
CITED2
DVL1
SKI
ANP
BNP
TNP