Supplementary Table 3. Genetic disorders associated with ASD

22q11.2 deletions
Angelman syndrome
CHARGE syndrome
de Lange syndrome
Fragile X syndrome
MED 12 disorders (including Lujan-Fryns syndrome)
Prader-Willi syndrome
PTEN-associated disorders (Cowden syndrome, Bannayan-Riley-Ruvalcaba syndrome)
Rett syndrome
Smith-Lemli-Opitz syndrome
Smith-Magenis syndrome
Sotos syndrome
Tuberous sclerosis

ASD, autism spectrum disorder; CHARGE, coloboma, heart defect, atresia choanae, retarded growth and development, genital hypoplasia, ear anomalies/deafness; MED 12, mediator of RNA polymerase II transcription, subunit 12 homolog; PTEN, phosphatase and tensin homolog