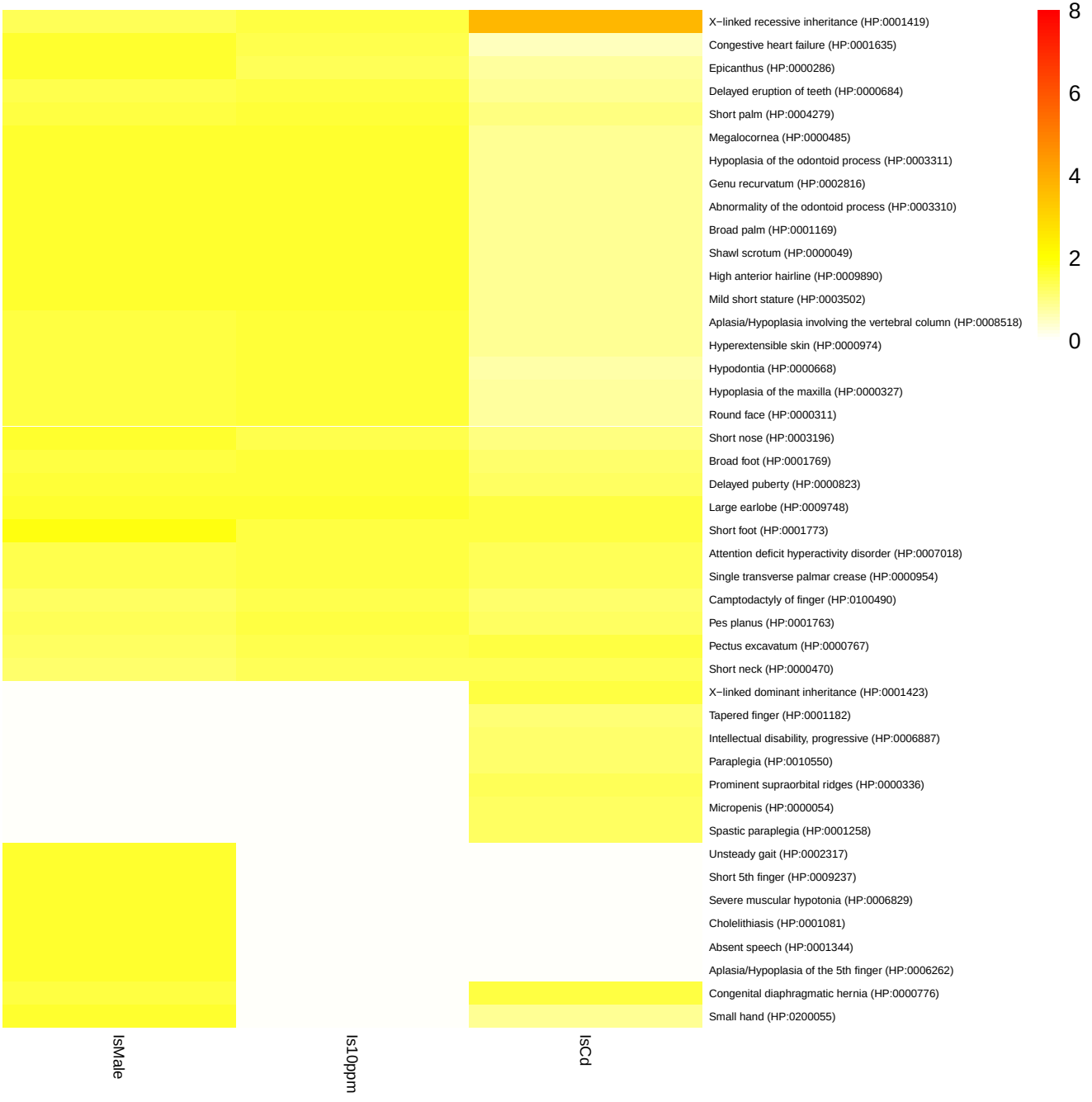
X−linked recessive inheritance (HP:0001419)
Congestive heart failure (HP:0001635)
Epicanthus (HP:0000286)
Delayed eruption of teeth (HP:0000684)
Short palm (HP:0004279)
Megalocornea (HP:0000485)
Hypoplasia of the odontoid process (HP:0003311)
Genu recurvatum (HP:0002816)
Abnormality of the odontoid process (HP:0003310)
Broad palm (HP:0001169)
Shawl scrotum (HP:0000049)
High anterior hairline (HP:0009890)
Mild short stature (HP:0003502)
Aplasia/Hypoplasia involving the vertebral column (HP:0008518)
Hyperextensible skin (HP:0000974)
Hypodontia (HP:0000668)
Hypoplasia of the maxilla (HP:0000327)
Round face (HP:0000311)
Short nose (HP:0003196)
Broad foot (HP:0001769)
Delayed puberty (HP:0000823)
Large earlobe (HP:0009748)
Short foot (HP:0001773)
Attention deficit hyperactivity disorder (HP:0007018)
Single transverse palmar crease (HP:0000954)
Camptodactyly of finger (HP:0100490)
Pes planus (HP:0001763)
Pectus excavatum (HP:0000767)
Short neck (HP:0000470)
X−linked dominant inheritance (HP:0001423)
Tapered finger (HP:0001182)
Intellectual disability, progressive (HP:0006887)
Paraplegia (HP:0010550)
Prominent supraorbital ridges (HP:0000336)
Micropenis (HP:0000054)
Spastic paraplegia (HP:0001258)
Unsteady gait (HP:0002317)
Short 5th finger (HP:0009237)
Severe muscular hypotonia (HP:0006829)
Cholelithiasis (HP:0001081)
Absent speech (HP:0001344)
Aplasia/Hypoplasia of the 5th finger (HP:0006262)
Congenital diaphragmatic hernia (HP:0000776)
Small hand (HP:0200055)
8
6
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IsMale
Is10ppm
IsCd