**Appendix 1: ICD-9 and ICD-10 developmental disabilities and related codes** (1)

|  |  |
| --- | --- |
| **Code** | **Label** |
| **ICD-9** |
| 299-299.99 | Pervasive development disorders (e.g., autism) |
| 317–317.99 | Mental retardation |
| 318–318.99 | Mental retardation |
| 319–319.99 | Mental retardation |
| 758.0–758.39 | Chromosomal anomalies for which a developmental disability is typically present  |
| 758.5 | Other conditions due to autosomal anomalies |
| 758.8, 758.89 | Other conditions due to chromosome anomalies (do not include 75881) |
| 758.9 | Conditions due to anomaly of unspecified chromosome |
| 759.5 | Tuberous sclerosis |
| 759.81 | Other and unspecified congenital anomalies: Prader-Willi syndrome |
| 759.821 | Other and unspecified congenital anomalies: de Lange syndrome (include only if 6 digits exist, i.e., do not include 759.82) |
| 759.827 | Other and unspecified congenital anomalies: Seckel syndrome (include only if 6 digits exist) |
| 759.828 | Other and unspecified congenital anomalies: Smith-Lemli-Opitz syndrome (include only if 6 digits exist) |
| 759.83 | Other and unspecified congenital anomalies: Fragile X syndrome |
| 759.874  | Other and unspecified congenital anomalies: Beckwith-Wiedemann syndrome (include only if 6 digits exist) |
| 759.875 | Other and unspecified congenital anomalies: Zellweger syndrome (include only if 6 digits exist) |
| 759.89 | Other and unspecified congenital anomalies: other (e.g., Menkes disease, Laurence-Moon-Biedl syndrome, Rubinstein-Taybi syndrome) |
| 760.71 | Fetal alcohol syndrome |
| 760.77 | Fetal hydantoin syndrome |
| **ICD-10** |
| F700 | Mild mental retardation with the statement of no, or minimal, impairment of behaviour |
| F701 | Mild mental retardation, significant impairment of behaviour requiring attention or treatment |
| F708 | Mild mental retardation, other impairments of behaviour |
| F709 | Mild mental retardation without mention of impairment of behaviour |
| F710 | Moderate mental retardation with the statement of no, or minimal, impairment of behaviour |
| F711 | Moderate mental retardation, significant impairment of behaviour requiring attention or treatment |
| F718 | Moderate mental retardation, other impairments of behaviour |
| F719 | Moderate mental retardation without mention of impairment of behaviour |
| F720 | Severe mental retardation with the statement of no, or minimal, impairment of behaviour |
| F721 | Severe mental retardation, significant impairment of behaviour requiring attention or treatment |
| F728 | Severe mental retardation, other impairments of behaviour |
| F729 | Severe mental retardation without mention of impairment of behaviour |
| F730 | Profound mental retardation with the statement of no, or minimal, impairment of behaviour |
| F731 | Profound mental retardation, significant impairment of behaviour requiring attention or treatment |
| F738 | Profound mental retardation, other impairments of behaviour |
| F739 | Profound mental retardation without mention of impairment of behaviour |
| F780 | Other mental retardation with the statement of no, or minimal, impairment of behaviour |
| F781 | Other mental retardation, significant impairment of behaviour requiring attention or treatment |
| F788 | Other mental retardation, other impairments of behaviour |
| F789 | Other mental retardation without mention of impairment of behaviour |
| F79.0 | Unspecified mental retardation with the statement of no, or minimal, impairment of behaviour |
| F791 | Unspecified mental retardation, significant impairment of behaviour requiring attention or treatment |
| F798 | Unspecified mental retardation, other impairments of behaviour |
| F799 | Unspecified mental retardation without mention of impairment of behaviour |
| F840 | Childhood autism |
| F841 | Atypical autism |
| F843 | Other childhood disintegrative disorder |
| F844 | Overactive disorder associated with mental retardation and stereotyped movements |
| F845 | Asperger's syndrome |
| F848 | Other pervasive developmental disorders |
| F849 | Pervasive developmental disorder, unspecified |
| Q851 | Tuberous sclerosis |
| Q860 | Fetal alcohol syndrome |
| Q861 | Fetal hydantoin syndrome |
| Q871 | Aarskog, Prader-Willi, de Lange, Seckel, etc.  |
| Q8723 | Rubinstein-Taybi syndrome (include only if all 5 digits) |
| Q8731 | Sotos syndrome (include only if all 5 digits) |
| Q878 | Other |
| Q900–Q939 except Q926 | All Down syndrome types, cri du chat, etc., except extra marker chromosomes |
| Q971 | Female with more than three X chromosomes |
| Q992 | Fragile X |
| Q998 | Other specified chromosome abnormalities |
| **DSM-IV[[1]](#footnote-1)** |
| 299-299.80 | Pervasive developmental disorders |
| 317–317.99 | Mental retardation |

1. DSM-IV was the active version at time of cohort creation (FY2009) [↑](#footnote-ref-1)