IV. Syndromes

A. Specified

1. Acrofacial dysostosis 1 (Nager type)
2. Apert
3. Al-Awadi/Raas-Rothschild/Schinzel phocomelia
4. Baller-Gerold
5. Bardet-Biedl (21 types)
6. Beals
7. CLOVES
8. Carpenter
9. Catel-Manzke
10. Cornelia de Lange (5 types)
11. Crouzon
12. Down
13. Ectrodactyly-ectodermal dysplasia-clefting
14. Fanconi pancytopenia
15. Freeman Sheldon
16. Fuhrmann
17. Goltz (focal dermal hypoplasia)
18. Gorlin (basal cell nevus syndrome)
19. Greig cephalopolysyndactyly
20. Hajdu-Cheney
21. Hemifacial microsomia (Goldenhar syndrome)
22. Holt-Oram
23. Lacrimoauriculodentodigital
24. Larsen
25. Laurin-Sandrow
26. Leri-Weill dyschondrosteosis
27. Liebenberg syndrome
28. Moebius sequence
29. Multiple synostoses (4 types)
30. Nail-patella
31. Noonan (2 types)
32. Oculodentodigital dysplasia
33. Orofaciodigital (18 types)
34. Otopalatodigital spectrum (filamin A)
35. Pallister-Hall
36. Pfeiffer
37. Pierre Robin (4 subtypes)
38. Poland
39. Proteus
40. Roberts
41. SC phocomelia
42. Rothmund-Thomson
43. Rubinstein-Taybi
44. Saethre-Chotzen
45. Split hand-foot malformation
46. Thrombocytopenia absent radius (TAR)
47. Townes-Brock
48. Trichorhinophalangeal (3 types)
49. Ulnar-mammary
50. VACTERLS association (3 types)

B. Other