This document enclosed two IPCCC code mappings used to identify patients with congenital non-cardiac comorbidity and preterm birth.

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| **Table 1: Codes to identify patients with premature birth** |
| 030102. Visceral heterotaxy (abnormal arrangement thoraco-abdominal organs), |
| 030109. Position or morphology of thoraco-abdominal organs abnormal |
| 030209. Lung anomaly |
| 030214. Functionally congenital single lung |
| 030305. Tracheobronchial anomaly |
| 030603. Intestines malrotated, |
| 102304. Hereditary disorder associated with heart disease |
| 140101. Chromosomal anomaly |
| 140103. Trisomy 18 - Edwards syndrome |
| 140104. Trisomy 13 - Pataus syndrome |
| 140105. 45XO - Turners syndrome |
| 140121. 22q11 microdeletion - CATCH 22 |
| 140200. Syndrome/association with cardiac involvement |
| 140206. DiGeorge sequence |
| 140210. Friedreich’s ataxia, |
| 140217. Marfan syndrome |
| 140219. Noonan syndrome |
| 140221. Pompe’s disease: glycogen storage disease type IIa, |
| 140228. Tuberous sclerosis |
| 140230. Williams syndrome (infantile hypercalcaemia) |
| 140232. Fetal rubella syndrome |
| 140234. Duchenne’s muscular dystrophy, |
| 140258. Muscular dystrophy, |
| 140262. Ehlers-Danlos syndrome |
| 140266. Alagille syndrome: arteriohepatic dysplasia |
| 140300. Non-cardiac abnormality associated with heart disease |
| 140304. Non-cardiothoracic / vascular abnormality (DESCRIBE) |
| 140306. Cystic fibrosis |
| 140307. Diaphragmatic hernia |
| 140308. Tracheo-oesophageal fistula |
| 140310. Omphalocoele |
| 140311. Duodenal stenosis/atresia |
| 140321. Sickle cell disease |
| 140323. Renal abnormality |
| 140328. Congenital coagulation disorder, |
| 140329. Thoracic / mediastinal abnormality |
| 140333. Microcephaly |
| 140347. Choanal atresia, |
| 140349. Tracheobronchial malacia |
| 140352. Hypothyroidism |
| 140391. Cerebral anomaly |
| 140392. Connective tissue disease, |
| 140409. Kyphoscoliosis |
| 140412. Cleft lip / palate |
| 140485. Loeys-Dietz Syndrome (transforming growth factor beta receptor (TGFBR) gene) |
| 140490. Von Willebrand disease |
| 140540.  Maternally derived fetal disease or syndrome associated with heart disease, |
| 140550. Major anomaly of gastrointestinal system |
| 140601. Multiple congenital malformations |
| 161001. Tracheal stenosis |
| 161009. Tracheal disease |
| 140102. Trisomy 21 - Downs syndrome |

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| **Table 2: Codes to identify patients with premature birth** |
| 102202. Premature birth |
| 102205. Premature birth 32-35 weeks, |
| 102206. Premature birth less than 32 weeks, |