Patient	Consan-guinity	Term at first ultrasound signs (WG)	Ultrasound signs	Pregnancy outcome (GA)	Additional postnatal / postmortem data	Follow-up (Age in month)	reason for refusing data reanalysis
Patient 6	No	30	Polyhydramnios HP:0001561 Short long bone HP:0003026 Macrocephaly HP:0000256	COP (40)	No	M35 No additional finding	not requested
Patient 7	No	16	Short fetal femur length HP:0011428 Single umbilical artery HP:0001195	COP (35)	Hypospadias HP:0000047	M35 No additional finding	not requested
Patient 15	No	20	Abnormal heart morphology HP:0001627	TOP (26)	Microcephaly HP:0000252 Agenesis of corpus callosum HP:0001274 Abnormal cortical gyration HP:0002536 Gray matter heterotopia HP:0002282	NA	loss of follow-up
Patient 17	No	20	Microcephaly HP:0000252 Mild fetal ventriculomegaly HP:0010952	COP (NA)	NA	NA	loss of follow-up
Patient 25	No	18	Tented philtrum HP:0011825 Pulmonary artery atresia HP:0004935 Ventricular septal defect HP:0001629	COP	NA	NA	not requested
Patient 27	No	20	Abnormal posterior cranial fossa morphology HP:0000932	COP (41)	No	NA	loss of follow-up
Patient 31	Yes	20	Intrauterine growth retardation HP:0001511 Dilatation of the bladder HP:0010955 Single umbilical artery HP:0001195	COP (35)	Anal atresia HP:0002023 Rectovaginal fistula HP:0000143 Long eyelashes HP:0000527 Hypertrichosis HP:0000998 Synophrys HP:0000664 Short phalanx of finger HP:0009803	M34 Neurodevelopmental delay HP:0012758	not requested
Patient 34	No	20	Agenesis of corpus callosum HP:0001274 Short middle phalanx of finger HP:0005819	TOP (35)	No	NA	loss of follow-up
Patient 35	No	23	Heterotaxy HP:0030853 Asplenia HP:0001746	COP (40)	No	NA	loss of follow-up
Patient 38	No	20	Short fetal femur length HP:0011428 Echogenic fetal bowel HP:0010943 Previous pregnancy: IUFD and echogenic fetal bowel	COP (38)	Intrauterine growth retardation HP:0001511	NA	loss of follow-up
Patient 41	No	20	Cleft lip HP:0410030 Fetal pyelectasis HP:0010945	COP (40)	No	NA	not requested
Patient 58	No	15	Polyhydramnios HP:0001561 Duodenal stenosis HP:0100867 Preaxial polydactyly HP:0100258	COP (33)	No	M10 No additional finding	not requested
Patient 59	No	21	Anomalous pulmonary venous return HP:0010772 Single umbilical artery HP:0001195	COP (36)	Congenital diaphragmatic hernia HP:0000776 Ventricular septal defect HP:0001629 Arachnodactyly HP:0001166 Abnormal facial shape HP:0001999	NA	not requested
Patient 60	No	16	Partial agenesis of the corpus callosum HP:0001338 Fetal pyelectasis HP:0010945	COP (NA)	NA	NA	not requested
Patient 62	No	20	Intrauterine growth retardation HP:0001511 Ebstein anomaly of the tricuspid valve HP:0010316 Talipes equinovarus HP:0001762	TOP (35)	Dacryocystocele HP:0030752 Abnormal facial shape HP:0001999 Polysplenia HP:0001748	NA	loss of follow-up

					Gray matter heterotopia HP:0002282 Abnormal cortical gyration HP:0002536		
Patient 64	No	10	Cystic hygroma HP:0000476 Intrauterine growth retardation HP:0001511 pulmonary artery hypoplasia Ventricular septal defect HP:0001629	TOP (33)	Abnormal facial shape HP:0001999 Retrognathia HP:0000278 Pterygium HP:0001059 Micromelia HP:0002983 Abnormal toe phalanx morphology HP:0010161 Single transverse palmar crease HP:0000954 Abnormal palate morphology HP:0000174	NA	loss of follow-up
Patient 68	Yes	21	Brachycephaly HP:0000248 Partial agenesis of the corpus callosum HP:0001338 Cerebellar vermis hypoplasia HP:0001320	COP (37)	NA	NA	loss of follow-up
Patient 72	No	25	Polyhydramnios HP:0001561 Lateral ventricular asymmetry HP:0100960 Persistent left superior vena cava HP:0005301 Right aortic arch with retroesophageal left subclavian artery HP:0011598 Ureteral duplication HP:0000073	COP (38)	Cerebellar vermis hypoplasia HP:0001320 Hypoplasia of the corpus callosum HP:0002079	NA	not requested
Patient 77	No	24	Unilateral Cleft lip HP:0410030 Fetal choroid plexus cysts HP:0011426	COP (39)	No	M24 No additional finding	not requested
Patient 80	No	27	Intrauterine growth retardation HP:0001511 Short fetal femur length HP:0011428 Single umbilical artery HP:0001195	COP (37)	Hypospadias HP:0000047, Bifid scrotum HP:0000048 Micropenis HP:0000054	M18 No additional finding	not requested
Patient 88	No	20	Esophageal atresia HP:0002032 Oligodactyly HP:0012165	COP (38)	Absent thumb HP:0009777 Abnormality of the ear HP:0000598 Micrognathia HP:0000347	M20 Neurodevelopmental delay HP:0012758	not requested
Patient 91	No	20	Median cleft upper lip HP:0000161 Hypoplastic spinal processes HP:0008460 Hemivertebrae HP:0002937 Micropenis HP:0000054	TOP (33)	NA	NA	loss of follow-up
Patient 104	No	20	Unilateral renal agenesis HP:0000122 Single umbilical artery HP:0001195, splenic cyst	COP (NA)	NA	NA	loss of follow-up
Patient 106	No	20	Polyhydramnios HP:0001561 Retrognathia HP:0000278 Fetal ascites HP:0001791 Hepatomegaly HP:0002240 Short fetal femur length HP:0011428	COP (39)	No	M17 No additional finding	not requested
Patient 107	No	10	Esophageal atresia HP:0002032 Atrioventricular canal defect HP:0006695 Double outlet right ventricle with subpulmonary ventricular septal defect and pulmonary stenosis HP:0011657 Omphalocele HP:0001539 Single umbilical artery HP:0001195	TOP (24)	Abnormal facial shape HP:0001999 Meckel diverticulum HP:0002245 Polysplenia HP:0001748	NA	not requested
Patient 110	No	16	Hydrops fetalis HP:0001789	TOP (26)	Pulmonary artery hypoplasia HP:0004971 Right aortic arch with retroesophageal left subclavian artery HP:0011598	NA	not requested

					Abnormal facial shape HP:0001999		
Patient 111	No	20	Hand clenching HP:0001188 Talipes equinovarus HP:0001762	TOP (26)	NA	NA	loss of follow-up
Patient 113	No	25	Oligohydramnios HP:0001562 Macroglossia HP:0000158 Fetal ascites HP:0001791	COP (37)	Congenital posterior urethral valve HP:0010957	No	not requested
Patient 119	No	20	Agenesis of corpus callosum HP:0001274	TOP (30)	Hypertrichosis HP:0000998 Macrocephaly HP:0000256 Missing ribs HP:0000921 Supernumerary vertebrae HP:0002946	NA	loss of follow-up
Patient 124	No	20	Polyhydramnios HP:0001561 Tetralogy of Fallot HP:0001636 Esophageal atresia HP:0002032	COP (36)	NA	NA	not requested
Patient 127	No	10	Increased nuchal translucency HP:0010880 Echogenic fetal bowel HP:0010943	COP (39)	No	NA	loss of follow-up
Patient 128	No	19	Agenesis of corpus callosum HP:0001274 Abnormal cortical gyration HP:0002536 Gray matter heterotopia HP:0002282 Interhemispheric cyst HP:0032327	TOP (30)	Polydactyly HP:0010442	NA	loss of follow-up
Patient 135	No	20	Polyhydramnios HP:0001561 Large for gestational age HP:0001520 Congenital diaphragmatic hernia HP:0000776	COP (33)	Neonatal death	NA	not requested
Patient 137	No	10	Cystic hygroma HP:0000476 Abnormality of ductus venosus blood flow HP:0010947	COP (37)	NA	NA	not requested
Patient 146	No	20	Agenesis of corpus callosum HP:0001274	TOP (30)	Bifid uvula HP:0000193 Retrognathia HP:0000278 Hypertrichosis HP:0000998 Abnormal facial shape HP:0001999 Abnormal cortical gyration HP:0002536	NA	not requested
Patient 148	No	20	Polyhydramnios HP:0001561 Esophageal atresia HP:0002032 Unilateral renal agenesis HP:0000122 Single umbilical artery HP:0001195	COP (38)	Ectopic kidney HP:0000086	NA	not requested

COP: Continuation of pregnancy, TOP: termination of pregnancy