

Type of CHD	n (%)	Antenatal diagnosis						Outcome					
		Aneuploidy screening (i.e. FTS or NIPT)		Confirmatory genetic testing (i.e. Amniocentesis, CVS, or postnatal)		Extracardiac fetal anomaly (%)		Fetal echocardiography		Lost to follow-up (%)	Second-trimester pregnancy loss (%)	TOP (%)	Live birth
		Number	High risk (%)	Number	Abnormal (%)	Performed	Concordance*	Number	TOP (%)	Number	Neonatal outcome	Number	Neonatal outcome
TGA	18 (16.5)	Total: 8 FTS: 2 NIPT: 3 Cont: 3	0 (0)	Total: 10 AN: 6 PN: 4	0 (0)	4 (22.2)	12	12 (100.0)	5 (27.8)	0 (0)	2 (11.1)	11 (61.1)	A: 0 B: 3 C: 0 D: 7 E: 1
DORV	10 (9.2)	Total: 5 FTS: 3 NIPT: 1 Cont: 1	2 (40)	Total: 5 AN: 3 PN: 2	1 (20.0)	0 (0)	4	3 (75.0)	3 (30.0)	0 (0)	2 (20.0)	5 (50.0)	A: 1 B: 0 C: 1 D: 3 E: 0
Tetralogy of Fallot	15 (13.8)	Total: 5 FTS: 2 NIPT: 3 Cont: 0	5 ^S (100)	Total: 14 AN: 11 PN: 6 [†] Cont: 0	6 (42.9) [‡]	4 (26.7)	13	10 (76.9)	2 (13.3)	0 (0)	3 (20.0)	10 (66.7)	A: 1 B: 2 C: 6 D: 1 E: 0
Truncus arteriosus	5 (4.6)	Total: 1 FTS: 1 NIPT: 0 Cont: 0	0 (0)	Total: 4 AN: 4 PN: 0	1 (25.0)	1 (20.0)	3	3 (100.0)	1 (20.0)	0 (0)	2 (40.0)	2 (40.0)	A: 2 B: 0 C: 0 D: 0 E: 0
Interrupted aortic arch	2 (1.8%)	Total: 0 FTS: 0 NIPT: 0 Cont: 0	-	Total: 0 AN: 0 PN: 0	-	0 (0)	0	-	1 (50.0)	0 (0)	1 (50.0)	0 (0)	A: 0 B: 0 C: 0 D: 0 E: 0
Coarctation of aorta	19 (17.4)	Total: 10 FTS: 5 NIPT: 3 Cont: 2	1 (10.0)	Total: 10 AN: 7 PN: 3	1 (10.0)	0 (0)	12	10 (83.3)	4 (21.1)	0 (0)	0 (0)	15 (78.9)	A: 2 B: 3 C: 4 D: 6 E: 0
UVH	15 (13.8)	Total: 6 FTS: 4 NIPT: 1 Cont: 1	0 (0)	Total: 6 AN: 5 PN: 1	1 (16.7)	1 (6.67)	7	7 (100.0)	5 (33.3)	1 (6.67)	6 (40.0)	3 (20.0)	A: 1 B: 1 C: 0 D: 0 E: 1

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		Number	High risk (%)	Number	Abnormal (%)	Number	Concordance* (%)	Performed	Concordance* (%)				
										Number (%)	Neonatal outcome		
Critical pulmonary valve stenosis	6 (5.5%)	Total: 4	0 (0)	Total: 4	0 (0)	0 (0)	5	5 (100.0)	1 (16.7)	0 (0)	5 (83.3)	A: 0 B: 2 C: 0 D: 3 E: 0	
		FTS: 3		AN: 3									
		NIPT: 0		PN: 1									
		Cont: 1											
AVSD	6 (5.5%)	Total: 4	2 (40.0)	Total: 4	3 (75.0)	2 (33.3)	5	5 (100.0)	1 (16.7)	0 (0)	5 (83.3)	A: 1 B: 0 C: 1 D: 3 E: 0	
		FTS: 1		AN: 1									
		NIPT: 3		PN: 3									
		Cont: 0											
Large ventricular septal defect	6 (5.5%)	Total: 2	0 (0)	Total: 4	1 (25.0)	1 (16.7)	5	5 (100.0)	1 (16.7)	0 (0)	5 (83.3)	A: 0 B: 1 C: 1 D: 3 E: 0	
		FTS: 1		AN: 2									
		NIPT: 1		PN: 2									
		Cont: 0											
Tricuspid atresia	4 (3.7%)	Total: 3	2 (66.7)	Total: 3	1 (33.3)	1 (25.0)	2	1 (50.0)	0 (0)	1 (25.0)	2 (50.0)	A: 0 B: 1 C: 1 D: 0 E: 0	
		FTS: 3		AN: 2									
		NIPT: 0		PN: 2†									
		Cont: 0											
Critical or severe aortic stenosis	3 (2.8%)	Total: 2	0 (0)	Total: 2	0 (0)	0 (0)	2	1 (50.0)	0 (0)	0 (0)	2 (66.7)	A: 0 B: 0 C: 0 D: 2 E: 0	
		FTS: 1		AN: 1									
		NIPT: 1		PN: 1									
		Cont: 0											
Total	109	Total: 50	12 (24.0)	Total: 66	15 (23.1)	14 (12.8)	70	62 (88.6)	24 (22.0)	2 (1.8)	65 (59.6)	A: 8 B: 13 C: 14 D: 28 E: 2	
		FTS: 26		AN: 45									
		NIPT: 16		PN: 25†									
		Cont: 8											

AN: antenatal, AVSD: atrioventricular septal defect, Cont: contingent, CVS: chorionic villus sampling, DORV: double outlet right ventricle, FTS: first trimester screening, NIPT: non-invasive prenatal testing, PN: postnatal, TGA: transposition of great arteries, TOP: termination of pregnancy, UVH: univentricular hearts

A: infant death, B: mild medical/developmental issues, C: major medical or developmental issues, D: no medical or developmental issues, E: postnatal lost to follow-up

* Where formal fetal echocardiography was performed, concordance was met when the obstetric fetal anatomical scan made the same major CHD diagnosis without missing any other major CHD

† Three cases of tetralogy of Fallot and one case of tricuspid atresia underwent postnatal genetic testing in addition to antenatal invasive genetic testing for the purpose of additional genetic testing

** One case of tetralogy of Fallot had a variant of unknown significance on chromosomal microarray analysis. These variants were also detected in phenotypically normal parents.

‡ One aneuploidy screening result was deemed high risk, as it comprised two consecutive inconclusive NIPT results.