Correction: *Genetic characterisation of 22q11.2 variations and prevalence in patients with congenital heart disease*

Hou H, Chen H, Wang X, *et al.* Genetic characterisation of 22q11.2 variations and prevalence in patients with congenital heart disease. *Arch Dis Child* 2020;105:367-74.

During production of this paper, two errors were introduced in the numbers of [table 1](#). The numbers highlighted in red should be 17.3 and 19.2 respectively instead of 1.7 and 1.9.

Table 1 Phenotypic characteristics of all patients with CHD and the percentage of 22q11.2 deletions or amplifications in each group

Phenotypic characteristics (heart disease)	Patients (n)	22q11.2 positive patients, n (%)
VSD	192	15 (7.8)
Isolated VSD	140	6 (4.3)
VSD with other defects	52	9 (17.3)
ASD	91	11 (12.1)
Isolated ASD	65	6 (9.2)
ASD with other defects	26	5 (19.2)
PDA	88	12 (13.6)
Isolated PDA	49	5 (10.2)
PDA with other defects	39	7 (17.9)
TOF	26	6 (23.1)
DCRV	7	2 (28.6)
DORV	4	1 (25)
RVOTS	26	4 (15.4)
PS	12	2 (16.7)
PH	100	18 (18)

ASD, atrial septal defect; CHD, congenital heart disease; DCRV, double-chamber right ventricle; DORV, double-outlet right ventricle; PDA, patent ductus arteriosus; PH, pulmonary hypertension; PS, pulmonary stenosis; RVOTS, right ventricular outflow tract stenosis; TOF, tetralogy of Fallot; VSD, ventricular septal defect.

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Arch Dis Child 2020;105:e7. doi:10.1136/archdischild-2018-316634corr1