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Vol.57 NO1 (2017)	Exome sequencing identifies a novel nonsense mutation of HOXD13 in a Chinese family with synpolydactyly	0
	Dynamics of gyrification in the human cerebral cortex during development	0
	Evaluation of the maxillofacial morphological characteristics of Apert syndrome infants	1
	Foxc2 *CreERT2* knock-in mice mark stage-specific *Foxc2* -expressing cells during mouse organogenesis	0
	Cartilage-hair hypoplasia associated with isolated hypoganglionosis: A case report	0
	Impact of the introduction of Non-invasive prenatal genetic testing on invasive tests: A single-center study in Japan	0
	Prenatal diagnosis of Fraser syndrome using routine ultrasound examination, confirmed by exome sequencing: Report of a novel homozygous missense *FRAS1* mutation	0
Vol.57 NO2 (2017)	Proposal for supplemental intake of folic acid to reduce the risk of neural tube defects	2
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	Magnetic resonance imaging based correlation analysis between calcarine sulcus development and isolated fetal ventriculomegaly	2
	Locus on chromosome 16 is significantly associated with increased tendency to lose pups in females of the RR/Sgn inbred mouse strain	0
	Miller-Dieker Syndrome with unbalanced translocation 45, X, psu dic(17;Y)(p13;p11.32) detected by fluorescence in situ hybridization and G-banding analysis using high resolution banding technique	0
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	Discordant ventriculo-arterial connections, or “transposition”, are not necessarily an essential part of isomerism	0
Vol.57 NO3 (2017)	Selective serotonin reuptake inhibitors and risk of major congenital anomalies for pregnancies in Japan: A nationwide birth cohort study of the Japan Environment and Children's Study	0
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	Variable presentation of Fraser syndrome in two fetuses and a novel mutation in *FRAS1*	0
	A Say-Barber-Biesecker-Young-Simpson variant of Ohdo syndrome with a *KAT6B* 10-base pair palindromic duplication: A recurrent mutation causing a severe phenotype mixed with genitopatellar syndrome	0
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	Bilateral choanal atresia in an adult woman with pycnodysostosis	0
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