Hindawi
Case Reports in Genetics
Volume 2021, Article ID 9897523, 2 pages
https://doi.org/10.1155/2021/9897523

Corrigendum

Corrigendum #2 to "Eye Manifestations of Shprintzen–Goldberg Craniosynostosis Syndrome: A Case Report and Systematic Review"

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Received 25 March 2021; Accepted 25 March 2021; Published 31 August 2021

In the article titled "Eye Manifestations of Shprintzen–Goldberg Craniosynostosis Syndrome: A Case Report and Systematic Review" [1], the authors have identified additional errors that require correction. These errors are resulting from the misinterpretation of data obtained from Doyle et al. [2].

Supplemental Table 1(a) should be corrected as follows: to remove "ectopia lentis" from patients 4, 7, and 8.

In the Discussion section, the sentence, "though ectopia lentis was reported in three patients, our patient did not exhibit any degree of apparent lens dislocation," should be deleted. In the abstract, "Ocular manifestations may include hypertelorism, downslanting palpebral fissures, proptosis, myopia, and ectopia lentis" should be changed to "Ocular manifestations may include hypertelorism, downslanting palpebral fissures, proptosis, and myopia."

Supplemental Table 1: Summary of 45 SGS patients with SKI gene mutations.

(a)

	Doyle,et al. [2012]									
Patient	1	2	3	4	5	6	7	8	9	10
SKI gene pathogenic variant	c.347G>A(p.Gly116Glu)	c.349G>C(p.Gly117Arg)	c.101G>A(p.Gly34Asp)	c.94C>G(p.Leu32Val)	c.94C>G(p.Leu32Val)	c.100G>A(p.Gly34Ser)	c.100G>T(p.Gly34Cys)	c.103C>T(p.Pro35Ser)	c.283_291del(p.Asp95_Ser97del)	c.62T>G(p.Leu21Arg)
Inheritance	de novo	de novo	de novo	de novo	de novo	de novo	de novo	de novo	de novo	de novo
Gender	F	M	M	M	F	M	F	M	M	F
Age(Years)	43	6	16	12	22	21	2	6	5	4
Ocular findings	Hypertelorism,down-slanting eyes, proptosis	Hypertelorism,down-slanting eyes	Hypertelorism,down-slanting eyes, proptosis	Hypertelorism,down-slanting eyes, proptosis	Hypertelorism,down-slanting eyes, proptosis	Hypertelorism, proptosis	Hypertelorism,down-slanting eyes, proptosis	Hypertelorism,down-slanting eyes, proptosis	Hypertelorism,down-slanting eyes, proptosis	Hypertelorism,down-slanting eyes, proptosis
Dysmorphic features	+	+	+	+	+	+	+	+	+	+
Cardiac anomalies	Mitral value prolapse, aortic root dilatation	Mitral value prolapse, aortic root dilatation	Mitral value prolapse, aortic root dilatation	aortic root dilatation	aortic root dilatation	aortic root dilatation	aortic root dilatation	-	Mitral value prolapse	Mitral value prolapse, aortic root dilatation, arterial tortuosity
Musculoskeletal anomalies	+	+	+	+	+	+	+	+	+	+
Neurological anomalies	+	+	+	+	+	+	+	+	+	+
Developmental delay	+	+	+	+	+	+	+	+	+	+
Other	Splenic artery aneurysm	Broad/bifiduvula	Club foot deformity	Cleft palate			Cleft palate	Cleft palate, Club foot deformity, Splenic artery aneurysm with spontaneous rupture		Broad/bifiduvula

References

[1] H. C. Jamie, L. Rachel, G. Rachel, N. C. Tahnee, H. Anna, and L. C. Natario, "Eye Manifestations of Shprintzen–Goldberg Craniosynostosis Syndrome: A Case Report and Systematic Review," *Case Reports in Genetics*, vol. 2020, Article ID 7353452, 2020.

[2] A. J. LopezCourcet, J. J. Doyle, S. L. Bessling et al., "Mutations in the TGF-β Repressor SKI Cause Shprintzen-Goldberg Syndrome with Aortic Aneurysm," *Nature Genetics*, vol. 44, no. 11, pp. 1249–1254, 2012.