



Correction

Correction: Spenger et al. Glutaric Aciduria Type I Missed by Newborn Screening: Report of Four Cases from Three Families. *Int. J. Neonatal Screen.* 2021, 7, 32

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There was an error in the original publication [1]. In patient 2 and 3, a mutation (p.Arg257Glu) is mistakenly documented. Instead, the mutation is p.Arg257Gln. A correction has been made to Section 2. Case Reports, 2.2. Case 2, Paragraph 3; Section 2. Case Reports, 2.3. Case 3, Paragraph 1; Section 3. Results, Paragraph 1; Section 4. Discussion, Paragraph 1:

Section 2. Case Reports, 2.2. Case 2, Paragraph 3: At the age of 10 years, genetic testing revealed previously described compound heterozygous variants in *GCDH* (p.Arg257Gln, p.Met405Val) [11,12], confirming the diagnosis of GA-1.

Section 2. Case Reports, 2.3. Case 3, Paragraph 1: After the diagnosis was made (compound heterozygous variants in *GCDH*: p.Arg257Gln and p.Met405Val), she was admitted during an infection with poor oral intake, and a clear deterioration of the movement disorder was observed.

Section 3. Results, Paragraph 1: Patients 2 and 3 were compound heterozygous for p.Arg257Gln and p.Met405Val, patient 1 was compound heterozygous for p.Gly241Val and p.Gly390Ala, and patient 4 was compound heterozygous for p.Arg257Trp and the novel variant p.Lys170Asn.

Section 4. Discussion, Paragraph 1: Patients 2 and 3 are compound heterozygous for a previously described severe mutation (p.Arg257Gln) with 0% residual activity [11] and a milder mutation (p.Met405Val) with 4–25% residual activity that is more prevalent in African patients [12].



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The authors apologize for any inconvenience caused and state that the scientific conclusions are unaffected. The original publication has also been updated.

Reference

1. Spenger, J.; Maier, E.M.; Wechselberger, K.; Bauder, F.; Kocher, M.; Sperl, W.; Preisel, M.; Schiergens, K.A.; Konstantopoulou, V.; Röschinger, W.; et al. Glutaric Aciduria Type I Missed by Newborn Screening: Report of Four Cases from Three Families. *Int. J. Neonatal Screen.* 2021, 7, 32. [CrossRef] [PubMed]