

## Corrigendum

## Corrigendum to “Frequent intragenic microdeletions of elastin in familial supravalvular aortic stenosis.” [Int. J. Cardiol., 274 (2019) 290–295]



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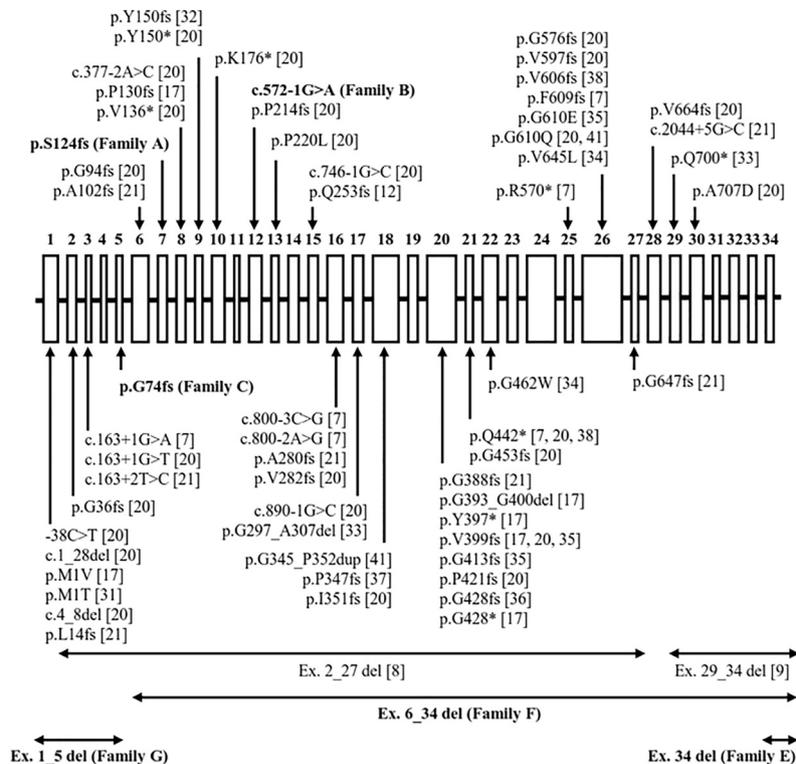
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The authors regret that there was an error in the description of microdeletion in the family F in Table 1 and the Results section. The microdeletion in the family F was detected in the exons 6–33 as shown in Fig. 2, not exons 5–33 as published in Table 1 and Results section.

In addition, we noted errata in the Fig. 3 of the manuscript. This figure summarized the previously reported data and the errata of the description of mutation and the numbering of references were noted. We incorporated the update for Fig. 3 and its legend as follows:

The authors would like to apologise for any inconvenience caused.



**Fig. 3.** ELN cDNA showing the exons and mutations detected. This figure summarizes previously reported and newly identified mutations for familial and sporadic SVAS. The numbers above open boxes indicate the exon numbers. The present findings are shown in bold letters. Although all the data besides this figure were shown by NM\_000501, our results and previously reported data in this figure were shown using NM\_001278939 as a reference sequence.

DOI of original article: <https://doi.org/10.1016/j.ijcard.2018.09.032>.

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