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## Corrigendum: Current phenotypic and genetic spectrum of syndromic deafness in Tunisia: paving the way for precision auditory health

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# Corrigendum: Current phenotypic and genetic spectrum of syndromic deafness in Tunisia: paving the way for precision auditory health

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## KEYWORDS

syndromic deafness (SD), spectrum of SDs, next-generation sequencing (NGS), early detection, under-diagnosis, Tunisia

**A Corrigendum on**

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In the published article, there was an error in the legend for Figure 4 as published. The gene names corresponding to color degradation for the Waardenburg syndrome pie chart are incorrect.

The corrected legend appears below: The SOX10 gene should be at the top (darkest blue), and the EDNRB gene should be at the bottom (lightest blue). Consequently, the number of patients should be 8 for the SOX10 gene and 2 for the EDNRB gene.

In addition, there was an error in the caption of Figure 4 in the published version of the manuscript: “FIGURE 4: Genetic and mutational spectrum of syndromic deafness in the Tunisian population. Deafness syndromes with known molecular etiologies are represented by pie charts. Each syndromic deafness is indicated with a different color. Color degradation has been adopted to specify

the gene(s) associated with each syndrome. The number of variants identified in each gene is provided.”

The corrected Figure 4 appears below:

The authors apologize for these errors and state that this does not change the scientific conclusions of the article in any way. The original article has been updated.

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