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## Three novel CYP21A2 mutations and their protein modelling in patients with classical 21-hydroxylase deficiency from northeastern Iran

Baradaran-Heravi, A.<sup>b</sup>, Vakili, R.<sup>a</sup>, Robins, T.<sup>d</sup>, Carlsson, J.<sup>e</sup>, Ghaemi, N.<sup>c</sup>, ARabi, A.<sup>b</sup>, Abbaszadegan, M.R.<sup>ab</sup>

<sup>a</sup> Division of Human Genetics, Bu-Ali Research Institute, **Mashhad University of Medical Sciences**, PO Box 91774, **Mashhad**, Iran

<sup>b</sup> Division of Human Genetics, Immunology Research Centre, Bu-Ali Research Institute, **Mashhad**, Iran

<sup>c</sup> Department of Paediatrics, Imam Reza Hospital, **Mashhad University of Medical Sciences (MUMS)**, **Mashhad**, Iran

<sup>d</sup> Department of Molecular Medicine and Surgery, Centre of Molecular Medicine (CMM) L8:02, Karolinska Institutet **University Hospital**, Stockholm, Sweden