Errata

Küçükemre-Aydın B, Öğrendil-Yanar Ö, Bilge I, Baş F, Poyrazoğlu Ş, Yılmaz A, Emre S, Bundak R, Saka N, Darendeliler F.

An easily missed diagnosis: 17-alpha-hydroxylase/17,20-lyase deficiency. Turk J Pediatr 2015; 57: 277-281, one correction is necessary in order to prevent misinterpretations of the nomenclature of the described mutation:

Page 279, left column, last paragraph, line 6: c.1307G>A should read c.1306G>A.