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Manuscript Type: Case Report

Point by point response to the Reviewers

Reviewer: 00631992

The authors present two brothers suffering from a rare genetic disorder with autosomal recessive inheritance, of nephrologic interest. The article is interesting, well written and certainly deserves to be published. Just a very few remarks: It would be appropriate to mention the possibility that point mutations that damage splice sites cause the phenomenon of “exon skipping" (see Tacheuchi et al., 2015).

Re: We edited the manuscript according to the reviewer's suggestions and we discussed and cited the reference (Tacheuchi et al., 2015) (page 8, lanes 8-11, new version of the manuscript).

At page. 4 (Introduction): line 6, "experienced" should be "experience"; lines 26 and 27, review the phrase “this mutation is a disease causative of GS.” At page. 8 (Discussion): line 7, "both patients, inherited" should be "both patients inherited"; line 11“The second mutation, inherited from the mother, carried by our patients, is a ...” would be better “The second mutation carried by our patients, inherited from the mother, is a ...”

Re: We edited the manuscript following all the reviewer's suggestions.

Reviewer: 00503254

No comments

Reviewer: 00503255

No comments