Author's response to reviews

Title: Identification of a novel de novo mutation in the NIPBL gene in an Iranian patient with Cornelia de Lange Syndrome: a case report

Authors:

    Hamid Galehdari (galehdari187@yahoo.com)
    Roya Monajemzadeh (Monajemzadeh@gmail.com)
    Gholamreza Mohammadian (Mohammadian2000@yahoo.com)
    Mohammad Pedram (m_Pedram_2007@yahoo.com)
    Habibolah Nazem (Hnazem@gmail.com)

Version: 10 Date: 4 December 2010

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Covering letter

Dear Editor of the JMCR

The manuscript: "Identification of a novel de novo mutation in the NIPBL gene in an Iranian patient with Cornelia de Lange Syndrome: a case report", MS: 9582203535278266

Was changed as follow:

1) Abstract:
   a. The gene name SMC1 was corrected.
   b. The phrase *are responsible* was replaced with *have been identified*.
   c. The phrase *presumably* was added in the first paragraph of the conclusion.
   d. line 8 - height was replaced to length.

2) Introduction
   a. The correct name *NIPBL* was used in the last sentence.
   b. Line 4 - estimated prevalence was changed to 1.6 - 2.2 per 100,000 births (Barisic et al 2008).

3) Declaration of competing interests was corrected.

4) Figures
   a. Better images of the patient’s hand were added.
   b. Figure legends were checked.

5) Case presentation
   a. Line 8 - cryptorchidism was corrected.
   b. Line 13 - length was corrected.
   c. The ethnicity of patient was added.
6) References were checked and rearranged.

Kind regards

Hamid Galehdari
Corresponding Author In behalf of the Co-Authors