

PEER-REVIEW REPORT

Name of journal: World Journal of Clinical Cases

Manuscript NO: 58819

Title: Discontinuous polyostotic fibrous dysplasia with multiple systemic disorders and unique genetic mutations: A case report

Reviewer's code: 03850089

Position: Editorial Board

Academic degree: MD, PhD

Professional title: Emeritus Professor

Reviewer's Country/Territory: Brazil

Author's Country/Territory: China

Manuscript submission date: 2020-08-25

Reviewer chosen by: Le Zhang

Reviewer accepted review: 2020-09-02 13:14

Reviewer performed review: 2020-09-05 22:21

Review time: 3 Days and 9 Hours

Scientific quality	<input type="checkbox"/> Grade A: Excellent <input type="checkbox"/> Grade B: Very good <input checked="" type="checkbox"/> Grade C: Good <input type="checkbox"/> Grade D: Fair <input type="checkbox"/> Grade E: Do not publish
Language quality	<input type="checkbox"/> Grade A: Priority publishing <input type="checkbox"/> Grade B: Minor language polishing <input checked="" type="checkbox"/> Grade C: A great deal of language polishing <input type="checkbox"/> Grade D: Rejection
Conclusion	<input type="checkbox"/> Accept (High priority) <input type="checkbox"/> Accept (General priority) <input checked="" type="checkbox"/> Minor revision <input type="checkbox"/> Major revision <input type="checkbox"/> Rejection
Re-review	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No
Peer-reviewer statements	Peer-Review: <input type="checkbox"/> Anonymous <input checked="" type="checkbox"/> Onymous Conflicts-of-Interest: <input type="checkbox"/> Yes <input checked="" type="checkbox"/> No

SPECIFIC COMMENTS TO AUTHORS

Manuscript ID: 58819: Discontinuous Polyostotic Fibrous Dysplasia with Unique Genetic Mutations: Case Report and Literature Review In the manuscript, the authors present a case of Polyostotic Fibrous Dysplasia (PFD) with unusual characteristics from a clinical, laboratory, and genetic research profile. In a review, it reports the genetic relationship of the syndrome with the GNAS gene, which was not observed in that case, but with a history of parents with inbreeding, a high increase in cortisol, and changes in HSPG and RIMS. It cited the alterations that can be observed in the described genetic alterations and conclude that they are responsible for what is observed. Also the patient were refractory to the use of bisphosphonates, which is described in the literature in some other cases. Turrion -Nieves AI, Medicina Clinica, 2020. I suggest that the title of the manuscript be modified and emphasized the observed because it is a different syndrome from the commonly observed. I suggest that the reference concerning CARE be placed at the end of the introduction as it is the method followed for writing the article. I request that the text of the manuscript be retyped (errors) and the English presented to me as I am not a native speaker of the English language at certain times or in parts of the article, the understanding was extremely difficult. In conclusion, I consider the manuscript important and should be published for the review work, which should be emphasized and highlighted.