|  |  |
| --- | --- |
|  | |
| Journal Name: | [**Asian Journal of Case Reports in Medicine and Health**](https://journalajcrmh.com/index.php/AJCRMH) |
| Manuscript Number: | **Ms\_AJCRMH\_134619** |
| Title of the Manuscript: | **Werner syndrome revealed by a severe metabolic acute pancreatitis: A unique clinical observation** |
| Type of the Article | **Case Report** |

|  |  |  |
| --- | --- | --- |
| PART 1: Comments | | |
|  | Reviewer’s comment **Artificial Intelligence (AI) generated or assisted review comments are strictly prohibited during peer review.** | Author’s Feedback *(Please correct the manuscript and highlight that part in the manuscript. It is mandatory that authors should write his/her feedback here)* |
| **Please write a few sentences regarding the importance of this manuscript for the scientific community. A minimum of 3-4 sentences may be required for this part.** | **The article has comprehensively portrayed characteristics f Werner Syndrome. This will help generate awareness amongst the physician about a rare entity. The author has described most of the clinical features associated with the syndrome.** | Absolutely yes. |
| **Is the title of the article suitable?**  **(If not please suggest an alternative title)** | **Yes** | Thank you |
| Is the abstract of the article comprehensive? Do you suggest the addition (or deletion) of some points in this section? Please write your suggestions here. | **Yes** | Thank you |
| Is the manuscript scientifically, correct? Please write here. | Yes | Thank you |
| **Are the references sufficient and recent? If you have suggestions of additional references, please mention them in the review form.** | Yes | Thank you |
| Is the language/English quality of the article suitable for scholarly communications? | yes | **Thank you** |
| Optional/General comments | **The diagnosis of WS is not supported by genetic testing in patient. The author must emphasize the fact that no testing was done, and the diagnosis was based on clinical suspicion.** | Indeed, the diagnosis was based on clinical findings, especially since the patient had a brother with the same symptoms.  Unfortunately, genetic testing could not be performed due to lack of resources. This was clearly stated in the medical observation. |

|  |  |  |
| --- | --- | --- |
| **PART 2:** | | |
|  | Reviewer’s comment | Author’s comment *(if agreed with the reviewer, correct the manuscript and highlight that part in the manuscript. It is mandatory that authors should write his/her feedback here)* |
| **Are there ethical issues in this manuscript?** | *(If yes, Kindly please write down the ethical issues here in detail)* | NO |