The present case report article describes a new case of an extremely rare inheritable autosomal recessive rheumatic disease, the progressive pseudorheumatoid dysplasia, associated to mutations in WISP3, a member of the connective tissue growth factor family.

This clinical description contains two points of originality with respect to other cases of PPD previously described in the literature. Firstly, the delayed-onset (15 years) is clearly later to that reported before, between three and six years. Secondly, the mutational analysis report a novel mutation in exon 4 of the WIPS3 gene consistent in a frameshift mutation (c.670dupA).

Although an English editing is recommended, the article is concisely written and the conclusions are clear. The methodology used for the genetic study is adequate and the radiological pictures in the figures are of good quality.

For all these reasons the article meets the criteria of quality and originality enough to be published in BMC Medical Genetics.
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