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# Author Correction: Analysis and visualisation of electronic health records data to identify undiagnosed patients with rare genetic diseases

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Correction to: *Scientific Reports* <https://doi.org/10.1038/s41598-024-55424-8>, published online 01 March 2024

The original version of this article contained an error in the title of the paper, where the term “Cluster” was erroneously added.

In addition, in the abstract,

“Data mining in the form of cluster analysis and visualisation, was performed on a database containing de-identified health records of 1.28 million patients across 3 major hospitals in Singapore, in a bid to improve the diagnostic process for patients who are living with an undiagnosed rare disease, specifically focusing on Fabry Disease and Familial Hypercholesterolaemia (FH).”

now reads:

“Data analysis in the form of visualisation and statistical testing, was performed on a database containing de-identified health records of 1.28 million patients across 3 major hospitals in Singapore, in a bid to improve the diagnostic process for patients who are living with an undiagnosed rare disease, specifically focusing on Fabry Disease and Familial Hypercholesterolaemia (FH).”

The original article has been corrected.



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