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Corrigendum: Heterogeneous phenotype of a Chinese Familial WHIM syndrome with CXCR4^{V340fs} gain- of-function mutation

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CXCR4 variant, gain-of-function, inborn error of immunity, WHIM syndrome,
heterogeneous phenotype

A Corrigendum on

**Heterogeneous phenotype of a Chinese Familial WHIM syndrome with
CXCR4^{V340fs} gain-of-function mutation**

By Huang Y, Li L, Chen R, Yu L, Zhao S, Jia Y, Dou Y, Zhang Z, An Y, Tang X, Zhao X and Zhou L
(2024) *Front. Immunol.* 15:1460990. doi: 10.3389/fimmu.2024.1460990

In the published article, there was an error. A typo was made.

A correction has been made to **Abstract**, *Results*, Paragraph 1. This sentence
previously stated:

“We provide and in-depth analysis of their clinical, genetic, immunological and
treatment characteristic, noting that these patients exhibited an atypical clinical
phenotype when compared to reported CXCR4R334X patients.”

The corrected sentence appears below:

“We provide an in-depth analysis of their clinical, genetic, immunological and
treatment characteristic, noting that these patients exhibited an atypical clinical
phenotype when compared to reported CXCR4R334X patients.”

1. In the published article, there was an error. A typo was made.

A correction has been made to **Introduction**, Paragraph 2. This sentence
previously stated:

“Thirty-seven distinct CXCR4 variants have been identified, which including eight nonsense variants, twenty-seven frameshift variants, and two missense variants.”

The corrected sentence appears below:

Thirty-seven distinct CXCR4 variants have been identified, which including eight nonsense variants, twenty-seven frameshift variants, and two missense variants.”

2. In the published article, there was an error. Omission of important information.

A correction has been made to **Results**, *Clinical manifestations of the family with WHIM syndrome*, Paragraph 3. This sentence previously stated:

“At the age of, P2 was hospitalized due to edema, and subsequent tests showed proteinuria, hyperlipidemia, and hypoproteinemia, leading to a diagnosis of nephrotic syndrome.”

The corrected sentence appears below:

“At the age of 4, P2 was hospitalized due to edema, and subsequent tests showed proteinuria, hyperlipidemia, and hypoproteinemia, leading to a diagnosis of nephrotic syndrome.”

3. In the published article, there was an error. A typo was made.

A correction has been made to **Results**, *Decreased surface CXCR4 expression on CD8+T cells and B cells from patients*, Paragraph 1. This sentence previously stated:

“Interestingly, P1-P4 exhibited relatively higher levels of CXCR4 lexpession, particularly in pediatric patients P1 and P2 (**Supplementary Figure SA**).”

The corrected sentence appears below:

“Interestingly, P1-P4 exhibited relatively higher levels of CXCR4 expression, particularly in pediatric patients P1 and P2 (**Supplementary Figure SA**).”

4. In the published article, there was an error. A typo was made.

A correction has been made to **Discussion**, Paragraph 4. This sentence previously stated:

“n our cohort, all four patients exhibited varying degrees of impaired CXCR4 internalization; pediatric patients showed reduced internalization, while adult patients demonstrated no internalization.”

The corrected sentence appears below:

“In our cohort, all four patients exhibited varying degrees of impaired CXCR4 internalization; pediatric patients showed reduced internalization, while adult patients demonstrated no internalization.”

The authors apologize for these errors and state that this does not change the scientific conclusions of the article in any way. The original article has been updated.

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