## CORRECTION

doi: 10.1136/ard.2004.025767corrl

Single nucleotide polymorphisms in the gene encoding the major histocompatibility complex class II transactivator
(CIITA) in systemic lupus erythematosus (Koizumi K, Okamoto H, Iikuni N, Nakamura T, Kawamoto M, Momohara S, Ichikawa N, Furuya T, Kotake S, Taniguchi A, Yamanaka H, Kamatani N. Ann Rheum Dis 2005;64:947-50.)
The authors have provided a new figure to replace figure 1 of this article. The figure is reproduced below.

Corrections printed in the journal also appear on the Annals website
http://www.annrheumdis.com and are linked to the original publication

A

| Promoter III | Coding sequence |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
| nt | nt | nt | nt | nt |
| $\mathrm{No}-155$ | No 1614 | No 2509 | No 2536 | No 2791 |
| (III) | (C1) | (C2) | (C2) | (C3) |
| $\mathrm{A} \rightarrow \mathrm{G}$ | $\mathrm{C} \rightarrow \mathrm{G}$ | $\mathrm{G} \rightarrow \mathrm{A}$ | $\mathrm{T} \rightarrow \mathrm{G}$ | $\mathrm{G} \rightarrow \mathrm{A}$ |

Allele frequency

| Total |  | 368 (92.0\%) | C | 128 (32.0\%) | G | 278 (69.5\%) | T | 128 (32.0\%) | G | 138 (34.5\%) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | A | 32 (8.0\%) | G | 272 (68.0\%) | A | 122 (30.5\%) | G | 272 (68.0\%) |  | 262 (65.5\%) |
| SLE | G | 192 (96.0\%) | C | 62 (31.0\%) | G | 136 (68.0\%) | T | 62 (31.0\%) | G | 70 (35.0\%) |
|  | A | 8 (4.0\%) | G | 138 (69.0\%) | A | 64 (32.0\%) | G | 138 (69.0\%) | A | 130 (65.0\%) |
| HD | G | 176 (88.0\%) | C | 66 (33.0\%) | G | 142 (71.0\%) | T | 66 (33.0\%) | G | 68 (34.0\%) |
|  | A | 24 (12.0\%) | G | 134 (67.0\%) | A | 58 (29.0\%) | G | 134 (67.0\%) |  | 132 (66.0\%) |

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| G | $37 \%$ | C | $35 \%$ | G | $81 \%$ | T | $29 \%$ | G | $35 \%$ |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| A | $63 \%$ | G | $65 \%$ | A | $19 \%$ | G | $71 \%$ | A | $65 \%$ |

B


Figure 1 SNPs in MHC2TA plll and coding region. (A) Genomic DNA was isolated from peripheral blood mononuclear cells, and PCR and direct sequencing were performed with eight oligonucleotide primer sets for genomic analysis of SNPs (arrows). (B) MHC2TA promoter elements and exon organisation. Arrows on the box represent upstream regulatory sequences for each promoter. Promoter pl is used primarily by dendritic cells, plll by B cells, and pIV for IFN $\gamma$-inducible CIITA expression by non-professional APCs.

