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Correction: Therapeutic potential of isochlorogenic acid A from *Taraxacum officinale* in improving immune response and enhancing the efficacy of PD-1/PD-L1 blockade in triple-negative breast cancer

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A Correction on

Therapeutic potential of isochlorogenic acid A from *Taraxacum officinale* in improving immune response and enhancing the efficacy of PD-1/PD-L1 blockade in triple-negative breast cancer

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There was a mistake in **Figure 7F** as published. The error was caused by the use of placeholder images during the typesetting process to maintain layout consistency; due to oversight, some placeholders were not replaced with the correct images. The corrected **Figure 7F** appears below.

The original version of this article has been updated.

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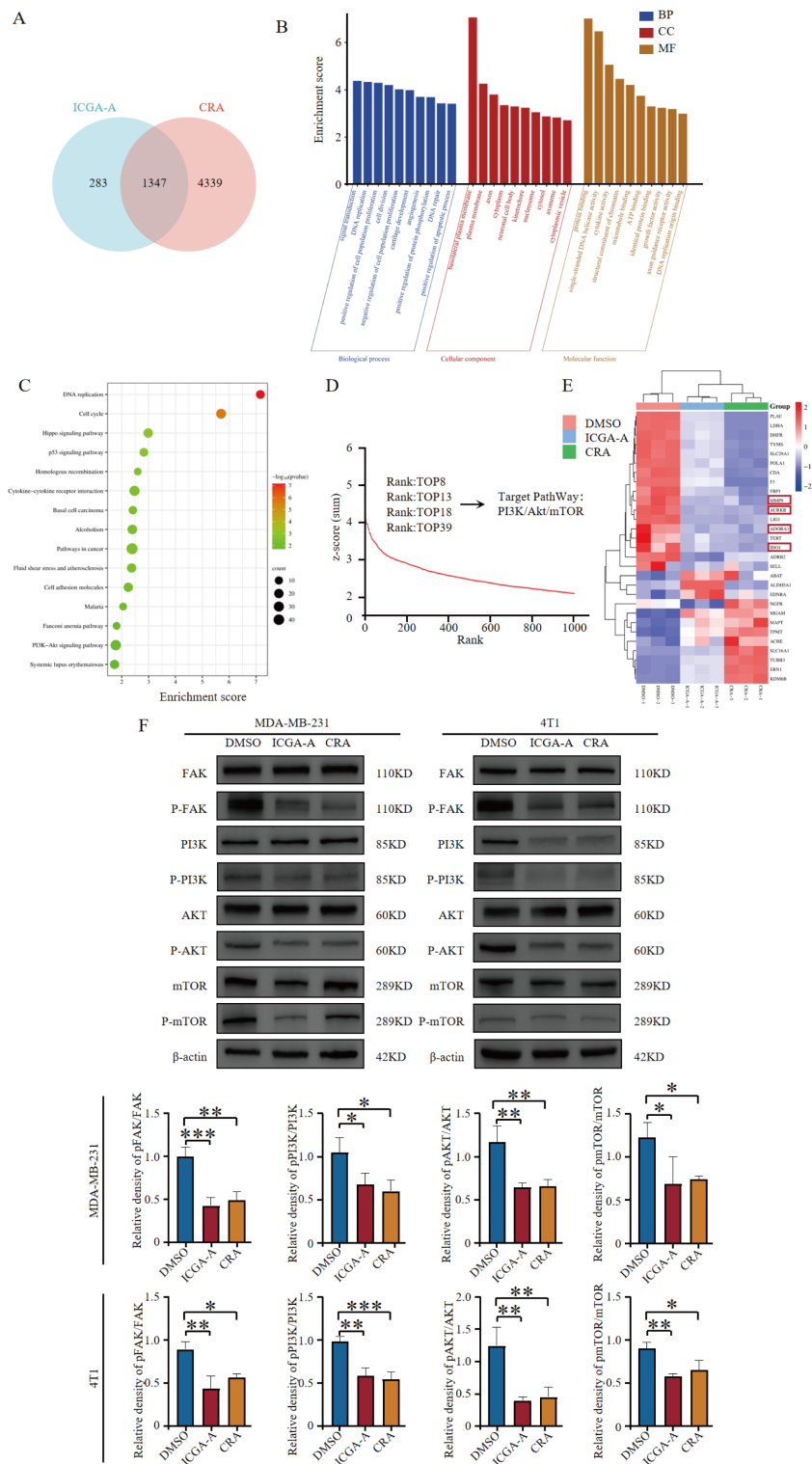


FIGURE 7
ICGA-A and CRA inhibited mRNA and protein expression levels of metabolic-associated proteins and pathways. **(A)** Venn diagram of DEGs in MDA-MB-231 cells treated with ICGA-A and CRA. **(B)** The top 10 GO terms in the BP, CC, and MF classifications of overlapping genes with MDA-MB-231. The x-axis represents the enriched terms, and the y-axis represents the enrichment score. **(C)** Top 20 KEGG pathways. KEGG pathway enrichment for the overlapping genes with MDA-MB-231. The x-axis represents the gene ratio ($p < 0.05$), and the y-axis represents the enriched terms. **(D)** The ranking of ICGA-A scored by HTS² in a library of over 20,000 compounds. **(E)** Heatmap of the intersecting genes between RNA-seq and network pharmacology targets. **(F)** ICGA-A and CRA decreased the phosphorylation levels of FAK, PI3K, AKT, and mTOR. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$ vs. DMSO.