

Correction to “Discovery of a Novel Class of Dual GPBAR1 Agonists—ROR γ t Inverse Agonists for the Treatment of IL-17-Mediated Disorders”

Bianca Fiorillo, Rosalinda Roselli, Claudia Finamore, Michele Biagioli,* Cristina di Giorgio, Martina Bordoni, Paolo Conflitti, Silvia Marchianò, Rachele Bellini, Pasquale Rapacciuolo, Chiara Cassiano, Vittorio Limongelli, Valentina Sepe,* Bruno Catalanotti, Stefano Fiorucci, and Angela Zampella

ACS Omega 2023, 8 (6), 5983–5994. DOI: [10.1021/acsomega.2c07907](https://doi.org/10.1021/acsomega.2c07907)

Cite This: ACS Omega 2023, 8, 45163–45163

Read Online

ACCESS |

Metrics & More

Article Recommendations

Amended Figure 5 and new caption.

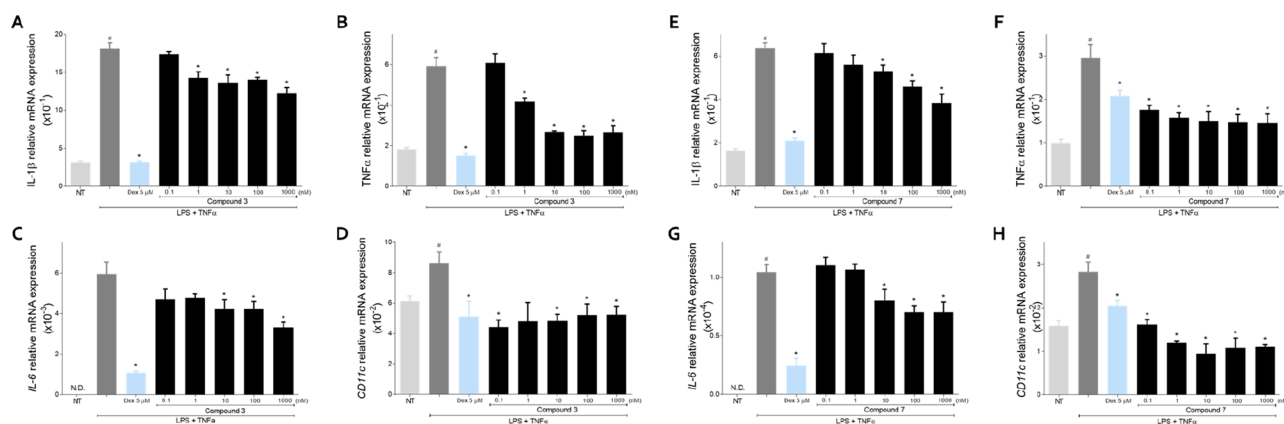


Figure 5. U937 cells activated with LPS (100 ng/mL) plus TNF α (100 ng/mL) for 24 h, alone or in combination with 3 or 7 at 0.1, 1, 10, 100, and 1000 nM. Dexamethasone (Dex) was used at 5 μ M. Quantitative real-time PCR analysis of expression of proinflammatory genes IL-1 β (A and E), TNF α (B and F), IL-6 (C and G) and M1 marker CD11c (D and H). These data are normalized to TBP mRNA expression. Data are derived from five replicates. Results represent mean \pm SEM # $p < 0.05$ vs NT group and * $p < 0.05$ vs LPS plus TNF α group. N.D. means not determined. Analysis of variance (ANOVA) was used for statistical comparisons.

Published: November 16, 2023

