Validation data for Jurkat-Lucia™ NFAT-CD16-Low Cells

https://www.invivogen.com/jurkat-lucia-nfat-adcc-adcp-cells

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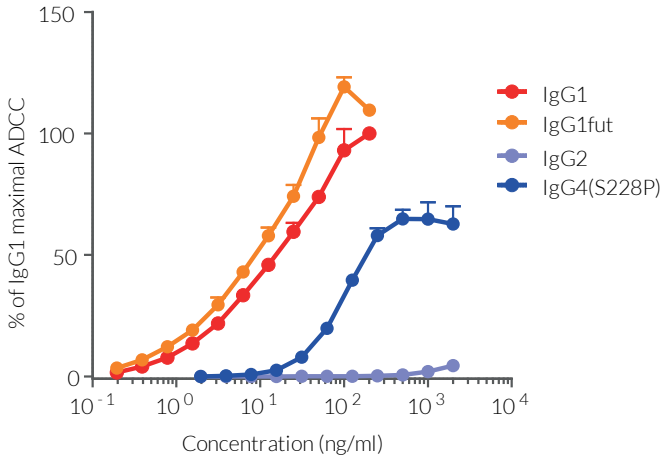
Version 22C10-NJ

Jurkat-Lucia™ NFAT-CD16-Low cells were engineered from the human T-lymphocyte Jurkat cell line and designed as reporter cells for antibody-dependent cellular cytotoxicity (ADCC). Nuclear translocation of NFAT (nuclear factor of activated T-cells), a transcription factor naturally expressed by Jurkat cells, is an early signaling event in ADCC induction. Jurkat-Lucia™ NFAT-CD16-Low cells stably express the cell surface Fc receptor CD16A (FcγRIIIA; F158 low-affinity allotype) and the Lucia luciferase reporter gene under the control of an ISG54 minimal promoter fused to six NFAT response elements. Human CD16A expression by Jurkat-Lucia™ NFAT-CD16 cells has been verified by flow-cytometry. These cells have been functionally tested with the Raji-Null target cell line (expressing CD20) and Anti-hCD20 IgG isotypes. Antibodies displaying lower EC_{50} have higher ADCC potency.

Jurkat-Lucia™ NFAT-CD16-Low cell responses to ADCC induction with anti-human CD20 isotypes and Raji-hCD20 target cells



Comparison of ADCC potency for native and engineered anti-human CD20 isotypes: Raji-Null cells were incubated with gradient concentrations of Anti-hCD20 or Anti-β-galactosidase (β-gal) mAbs for 1 hour. Jurkat-Lucia™ NFAT-CD16-Low effector cells were then co-incubated with targets cells for 6 hours. NFAT activation, reflecting the induced ADCC response, was assessed by determining Lucia luciferase activity in the supernatant using QUANTI-Luc™. Percentages of the maximal response normalized to the IgG1 isotype are shown.

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