Validation data for HEK-Blue[™] IL-1β cells

https://www.invivogen.com/hek-blue-il1b

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Version 21H18-MM

HEK-Blue^M IL-1 β cells allow the detection of bioactive human interleukin-1 β (hIL-1 β) by monitoring NF-kB and AP-1 activation. These human embryonic kidney HEK293 derived cells express an NF- κ B/AP-1-inducible SEAP (secreted embryonic alkaline phosphatase) reporter gene. HEK-Blue^M IL-1 β cells display high sensitivity to human IL-1 β (detection range: 0.01-100 ng/ml) and low sensitivity to murine IL-1 β (detection range: 10-1000 ng/ml) (Figure 1). They can detect both IL-1 α and IL-1 β , as these cytokines bind to the same receptor, IL-1R1. Of note, HEK-Blue^M IL-1 β cells show no response to human or murine TNF- α , flagellin, or Poly(I:C) (Figure 2). These cells can be used for screening antibodies targeting the IL-1 β /IL-1R1 pathway (Figure 3).

Cellular response to IL-1β

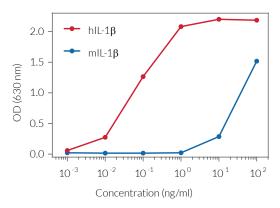


Figure 1. Dose-response of HEK-Blue^T IL-1 β cells to recombinant IL-1 β . Cells were stimulated with increasing concentrations of recombinant human (h) or murine (m) IL-1 β . After overnight incubation, the NF- κ B/AP-1 response was determined using QUANTI-Blue^T Solution, a SEAP detection reagent, and reading the optical density (OD) at 630 nm.

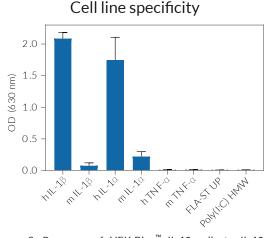


Figure 2. Response of HEK-Blue[™] IL-1 β cells to IL-1 β and IL-1 α . Cells were stimulated with 1 ng/ml of hIL-1 β , mIL-1 β , hIL-1 α , mIL-1 α , 100 ng/ml of hTNF- α , mTNF- α , ultrapure flagellin from *S. typhimurium* (FLA-ST UP) or 300 ng/ml of Poly(I:C) HMW. After overnight incubation, SEAP activity was assessed using QUANTI-Blue[™] Solution. OD at 630 nm is shown as mean ± SEM.

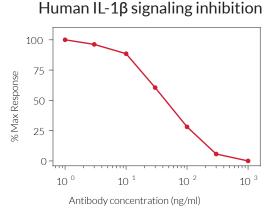


Figure 3. Dose-dependent inhibition of HEK-Blue^T IL-1 β cellular response using a neutralizing antibody against hlL-1 β . The anti-hlL1 β antibody was incubated with the cells for 30 minutes prior to the addition of hlL-1 β (1 ng/ml). After overnight incubation, SEAP activity in the cell culture supernatant was assessed using QUANTI-Blue^T Solution. Data represent % of maximal reporter activity without the anti-hlL1 β antibody.

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