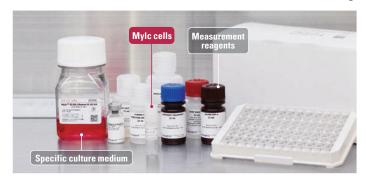


Mylc ELISA(human IL-6)kit

Condition setting has already been done. You can start evaluation studies immediately.



POINT You high

You can readily perform high quality evaluation.

You can readily use a large quantity of homogeneous Mylc cells for evaluation without troublesome condition setting.

POINT 2

Free from complicated condition setting.

Because the kit is delivered with the optimum conditions for evaluation, you can readily start evaluation.

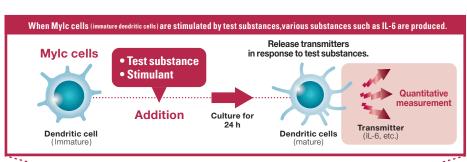


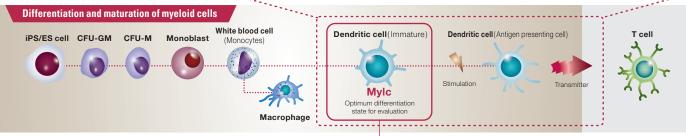
You can start immediately with this All-in-One kit.

This kit contains all the materials necessary for evaluation: from blood cells to measurement reagents. You can start evaluation immediately after opening the box.

Principle of evaluation

A test substance is added to the Mylc cells, and the produced transmitter (IL-6) is quantified. In 24 hours, the intensity of the stimulation by the test substance can be evaluated numerically, which enables judgment of the function.





Homogeneous blood cells with identical genetic information

The same genetic information and conditions can be always reproduced because they are produced in our laboratory. This feature contributes to a drastic improvement in the accuracy and applicability of pharmaceutical evaluation.

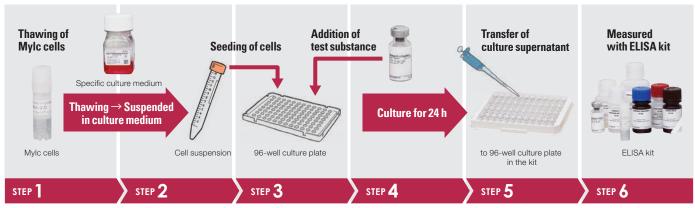
We can control the differentiation stage to the optimum conditions

Differentiation stage (maturity) can be minutely controlled with our high level of regenerative medicine technologies and product control. Research and evaluation of high accuracy are available.

Stable mass-production and supply are possible

You can obtain blood cells in the sufficient quantities in ideal condition any time. You can conduct research and development and efficacy evaluation with no delay and accelerate your development.

Assay procedure



Thawing of Mvlc cells

Frozen cells in a vial are thawed in a water bath.

Preparation of Mvlc cells

After washing by centrifugation, the cells are suspended in a specific culture medium.

Seeding of cells

Cells are seeded at a certain density in 96-well culture plates you provide.

Addition of test substance

Add test substance with the stimulant. and culture for 24 h.

Collection of culture supernatant in culture medium

Transfer culture supernatant from each well to the 96-well culture plate in the kit.

Measurement of IL-6

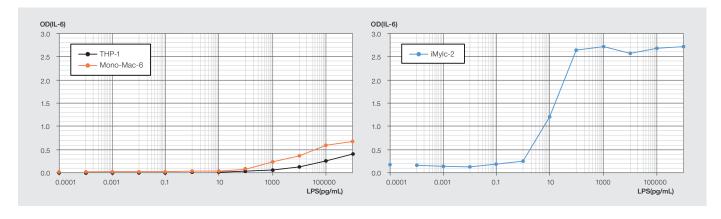
Quantified in a colorimetric ELISA assay according to the method attached to the kit.

In evaluation based on alternative methods to animal testing, Mylc cells respond with high sensitivity and stability.

Mylc, which shows similar response to in vivo cells, is superior in sensitivity in sensitization and stability compared with model cells (Mono-Mac-6, etc.) used in alternative methods to animal testing. In a comparison study in IL-6 production assay, Mylc-02 (right figure), used in this study, shows higher response at lower concentrations than Mono-Mac-6 (left figure).

Comparison study with the Mono-Mac-6

Comparison study in IL-6 production assay, in which a pyrogen (LPS) was added (expressed in OD) In Mylc-2 (right figure), used in this study, the production was detectable at lower concentrations than in Mono-Mac-6 (left figure), indicating a higher response



Contents of kit*

Cells: Mylc cells (frozen in storage medium, 1 screw vial)

Specific culture medium: Specific for culture of Mylc cells for measurement and evaluation (frozen product, 50 mL bottle × 1)

Stimulant: NiSO₄ solution for control stimulation (refrigerated, 1 vial)

Measurement reagent: ELISA kit for measurement of IL-6 produced in supernatant

* Please provide the following instruments and materials by yourself in advance. Sterilized centrifuge tubes / Pipettes and pipette tips / 96-well culture plate Centrifuge, CO2 incubator / other necessary materials and instruments

Evaluation and research in a common laboratory with Kyoto University

Our headquarters and research laboratory are located in Kyoto-University Katsura Venture Plaza, where Kyoto University nurtures the creation of new businesses utilizing new ideas/technologies and intellectual properties



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