

#### Monocyte activation test

Commonly used acronym: MAT

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# **Contact person**

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## **Organisation**

Name of the organisation Sciensano
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#### SCOPE OF THE METHOD

The Method relates to	Human health
The Method is situated in	Regulatory use - Routine production
Type of method	In vitro - Ex vivo
Specify the type of cells/tissues/organs	Whole human blood cells

#### **DESCRIPTION**

## **Method keywords**

Pyrogen test alternative to rabbit pyrogen test european pharmacopoeia endotoxins and non-endotoxin pyrogens

#### Scientific area keywords

quality control of injectable medicines

#### **Method description**

The monocyte-activation test (MAT) is used to detect or quantify substances that activate human monocytes or monocytic cells to release endogenous mediators such as proinflammatory cytokines, for example tumour necrosis factor alpha (TNF?), interleukin-1 beta (IL-1?) and interleukin-6 (IL-6). These cytokines have a role in fever pathogenesis. Consequently, the MAT will detect the presence of pyrogens (endotoxins on non-endotoxin pyrogens) in the test sample. The MAT is suitable, after a product-specific validation, as a replacement for the rabbit pyrogen test (see European pharmacopoeia

chapter 2.6.30). Although 3 methods (quantitative test, semi-quantitative test or reference lot comparison test) are described in the Ph. Eur.

## Lab equipment

Biosafety cabinet; CO2-incubator; ELISA plate reader.

#### Method status

Still in development History of use

# PROS, CONS & FUTURE POTENTIAL

# **Advantages**

No rabbits; ex-vivo test.

# Challenges

Proper storage of whole human blood (store only in nitrogen tank, not at minus 80°C).

# REFERENCES, ASSOCIATED DOCUMENTS AND OTHER INFORMATION

#### References

User Manual of the PyroDetect System Monocyte-Activation Test (MAT) from Merck; Monocyte Activation Test guideline from European Pharmacopoeia.

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