

# Direct Peptide Reactivity Assay

*Commonly used acronym: DPRA*

*Created on: 16-04-2019 - Last modified on: 22-02-2022*

## SCOPE OF THE METHOD

|                                  |   |
|----------------------------------|---|
| <b>The Method relates to</b>     | Human health  |
| <b>The Method is situated in</b> | Basic Research, Regulatory use - Routine production |
| <b>Type of method</b>            | In chemico  |
| <b>This method makes use of</b>  | Animal derived cells / tissues / organs             |

## DESCRIPTION

### Method keywords

toxicology

OECD

AOP

Molecular initiating event

### Scientific area keywords

Skin Sensitisation

in vitro

Toxicology

OECD

AOP

molecular initiating event

## **Method description**

The DPRA is an *in chemico* method which quantifies the remaining concentration of cysteine- or lysine-containing peptide following 24 hours incubation with the test chemical at 25 +/-2,5°C. The synthetic peptides contain phenylalanine to aid in the detection. Relative peptide concentration is measured by highperformance liquid chromatography (HPLC) with gradient elution and UV detection at 220 nm.

Cysteine and lysine peptide percent depletion values are then calculated and used in a prediction model (see paragraph 29) which allows assigning the test chemical to one of four reactivity classes used to support the discrimination between sensitisers and non-sensitisers.

## **Lab equipment**

HPLC UV

## **Method status**

History of use

Internally validated

Validated by an external party (e.g. OECD, EURL ECVAM,...)

## **PROS, CONS & FUTURE POTENTIAL**

### **Advantages**

Validated methodology (EURL ECVAM) ;  
AOP based ;  
High throughput ;  
Low cost ;  
*In chemico*.

## **Challenges**

The test method described in this Test Guideline is an *in chemico* method that does not encompass a metabolic system.

## **Future & Other applications**

The methodology behind AOP and MIE can be applied to other toxicological endpoints.

## **REFERENCES, ASSOCIATED DOCUMENTS AND OTHER INFORMATION**

### **References**

OECD, TG 442C, OECD GUIDELINE FOR THE TESTING OF CHEMICALS, *In Chemico* Skin Sensitisation: Direct Peptide Reactivity Assay (DPRA)

### **Associated documents**

## **PARTNERS AND COLLABORATIONS**

### **Organisation**

**Name of the organisation** Sciensano

**Department** Chemical and physical health risks

**Specific Research Group or Service** Medicines and health products

**Country** Belgium

## Geographical Area

Brussels

Region

*Coordinated by*



*Financed by*

