

LipidTOX assay in primary rat hepatocytes

Commonly used acronym: LipidTOX

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Organisation

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Department Pharmaceutical and Pharmacological Sciences

Specific Research Group or Service In Vitro Toxicology and Dermato-Cosmetology

Country Belgium

Geographical Area Brussels Region

SCOPE OF THE METHOD

| The Method relates to | Human health |
|---|---|
| The Method is situated in | Basic Research, Translational - Applied Research |
| Type of method | In vitro - Ex vivo |
| Species from which cells/tissues/organs are derived | Rat |
| Type of cells/tissues/organs | Primary rat hepatocytes |

DESCRIPTION

Method keywords

Steatosis

Phospholipidosis
Hepatocytes
neutral lipids
phospholipids
Hepatotoxicity

Scientific area keywords

Hepatotoxicity cytotoxicity Steatosis Phospholipidosis

Method description

The method detects two facets of drug-induced cytotoxicity i.e. the intracellular accumulation of phospholipids and of neutral lipids, i.e. phospholipidosis and steatosis respectively. The assay makes use of a kit containing an aqueous, red-fluorescent formulation of labelled phospholipids (LipidTOX™ Red phospholipid stain, excitation/emission ~595/615 nm) which is up taken by the cells upon incubation with a phospholipidosis-inducing compound. The second component of the kit is a selective green-fluorescent stain for neutral lipids (LipidTOX™ Green neutral lipid stain, excitation/emission ~495/505 nm), which can be used sequentially on fixed cells for the analysis of steatosis or can be used independently for single-parameter analysis (Nioi et al. 2007). Additionally, use of VECTASHIELD® Mounting Medium containing 4',6-diamidino-2-phenylindole (DAPI) which binds directly to DNA and produces upon excitation a blue fluorescence, enables intracellular localisation of the lipids.

Lab equipment

Fluorescence microscope

Method status

History of use

REFERENCES, ASSOCIATED DOCUMENTS AND OTHER INFORMATION

References

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