

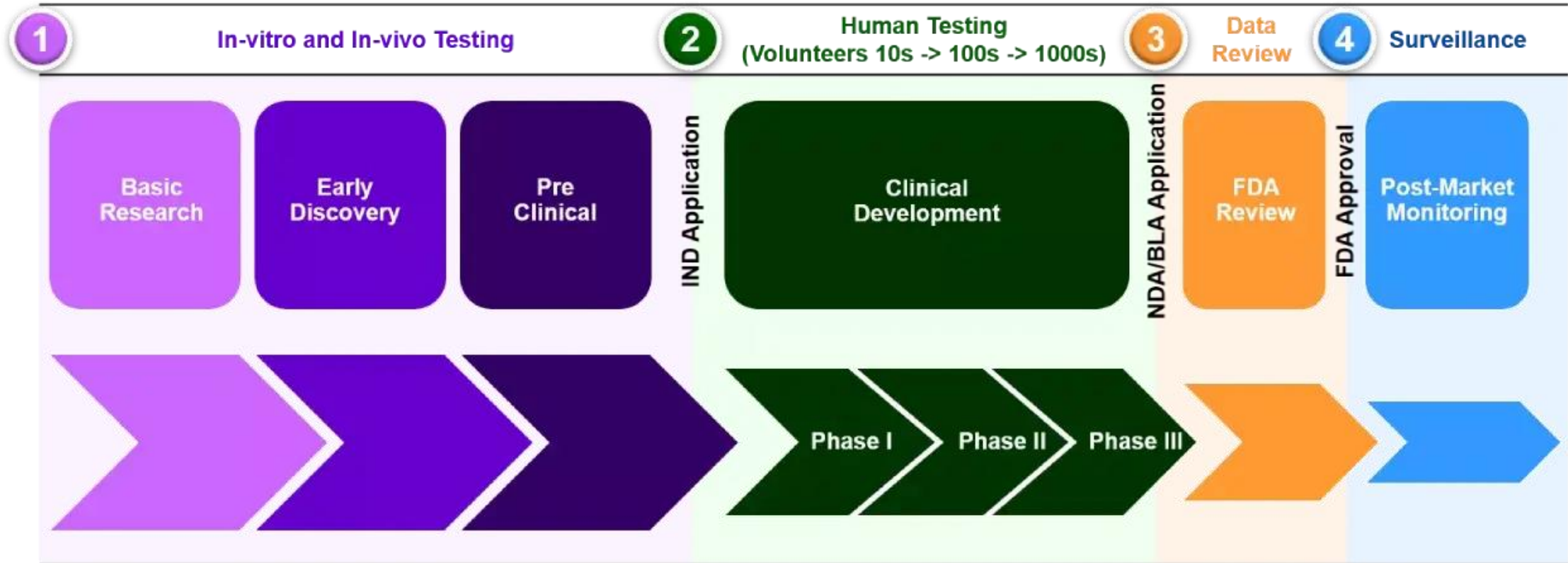
In vitro Toxicity Evaluation

Toxicology Seminar
Autumn 2024

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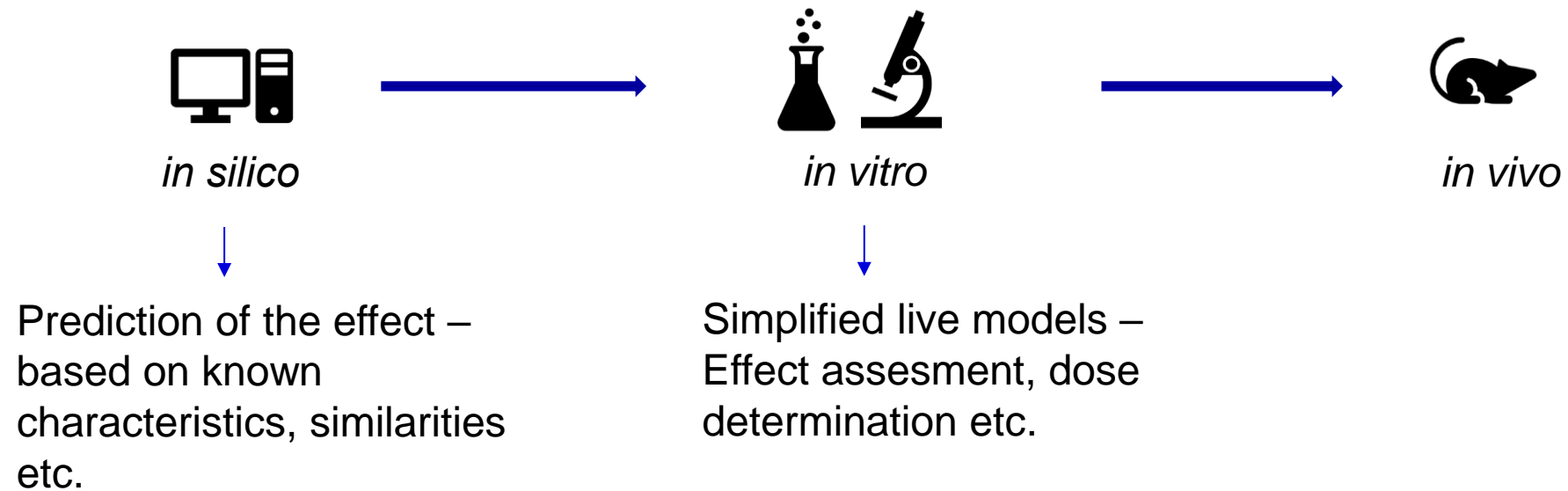
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Phases And Stages of Drug Development



Adapted from: PANDEY, Abhay. Phases of Drug Development Process, Drug Discovery Process. *NorthEast BioLab* [online]. 5. červen 2020 [vid. 2023-10-16]. Dostupné z: <https://www.nebiolab.com/drug-discovery-and-development-process/>

Position of *in vitro* methods among the processes of toxicity testing



Example of testing strategy – e.g. skin sensitization test

a. *in silico* (structure characteristics, pKa, log P etc.)

b. *in vitro* – human skin models

c. *in vivo* – animal model – usually albino rabbit

In vitro models

= living system simplified as compared with the *in vivo* model

Types of *in vitro* models

- Subcellular models (e.g. isolated mitochondria)
 - **Cell cultures**
 - Tissue cultures
 - Isolated organs
-
- 2D vs. 3D models



Cell line

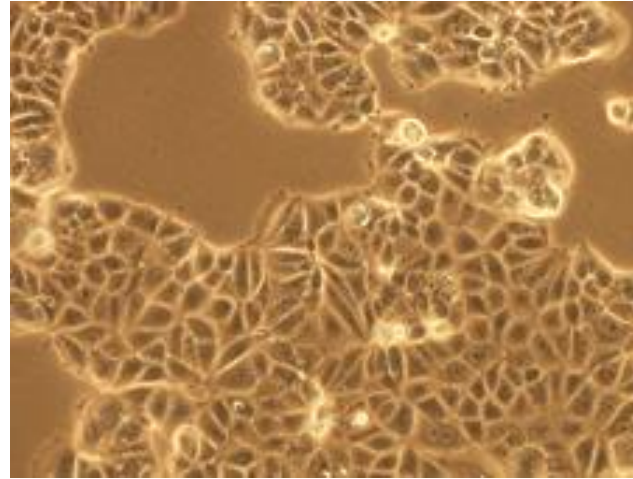
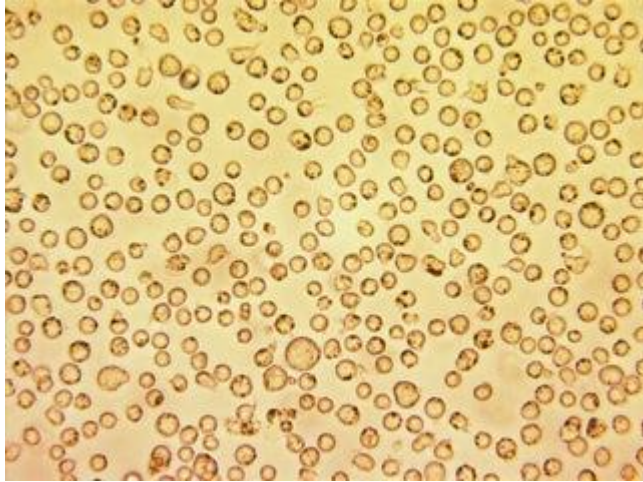
- Permanent cell lines
(immortalized – they will proliferate “indefinitely”)
- Single cell type
- Fully adapted to *in vitro* conditions
- They are derived from **tumor cells** or transformed from normal cells by physical or chemical mutagens

- Sources cell lines:
 - ATCC (American Tissue and Cultures Collection)
 - ECACC (European Collection of Cell Cultures)

In suspension

x

Adherent



Culture conditions



- !! Sterile conditions !! Risk of contamination
- Laboratory equipment
 - incubator, flowbox etc.



Culture conditions

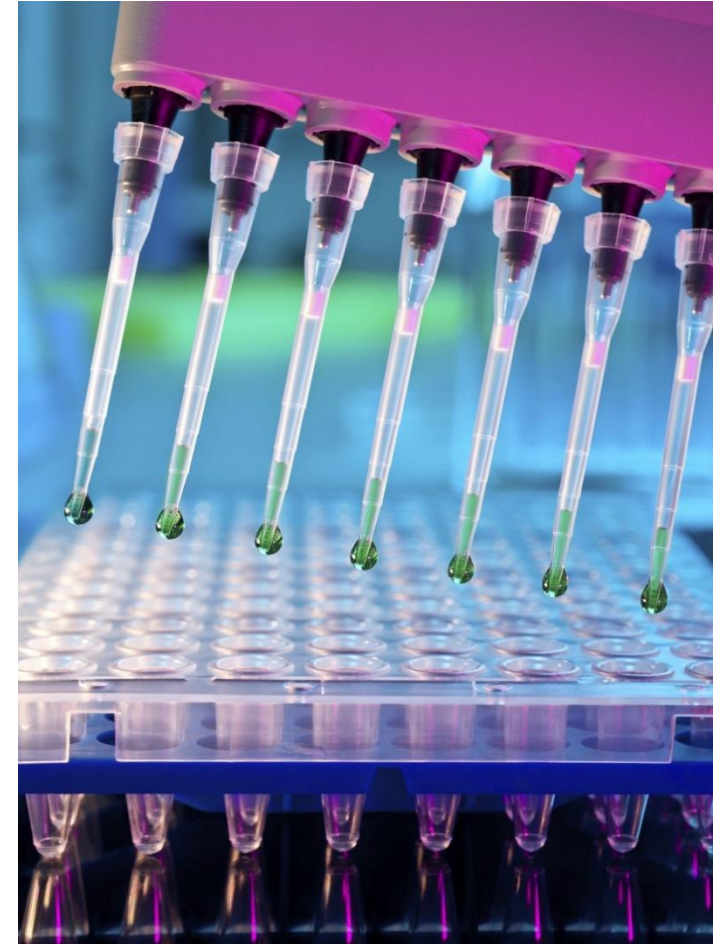


In vivo x *in vitro* models

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<p><i>In vivo</i></p> 	<p>Possible toxicokinetic testing</p> <p>Monitoring of the effect of systemic regulation</p> <p>In vivo models can not be fully eliminated !!</p>	<p>Financial aspects, time-consuming</p> <p>Ethical aspects</p> <p>Interindividual differences</p>
<p><i>In vitro</i></p> 	<p>Testing a larger number of compounds in a short time-period</p> <p>Plenty of biological material as a model</p> <p>Reproducibility</p> <p>Possibility of using human cell cultures</p> <p>Determination of organ-specific toxicity (eg. hepatotoxicity, nephrotoxicity etc.)</p>	<p>No information about systemic regulation</p> <p>For the replacement of in vivo model validation techniques are required</p> <p>The problem with the extrapolation of data</p> <p>Not all cell types can be cultured <i>in vitro</i></p> <p>Culturing under non-physiological conditions (culture media, cell lines in the absence of tissue context)</p>

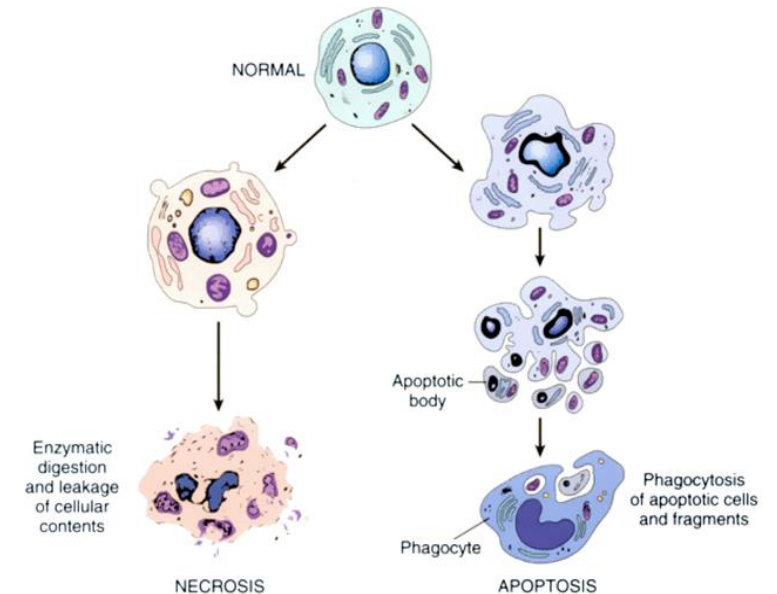
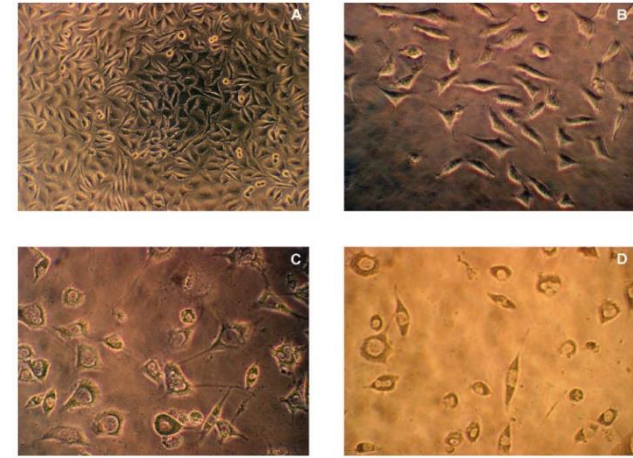
In vitro cell culture model for the evaluation of toxicity

- **One cell type** – homogenous properties e.g. expression of specific receptors, overexpression of cell cycle regulators etc.
- Enable us to study **the molecular basis of the toxic effect** – how the potential toxic substances affect their biological targets
- The results correspond to the effect of the substances without any interactions with other cell types or tissues



Cytotoxicity

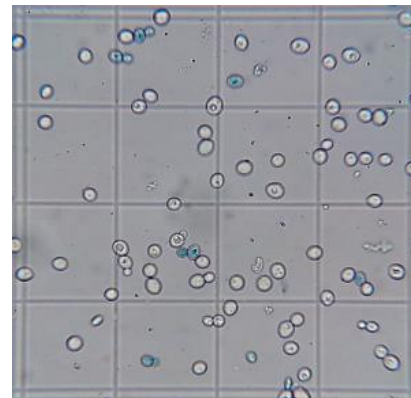
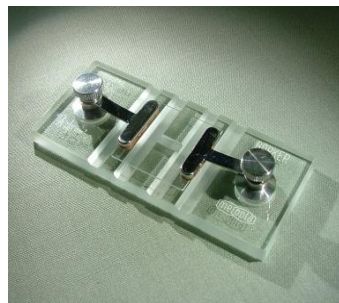
- It means the response of cells to the effect of toxic substance
- Cytotoxic effect:
 - Changes of cell morphology (augmentation, multinuclear cells, granularity of cell surface etc.)
 - Changes in cell metabolism
 - Inhibition of proliferation (changes in cell cycle progression etc.)
 - Cell death (apoptosis, necrosis, autophagy etc.)



Dye-exclusion test

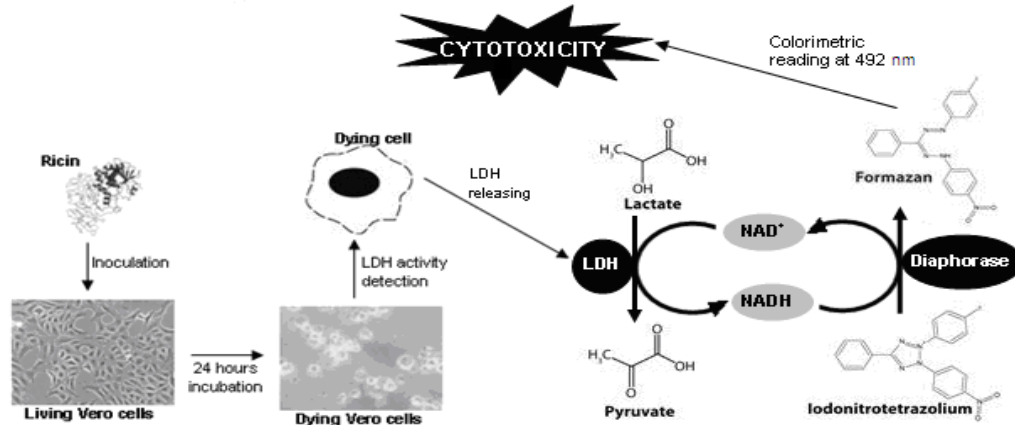


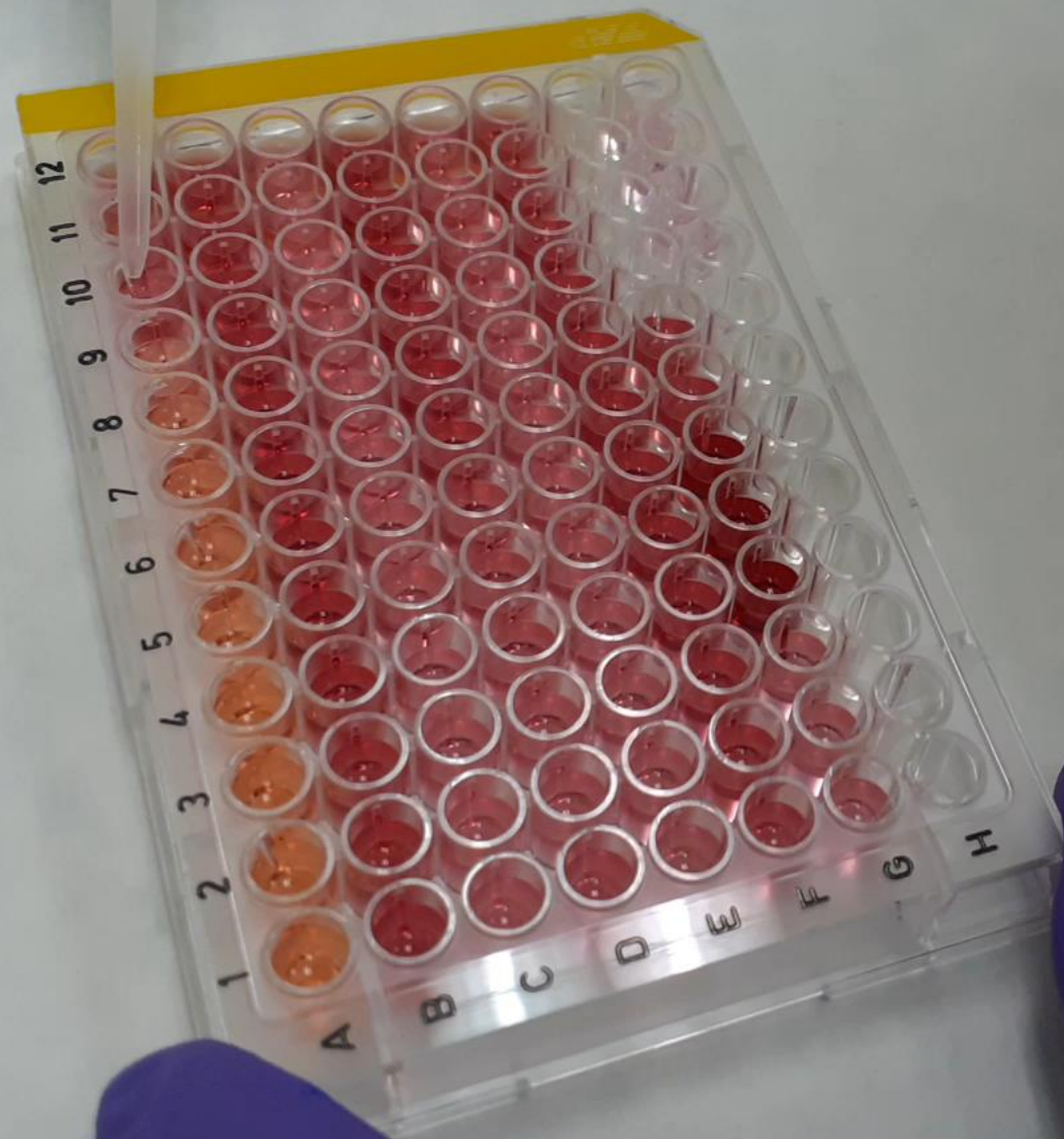
- Evaluating of the cell viability using intracellular cell dyes e.g. erythrosin B, trypan blue, neutral red
- Viable (live) cells stay unstained x dead cells are stained
→ because of the penetration of the dye through impaired cell membrane
- Hemocytometer (e.g. Bürker chamber) + microscope



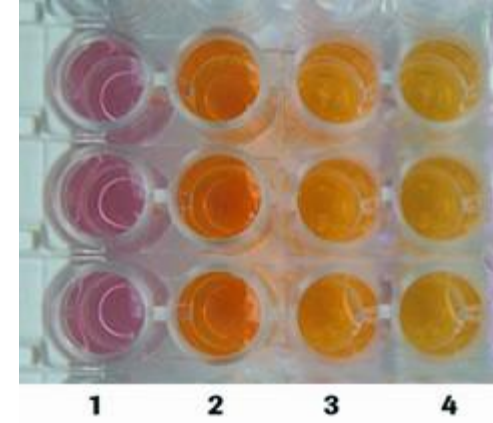
Cell viability

- Another method:
 - **Lactate dehydrogenase (LDH) analysis** – impaired membrane integrity, then we can detect LDH extracellularly





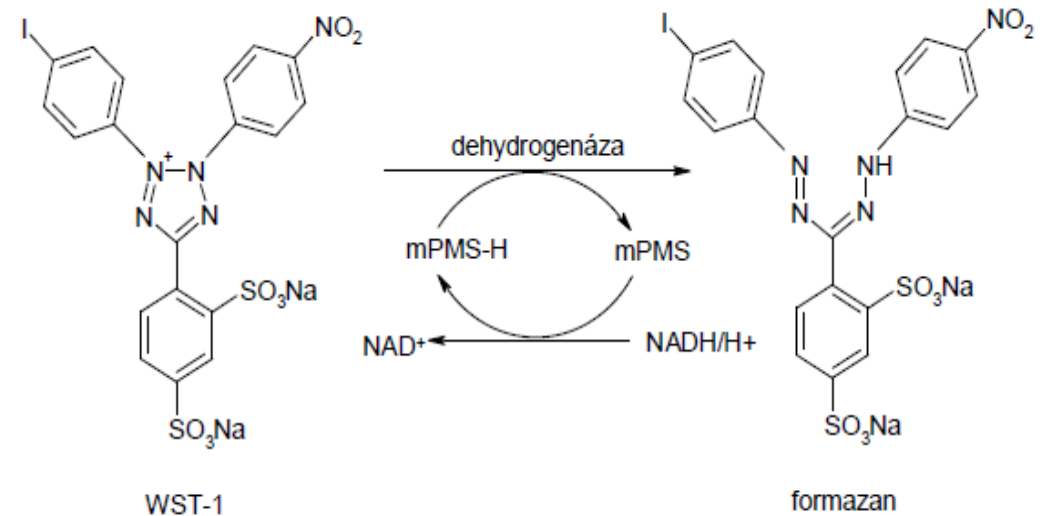
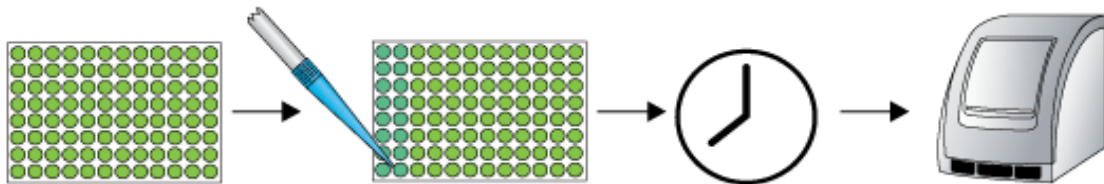
Cell viability and proliferation



– Tetrazolium salts

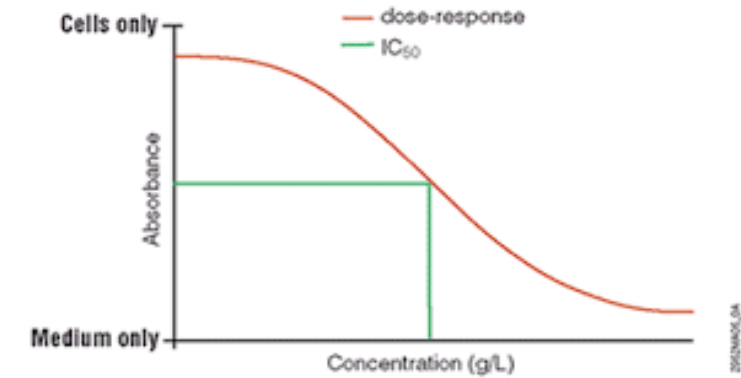
- Enzymatic reduction of TS (changes of the color) – changes of the color correlate with the intensity of cell metabolic activity

– MTT; XTT; WST-1 analysis

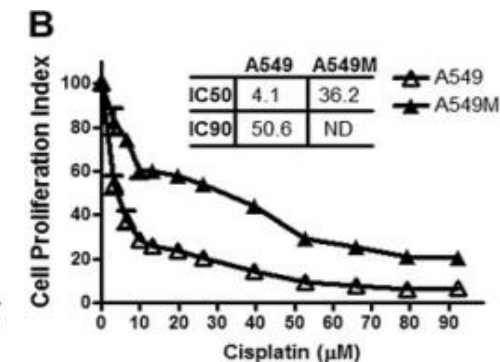
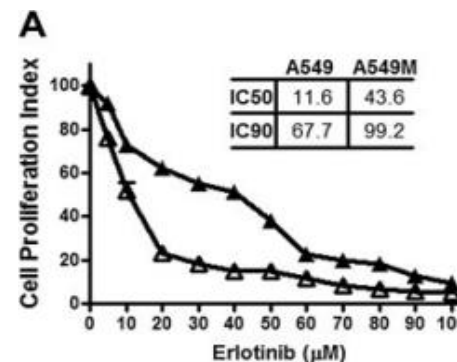


Resulting parameters

Inhibitory concentration	Lethal concentration	Effective concentration	Toxic concentration
IC ₁₀	LC ₁₀	EC ₁₀	TC ₁₀
IC ₅₀	LC ₅₀	EC ₅₀	TC ₅₀
IC ₉₀	LC ₉₀	EC ₉₀	TC ₉₀



- Enable us to compare the effect of different substances
- Depend on the time of exposition, selected model etc.
- *Dose-response curve*



In vitro toxicity tests – validated methods

Validated by **OECD**

(Organisation for Economic

Co-operation and

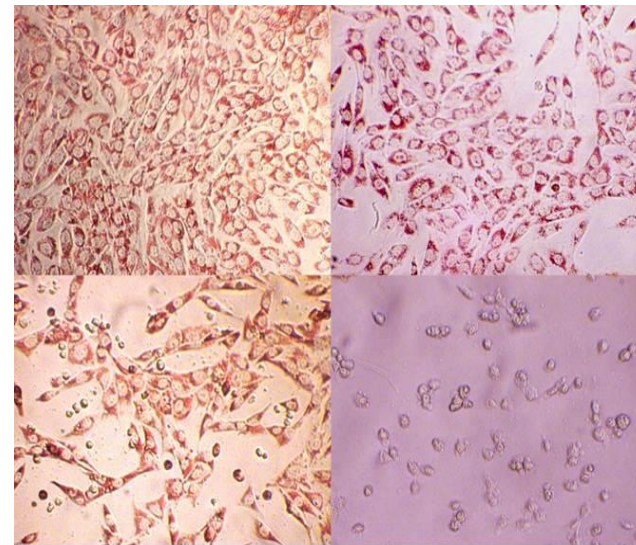
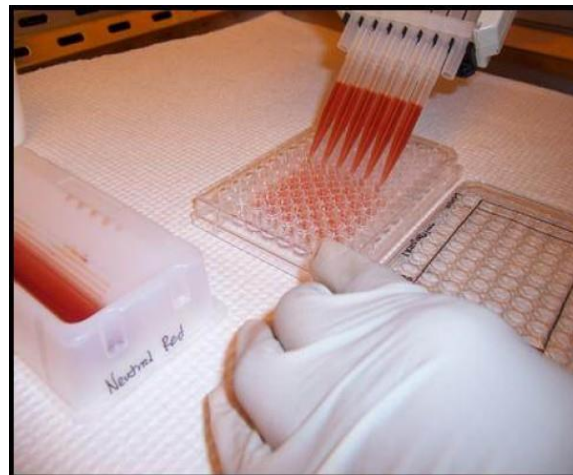
Development)

- Cytotoxicity
- Genotoxicity
- Eye irritation
- Phototoxicity

- Cardiotoxicity
- Nephrotoxicity
- Hepatotoxicity
- Reproductive toxicity
- Ecotoxicity

Cytotoxicity

- E.g. Biological evaluation of medical devices
 - = Tests for *in vitro* cytotoxicity
- Fibroblast cell line 3T3 - Neutral Red Uptake Test (NRU)
- Incorporation of NR into the lysosomes of live cells – spectrophotometrical determination of changes in color intensity



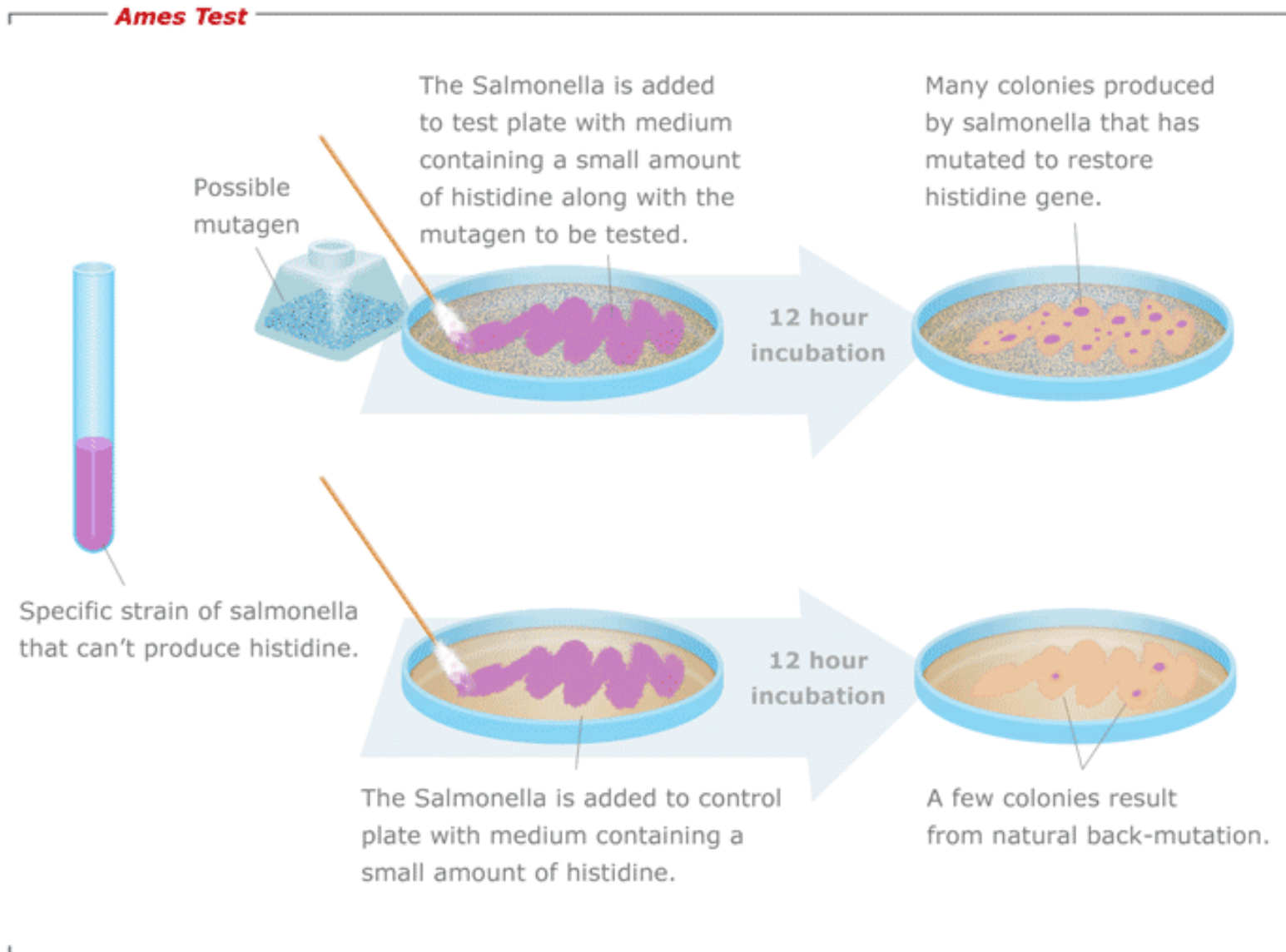
Genotoxicity

- Principle: detection of mutation – DNA damage (changes in genetic information)

Standard testing series:

- The bacterial reverse mutation test (**Ames Test**)
- *In Vitro* Mammalian Chromosome Aberration Test (changes in cell ploidy, detection of polyploidy)
- *In vitro* mammalian cell gene mutation test
- Etc.

Ames test



Eye Irritation/Corrosion

- Methods for testing the effect of potential ocular corrosive or irritant substances

Examples of *in vitro* test methods:

- **BCOP** (Bovine Corneal Opacity and Permeability)
- **ICE** (isolated chicken eye) test
- **HET-CAM** (Hen's Egg Test - Chorioallantoic Membrane)
 - Fertilized eggs – substances are applied on the membrane – changes are detected
 - e.g. hemorrhage, koagulation, lysis ...



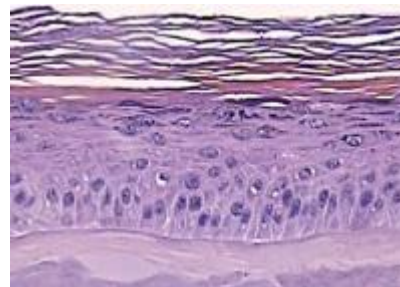
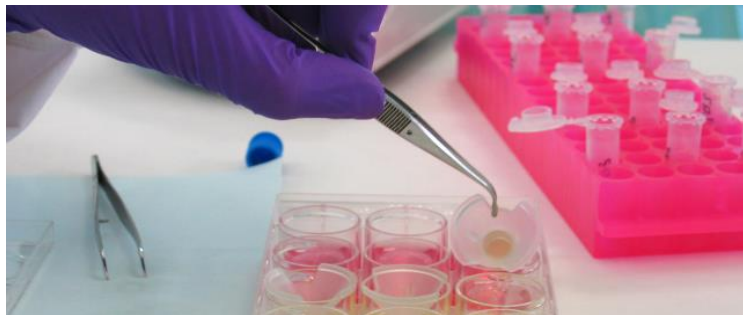
In vitro

Skin Corrosion Test: human skin model

- Episkin, Epiderm, SkinEthic....
- (reconstructed human epidermis – human keratinocytes – with functional stratum corneum)

Principle:

- Application of tested substance on the surface of human skin model - MTT analysis – detection of changes in cell viability



Reproductive toxicity

Embryonic Stem Cell test (EST test)

= evaluation of embryotoxic potential of tested substances

Principle of the test: determination of inhibited differentiation of embryonic stem cells (ESC)
and inhibition of cell proliferation (ESC and 3T3 fibroblasts)

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Thank you for your attention