## In Vitro Ocular Irritation Assay

## Need

Assessing the risk of eye injury is a primary consideration in determining the safety of ophthalmologic drugs or cosmetics. It is also of concern in the case of accidental exposures to chemicals, formulations, and other products. Early ocular testing is important because:

- Early assessment can determine a product's relative risk for causing eye irritation and/or serious eye damage.
- Initial testing may identify chemicals or mixtures with a low potential for eye injury and obliviate the need for further *in vitro* or *in vivo* testing.

## Solution

LifeNet Health offers chemical testing services with the validated EpiOcular™ Eye Irritation Test (EIT) for the assessment of potential ocular irritation and/or serious eye damage of client's test articles, operating in full compliance with the OECD 492 guideline. The health of the corneal tissue is assessed by measuring tissue viability immediately following exposure and a post-treatment incubation period.



Accurate & reliable data



Fast turnaround times



OECD methodbased studies



Collaborative

## **Testing Parameters**

| ASSAY PARAMETERS                | PROTOCOL  |
|---------------------------------|---|
| Model                           | MatTek EpiOcular™ (OCL-200)   |
| Replicates                      | 3   |
| <b>Test Article Formulation</b> | Liquid/Solid (tested neat or as provided)                                   |
| Negative Control                | Sterile deionized water   |
| Positive Control                | Methyl Acetate (applied neat)   |
| Exposure Time                   | 30 minutes (liquid test articles) and/or 6 hours (solid test articles)      |
| Post Exposure Soak              | 12 minutes (liquid test articles) and/or 25 minutes (solid test articles)   |
| Post Exposure Recovery          | 2 hours (liquid test articles) and/or 18 hours (solid test articles)        |
| Viability Assessment            | MTT   |
| Time to Complete                | 2-3 weeks from initiation   |
| Regulatory                      | Non-GLP or GLP  |
| Deliverables                    | Full Report including Tissue Viability, UN GHS classification (if possible) |



| TISSUE VIABILITY<br>(% NEGATIVE CONTROL) | UN GHS CATEGORIZATION                                       |
|--|---|
| <u> </u>                                 | No prediction can be made (UN GHS Category 2 or Category 1) |
| <b>&gt; 60%</b>                          | Non-Irritant (UN GHS No Category)                           |

Irritancy categorization based on tissue viability after exposures and post-exposure recovery.

