

# Alkaline In Vitro Comet Assay in C3a cells

*Commonly used acronym: in vitro comet*

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## SCOPE OF THE METHOD

<b>The Method relates to</b>	Human health
<b>The Method is situated in</b>	Basic Research
<b>Type of method</b>	In vitro - Ex vivo
<b>This method makes use of</b>	Human derived cells / tissues / organs
<b>Specify the type of cells/tissues/organs</b>	C3a Cells (Hepatocellular Carcinoma Cells, the C3A cell line is a clonal derivative of Hep G2 cells )

## DESCRIPTION

### Method keywords

DNA damage

DNA strand breaks

single cell gel electrophoresis assay

comet

automated scoring

### **Scientific area keywords**

in vitro toxicology

genotoxicity

hazard assessment

### **Method description**

The in vitro Alkaline Comet Assay is a microgel electrophoresis technique which allows to measure DNA damage (single and double strand breaks, alkali labile sites, incomplete excision repair sites and cross links) cell by cell. Cells are mixed with 0.8% Low Melting Point Agarose which is spread as a gel onto a microscope slide. The cells are lysed with high salt concentrations and detergents. The remaining nuclear DNA is then denaturated in alkali buffer pH>13 and electrophoresed in the same buffer. The negatively charged DNA fragments migrate out of the nucleus, towards the positive pole whereas undamaged supercoiled DNA does not migrate. After electrophoresis and neutralization of the slides, cells are dried and stained with GelRed, a fluorochrome intercalating agent. A fluorescent microscope equipped with an image analysis system is used to capture and quantify DNA damage in the single cells. DNA damage is expressed as percentage of DNA in the tail.

### **Lab equipment**

- Standard equipment for working with cell cultures;
- Electrophoresis Chamber with Power Supply Circulating Pump;
- Fluorescence Microscope;
- Software for automated imaging.

### **Method status**

History of use

Published in peer reviewed journal

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

### **Advantages**

- Identify DNA damage at the single cell level;
- Sensitivity for detecting low levels of DNA damage;
- Only small numbers of cells per sample needed;
- Fast, cheap & easy;
- Applicable on many cell types.

### **Challenges**

- Most of the damage is repaired (no fixed mutations are detected, regulatory genotoxicity testing in vitro usually relies on mutagenicity test not indicator tests ) but this gives an opportunity to study DNA repair;
- Quality of protocol and experimental performance is of crucial importance;
- Suitable statistical analysis;
- Incomplete metabolic capacities C3a cell line (Problems with indirect mutagens).

### **Modifications**

- Use of a cooling system for the electrophoresis device to minimize slide/slide differences;
- Improvement of the imaging system (upgrade software metasystems);
- Use of other cell lines (e.g. TK6 cells).

### **Future & Other applications**

- High Throughput analysis (48 well slides Trevigen);

- Use of lesion specific enzymes to make comet more sensitive (eg FPG) and more specific (eg epigenetic studies).

## REFERENCES, ASSOCIATED DOCUMENTS AND OTHER INFORMATION

### References

Collins AR, Azqueta Oscoz A, Brunborg G, Gaivão I, Giovannelli L, Kruszewski M, Smith CC, Stetina R (2008) The comet assay: topical issues. *Mutagenesis* 23:143–151.

Tice RR, Agurell E, Anderson D, Burlinson B, Hartmann A, Kobayashi H, Miyamae Y, Rojas E, Ryu JC, Sasaki YF (2000) Single cell gel/comet assay: guidelines for in vitro and in vivo genetic toxicology testing. *Environ Mol Mutagen* 35:206–221.

Azqueta A. and Collins A. R . (2013) The essential comet assay: a comprehensive guide to measuring DNA damage and repair. *Arch. Toxicol .*, 87, 949–968.

Investigation into the genotoxicity of water extracts from Hypoxis species and a commercially available Hypoxis preparation., Verschaeve, Luc, Mertens Birgit, Ndhlala A R., Anthonissen R, Gorissen B, and Van Staden J , *Phytother Res*, 2013 Mar, Volume 27, Issue 3, p.350-6, (2013).

### Associated documents

[The essential comet assay a comprehensive guide to measuring DNA damage and repair.pdf](#)

[Single Cell GelComet Assay Guidelines for In Vitro and In Vivo Genetic Toxicology Testing.PDF](#)

[the comet assa topical issues Mutagenesis-2008-Collins-143-51.pdf](#)

[Investigation into the genotoxicity of water extracts from Hypoxis species and a commercially available Hypoxis preparation.pdf](#)

### Other remarks

In addition to genotoxicity testing we also apply the in vitro comet assay to examine

the antigenotoxic properties of test substances (mostly plant extracts). A test solution is considered antigenotoxic when the genetic damage caused by the combined treatments (extracts and known mutagen) is substantially lower compared to the damage induced by the mutagen alone.

## PARTNERS AND COLLABORATIONS

### Organisation

**Name of the organisation** Sciensano

**Department** Chemical and physical health risks

**Country** Belgium

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