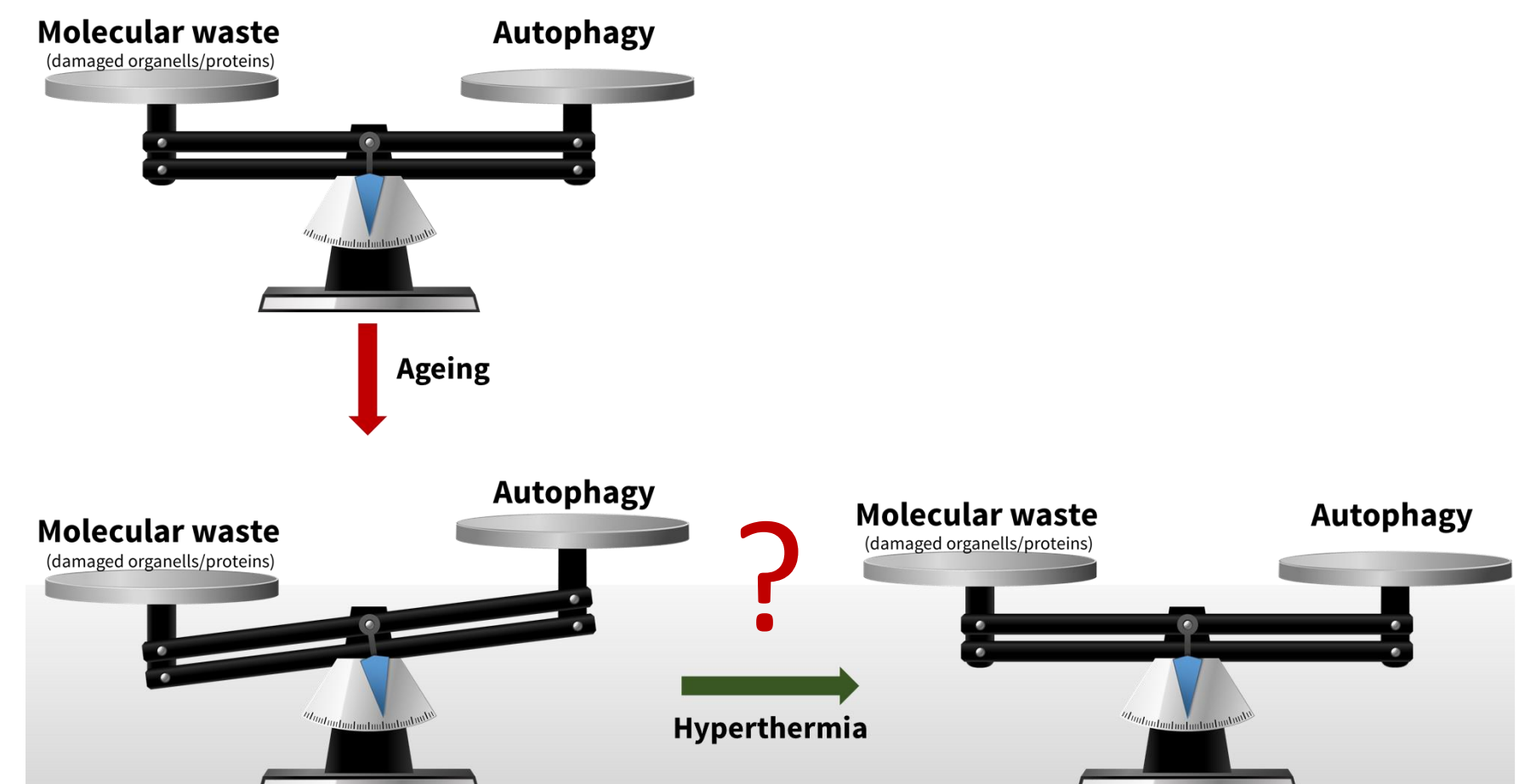


Hochecker Barbara¹, von Ardenne Alexander², Molinski Noah², Schniertshauer Daniel¹, Bergemann Jörg¹
Albstadt-Sigmaringen University, Germany¹; Von Ardenne Institute of Applied Medical Research GmbH, Germany²

1 Introduction

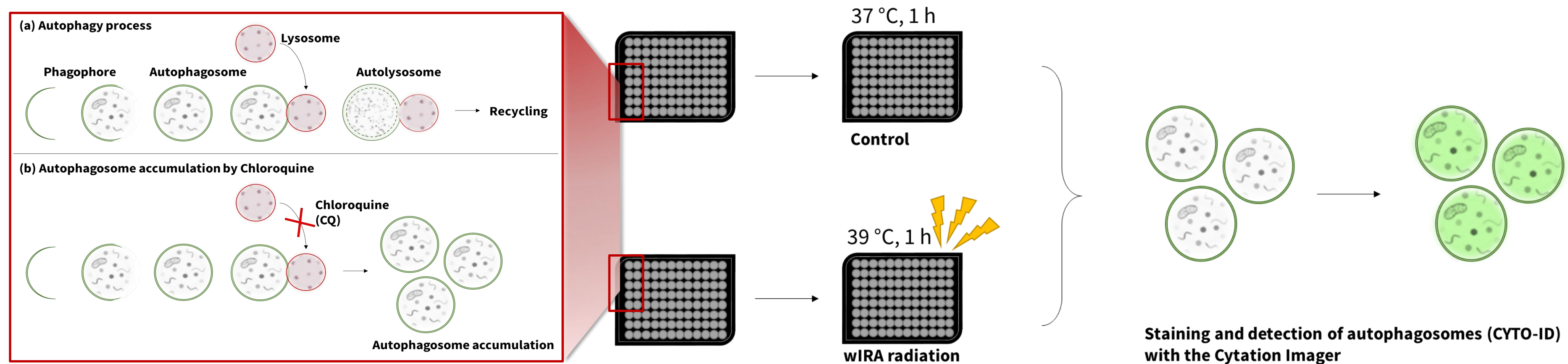
Autophagy is an intracellular degradative mechanism that removes damaged cell organelles and is critical for maintaining cellular homeostasis under stress conditions, such as age-related cellular dysfunctions [1]. It is well known that damaged proteins and organelles accumulate in aged cells due to these dysfunctions [2]. To counteract this large amount of molecular waste, an increase in the autophagic process is required. A promising option to activate autophagy is hyperthermia with water-filtered infrared A (wIRA) radiation. [3-4]

→ Does hyperthermia have a positive effect on autophagy in human skin fibroblasts?



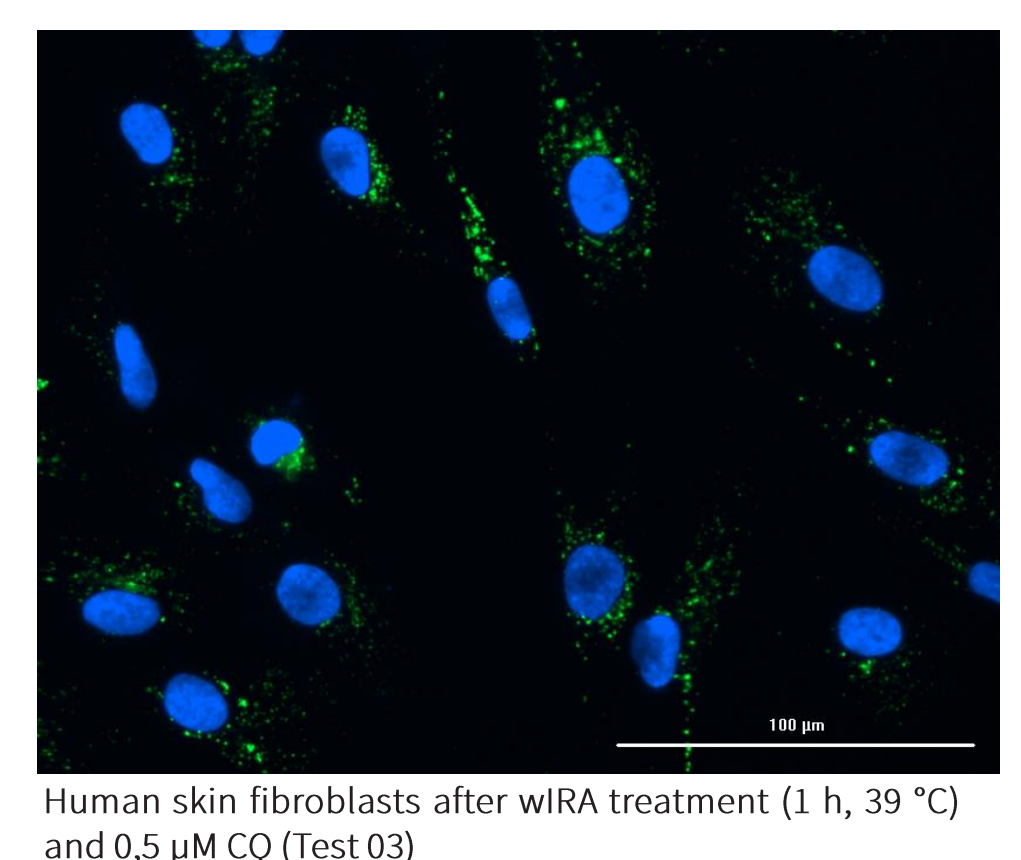
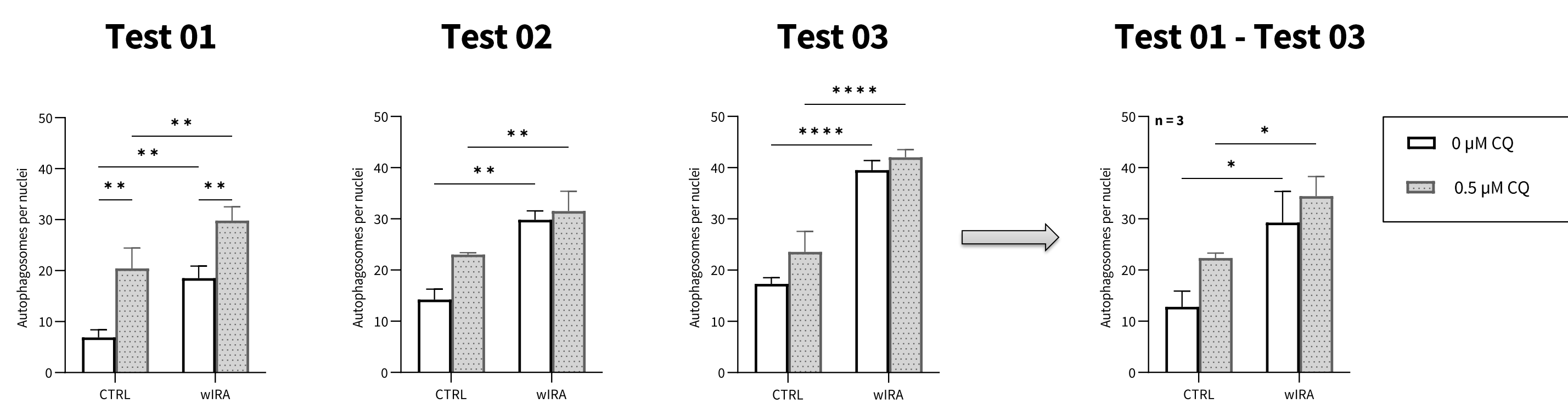
2 Method

We investigated the effect of one-hour hyperthermia treatment with wIRA radiation at 39 °C on autophagy in human skin fibroblasts. For this purpose, some cells of both groups (control/wIRA) were inhibited with chloroquine (CQ) prior to wIRA treatment. CQ acts as a test control by inhibiting the fusion of lysosomes with autophagosomes, resulting in the accumulation of autophagosomes. The cells were treated with wIRA at 39 °C for 1 h or incubated for 1 h at 37 °C (control). After treatment, plates were subjected to the CYTO-ID® Autophagy Detection Kit 2.0 and read with the Cytation Imager.



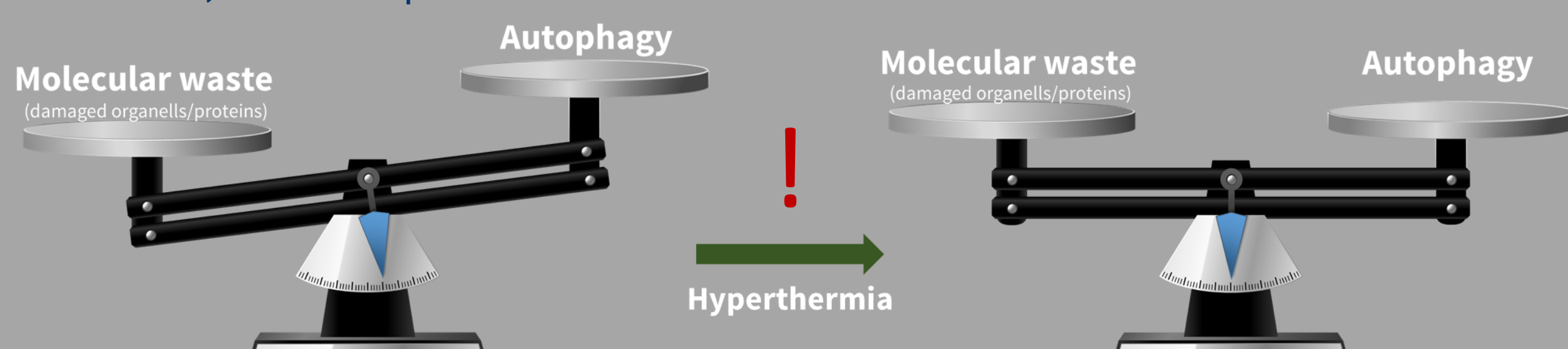
3 Results

Hyperthermia leads to a stronger induction of autophagosome formation per nuclei compared to the control group. All three performed experiments as well as the grouped data show significantly increased induction of autophagosomes per nuclei. Statistical analysis was conducted as two-way ANOVA with subsequent Tukey's multiple comparisons test. (ns $P \geq 0.5$, * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$ and **** $P < 0.0001$)



4 Conclusion

Our data suggest that hyperthermia has a positive effect on autophagy in human skin fibroblasts and may be a potential way to interfere with skin aging. However, further experiments are needed to confirm these results.



Quellenangaben

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