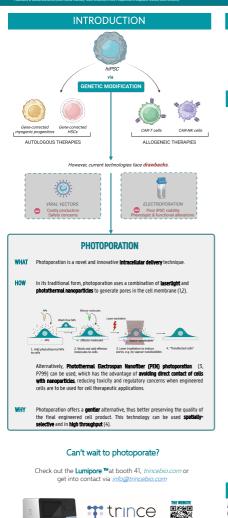
EFFICIENT AND GENTLE NON-VIRAL ENGINEERING OF iPSCs BY PHOTOPORATION

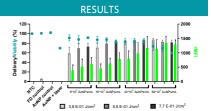


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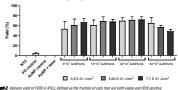


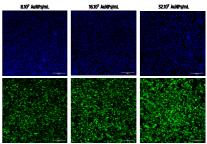
METHODS

- 60 nm cationic could nanonarticles (PDDAC) were used as photothermal sensitizers
- FITC-labelled dextran of 10kDa (FD10) was used a model caroo molecule (2 mg/mL).
- Delivery efficiency and relative mean fluorescence intensity were quantified using flow cytometry.
- Viability was measured by means of the metabolic activity assay CellTiter-Glo.



ft y-axis, grey bars) and ---avis, blue dots) was





Report 1 Confocal microscopy mages of iPSCs after AuNP-; with Hoechst 33342 (blue channel, top row). Increasing co with FDIO (green channel, of AuNPs: (from left to riot tw). Nuclei were wed Scale har 25

CONCLUSION and FUTURE PERSPECTIVES

AuNP-Photoporation can reach high delivery percentages while maintaining high viability in IPSCs Other cargo molecules and photosensitizers technology for the transfection of iPSCs. will be tested to further discover this promising

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