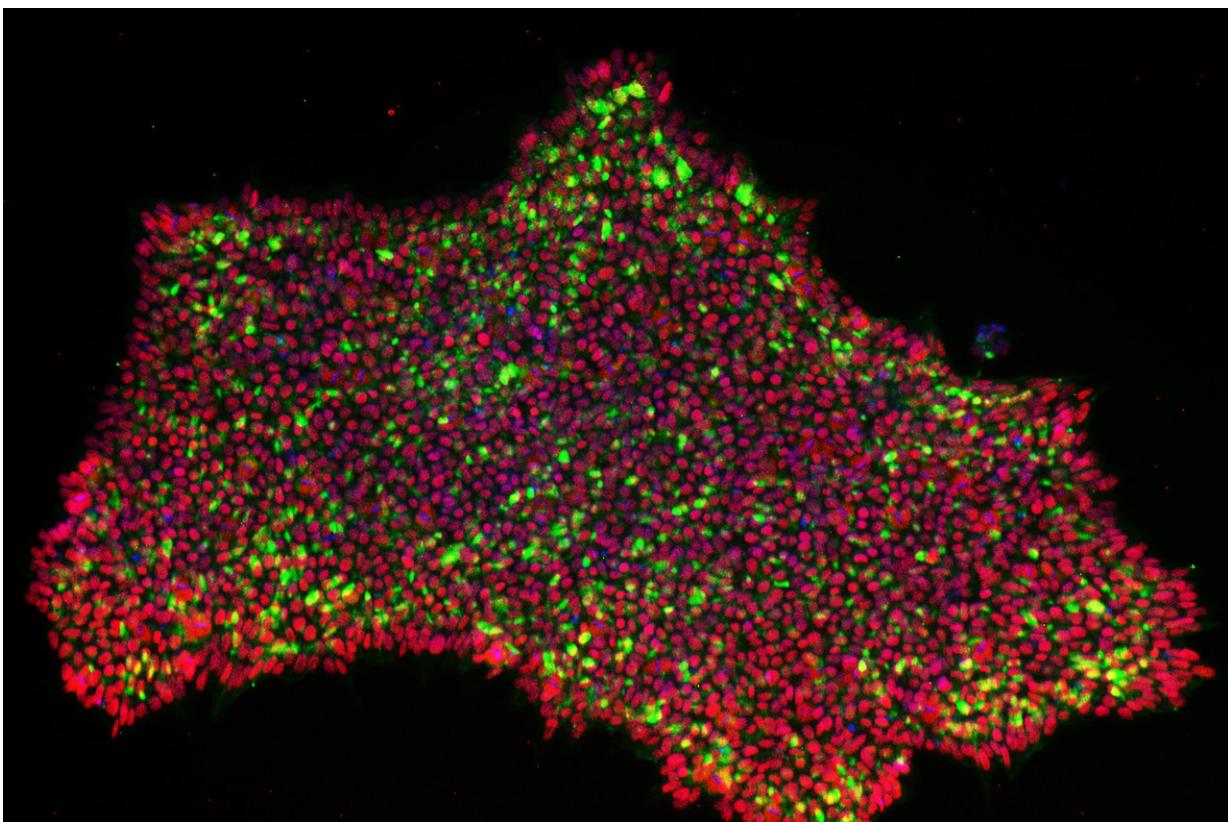


# Researchers develop a new method for turning skin cells into pluripotent stem cells

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CRISPRa reprogrammed induced pluripotent stem cell colonies stained for pluripotency marker expression. Credit: Otonkoski Lab / University of Helsinki

Our bodies consist of many different kinds of cells, each with their own role. The Japanese scientist Shinya Yamanaka had made earlier the

discovery, earning the Nobel Prize in 2012, that cells from adult skin can be converted to cells typical of early embryos, so-called induced pluripotent stem cells (iPSC). This process is called reprogramming.

Up until now, reprogramming has only been possible by introducing the critical genes for the conversion, called Yamanaka factors, artificially into skin cells where they are not normally active at all. Professor Timo Otonkoski at the University of Helsinki and Professor Juha Kere at Karolinska Institutet and King's College London have now succeeded in converting [skin cells](#) into pluripotent stem cells by activating the cell's own genes. This was achieved by using gene editing technology—called CRISPRa—that can be directed to activate genes. The method utilizes a blunted version of the Cas9 'gene scissors' that does not actually cut DNA, and can therefore be used to activate gene expression without mutating the genome.

"CRISPR/Cas9 can be used to activate genes. This is an attractive possibility for [cellular reprogramming](#) because multiple genes can be targeted at the same time. Reprogramming based on activation of endogenous genes rather than overexpression of transgenes is also theoretically a more physiological way of controlling cell fate and may result in more [normal cells](#). In this study, we show that it is possible to engineer a CRISPR activator system that allows robust reprogramming of iPSC," tells Professor Otonkoski.

An important key for the success was also activating a critical genetic element that was earlier found to regulate the earliest steps of [human embryo development](#) after fertilization. "Using this technology, [pluripotent stem cells](#) were obtained that resembled very closely typical early embryonal [cells](#)," Professor Kere says.

The discovery also suggests that it might be possible to improve many other reprogramming tasks by addressing genetic elements typical of the

intended target cell type.

"The technology may find practical use in bio banking and many other tissue technology applications," says Ph.D. student, MSc Jere Weltner, the first author of the article published in *Nature Communications*. "In addition, the study opens up new insights into the mechanisms controlling early embryonic gene activation."

**More information:** Jere Weltner et al, Human pluripotent reprogramming with CRISPR activators, *Nature Communications* (2018).  
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