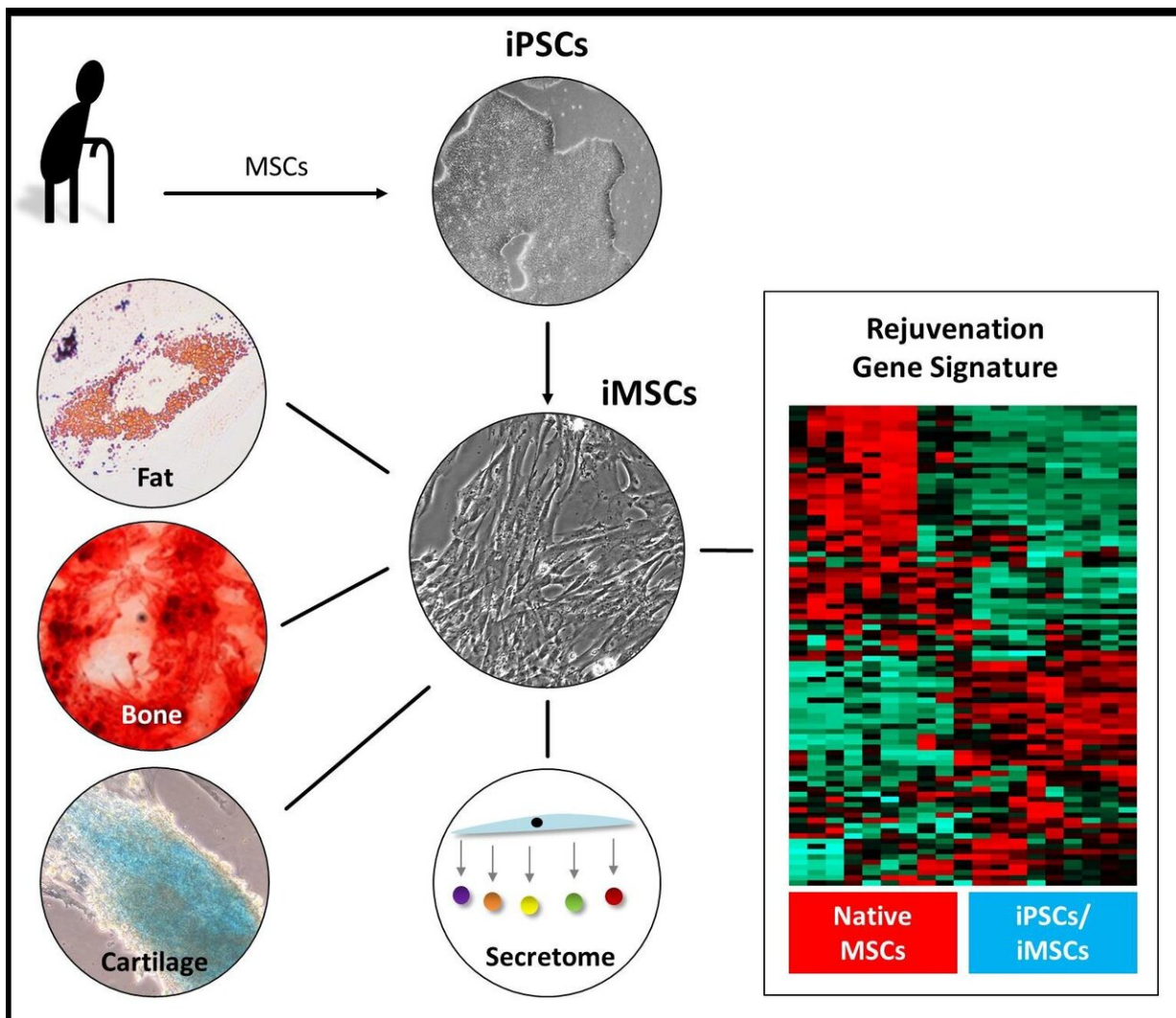


# Human iPSC-derived MSCs from aged individuals acquire a rejuvenation signature

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MSCs from elderly patients were reprogrammed into iPSCs and further differentiated into iMSCs. These iMSCs are endowed with typical MSC features such as making bone, cartilage and fat. The iMSCs acquired a gene signature

associated with rejuvenation and secreted factors present in native MSCs, thereby emphasizing their clinical potential. Credit: Spitzhorn LS, et. al.

The use of primary mesenchymal stem cells (MSCs) is fraught with aging-related shortfalls such as limited expansion and early senescence. Human induced pluripotent stem cells (iPSCs) derived MSCs (iMSCs) comprise a useful, clinically relevant source of MSCs that circumvent these aging-associated drawbacks. The importance of this concept is manifested by the successful phase 1 clinical trial for the treatment of GvHD by Cynata Therapeutics and the National Health Service of the U.K. A phase 2 trial is planned for 2019. This trial is independent of the work carried out by Spitzhorn et al.-described below.

A collaborative study coordinated by the Institute for Stem Cell Research and Regenerative Medicine analysed the acquisition of rejuvenation-associated hallmarks in iMSCs. In their study, the team compared cellular features, transcriptomes and secretomes of iMSCs differentiated from embryonic stem cells (ESCs-H1) and iPSCs, emanating from MSCs isolated young and elderly individuals. The generated iMSCs (irrespective of source) met the criteria set out for MSCs and dendrogram analyses confirmed that the transcriptomes of all iMSCs clustered together with the parental MSCs and distinct from pluripotent stem cells.

Irrespective of donor age and initial cell type, iMSCs acquired a rejuvenation-associated 50-gene comprising signature which is also expressed in [pluripotent stem cells](#) but not in the parental MSCs.

Significantly, in terms of [regenerative medicine](#), iMSCs acquired a secretome similar to that of primary MSCs, thus highlighting their ability to act via paracrine signalling.

The iMSC concept has enabled circumventing the drawbacks associated with the use of adult MSCs and thus provide a promising tool for use in various clinical settings in the future.

**More information:** Lucas-Sebastian Spitzhorn et al, Human iPSC-derived MSCs (iMSCs) from aged individuals acquire a rejuvenation signature, *Stem Cell Research & Therapy* (2019). [DOI: 10.1186/s13287-019-1209-x](https://doi.org/10.1186/s13287-019-1209-x)

Provided by Heinrich-Heine University Duesseldorf

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