

## Human Dental Pulp Stem Cells as a Patient-in-a-Dish Model for Charcot-Marie-Tooth Disease Type 1A

**Commonly used acronym:** DPSC-SC CMT1A Created on: 20-08-2024 - Last modified on: 23-08-2024

## Contact person

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#### Organisation

Name of the organisation University of Hasselt (UHasselt) Department BIOMED Specific Research Group or Service Team FIERCE Country Belgium Geographical Area Flemish Region

#### Partners and collaborations

University of Hasselt (UHasselt)

# SCOPE OF THE METHOD

The Method relates to	Human health
The Method is situated in	Basic Research, Translational - Applied Research
Type of method	In vitro - Ex vivo
Specify the type of cells/tissues/organs	Dental pulp stem cells

## DESCRIPTION

#### Method keywords

Disease modeling mesenchymal stem cell Peripheral neuropathy Lentiviral transduction CRISPR-Cas9 cellular differentiation Patient-derived

#### Scientific area keywords

Charcot-Marie-Tooth disease type 1A demyelination basic research Schwann cells adult stem cells Human Stem cells

#### Method description

Dental pulp stem cells (DPSC) are mesenchymal stem cells residing within the inner mucoid core (dental pulp) of teeth, responsible for tissue turnover and regeneration. Since third molars, or wisdom teeth, are frequently extracted for orthodontic reasons, DPSC are a highly accessible stem cell source. DPSC exhibit high proliferation rates and can be cryopreserved for long periods, rendering them suitable for biobanking. In addition, DPSC are embryonically derived from the neural crest lineage, sharing their origin with myelinating Schwann cells. Hence, we have developed a protocol to differentiate human DPSC towards functional Schwann cells called DPSC-SC. Furthermore, we have implemented these DPSC-SC as a novel research model for Charcot-Marie-Tooth disease type 1A (CMT1A). CMT1A is the most common demyelinating peripheral neuropathy. Previous research has evidenced that CMT1A animal models lack translatability and that human models are necessary to bridge the gap between preclinical and clinical research. We are using human DPSC-SC for CMT1A modeling by mimicking the disease using lentiviral transduction and CRISPR-Cas9 while also building a biobank of patient-derived DPSC-SC.

#### Lab equipment

Laminar Flow Cabinet, Incubator, Centrifuge

#### Method status

Still in development History of use Internally validated Published in peer reviewed journal

# **PROS, CONS & FUTURE POTENTIAL**

#### Advantages

- Human disease model,
- Highly accessible,
- Cost-effective,
- Unique DPSC differentiation potential allows for derivation of more mature Schwann cells,
- Possibility of genetically engineering cells,
- Novel patient-derived biobank representing the hetereogeneity of the diesase,
- Drug screening.

#### Future & Other applications

DPSC are currently being used for multiple regenerative applications in many fields of science including dentistry, oncology, and cardiology. DPSC-SC have potential in all research fields related to peripheral neuropathies. Our CMT1A model will undergo further optimization and will be used for generating 3D co-cultures with neuronal cells to investigate myelination defects.

## **REFERENCES, ASSOCIATED DOCUMENTS AND OTHER INFORMATION**

#### References

Martens W, Sanen K, Georgiou M, et al. Human dental pulp stem cells can differentiate into Schwann cells and promote and guide neurite outgrowth in an aligned tissue-engineered collagen construct in vitro. FASEB J. 2014;28(4):1634-1643. doi:10.1096/fj.13-243980

#### Associated documents

The FASEB Journal - 2013 - Martens - Human dental pulp stem cells can differentiate into Schwann cells and promote and (2).pdf

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