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Lu S<sup>1</sup>, Lin A<sup>2</sup>, Chen K<sup>2</sup>, Chang L S<sup>2</sup>

1. Department of Urology, National Yang-Ming University, School of Medicine; Department of Urology, Taipei City Hospital, 2. Department of Urology, Taipei-Veterans General Hospital and National Yang-Ming University, School of Medicine

# CHARACTERIZATION OF SMOOTH MUSCLE DIFFERENTIATION OF PURIFIED HUMAN SKELETAL MUSCLE DERIVED CELLS FOR UROLOGICAL REGENERATION

#### Hypothesis / aims of study

The purpose of this study is to characterize the smooth muscle differentiation of purified human skeletal muscle derived cells (hMDCs) for bladder reconstitution and management of stress urinary incontinence.

#### Study design, materials and methods

The isolation and purification of hMDCs were conducted by modified preplate technique and Dynal CD34 cell selection. Smooth muscle cell differentiation was induced by the use of smooth muscle induction medium (SMIM) and low serum medium. The gene expressions at the mRNA and protein levels of undifferentiated and differentiated hMDCs were tested by RT-PCR, western blot and immunofluorescence studies.

#### Results

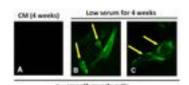
Western blot and immunofluorescence studies demonstrated the purified hMDCs cultured in SMIM for 4 weeks expressed significant amount of smooth muscle myosin heavy chain (MHC) and alpha-smooth muscle actin (ASMA). The cells cultured in low serum medium for 4 weeks also expressed ASMA, while the control group did not. RT-PCR analysis showed increased gene expression of smooth muscle markers, such as ASMA, Calponin, SM22, Caldesmon, Smoothelin, and MHC when purified hMDCs were exposed to SMIM for 2 and 4 weeks when compared to the controls.

#### Interpretation of results

The smooth muscle differentiation of purified human MDCs was justified. These cells may be used for potential biomaterials for bladder reconstitution and the treatment for stress urinary incontinence in the future.

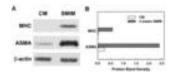
## Concluding message

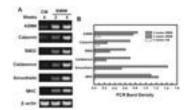
We confirmed the smooth muscle differentiation capability of purified hMDCs. The gene expression of smooth muscle differentiation of purified hMDCs was characterized. These cells may be potential biomaterials for human tissue regeneration.



MHC SMIM (4 weeks)

ASMA





# References

- 1. 7. Shing-Hwa Lu, An-Hang Yang, Chou-Fu Wei, Kuang-Kuo Chen, and Luke S. Chang. Isolation and characterization of human muscle derived cells. Urology 2009 Aug; 74(2):440-45
- 2. 3. Shing-Hwa Lu, An-Hang Yang, Kuang-Kuo Chen, Han Sun Chiang, and Luke S. Chang. Purification of human muscle derived cells using immunoselective method for urological regeneration. BJU Int; 2010 Jun; 105(11):1598-603.
- 3. 2. Shing-Hwa Lu, An-Hang Yang, Chou-Fu Wei, Han Sun Chiang, and Michael B. Chancellor. Multi-potent differentiation of the human purified muscle-derived cells: potential for tissue regeneration. BJU International, 2010 March;105(8):1174-1180.

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Is this a clinical trial?	No
What were the subjects in the study?	HUMAN
Was this study approved by an ethics committee?	Yes
Specify Name of Ethics Committee	IRB of Taipei-Veterans General Hospital in Taipei Taiwan with the
	IRB number: VGHIRB No: 92-11-07A
Was the Declaration of Helsinki followed?	Yes
Was informed consent obtained from the patients?	Yes