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## **Supplemental Information**

### **Reprogramming of Urine-Derived Renal Epithelial Cells into iPSCs Using srRNA and Consecutive Differentiation into Beating Cardiomyocytes**

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## SUPPLEMENTARY DATA

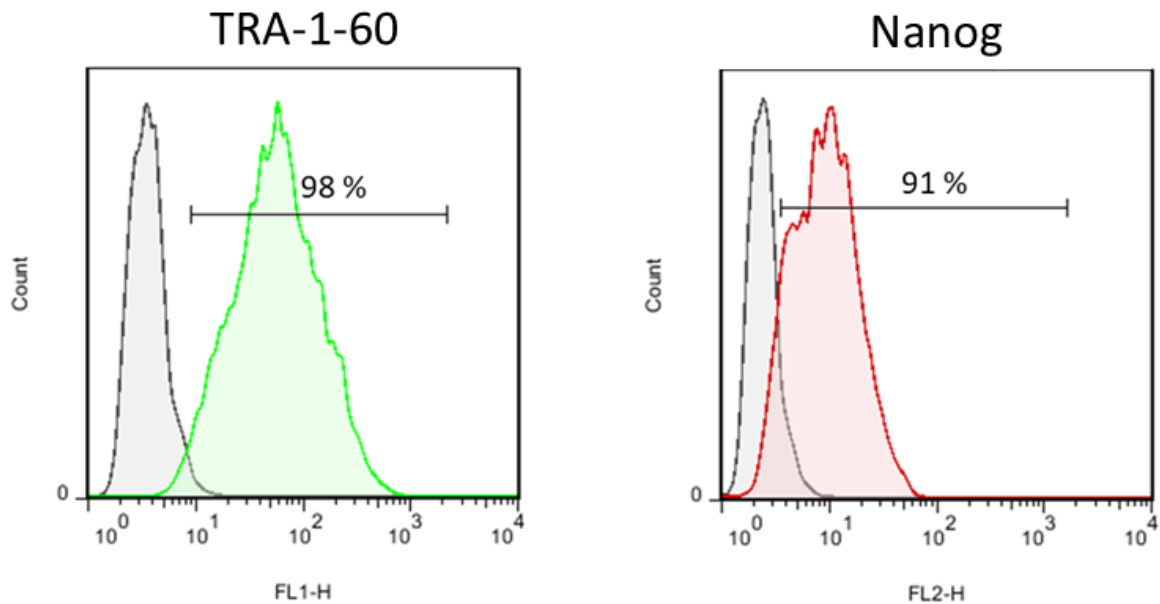
**TABLE 1**

**Table 1: List of primers used for qRT-PCR analysis.**

Gene	Forward primer (5'→3')	Reverse primer (5'→3')
<b>pluripotency marker</b>		
GAPDH	TCAACAGCGACACCCACTCC	TGAGGTCCACCACCCTGTTG
Oct4 <sup>33</sup>	AGCGAACCAGTATCGAGAAC	TTACAGAACCACACTCGGAC
Sox2 <sup>33</sup>	AGCTACAGCATGATGCAGGA	GGTCATGGAGTTGTACTGCA
Nanog <sup>33</sup>	TGAACCTCAGCTACAAACAG	TGGTGGTAGGAAGAGTAAAG
Lin28	CTTCTTCTCCGAACCAACC	CAGCCACCTGCAAACCTG
E-Cadherin	TATACCCTGGTGGTTCAAGC	CACCTGACCCTTGTACGTG
Klf4 <sup>33</sup>	TCTCAAGGCACACCTGCGAA	TAGTGCCTGGTCAGTTCATC
cMyc <sup>33</sup>	ACTCTGAGGAGGAACAAGAA	TGGAGACGTGGCACCTCTT
<b>srRNA specific marker</b>		
nsP2	TCCACAAAAGCATCTCTCGCCG	TTTGCAACTGCTTCACCCACCC
nsP4	TTTTCAAGCCCCAAGGTCGCAG	TGTTCTGGATCGCTGAAGGCAC
<b>cardiomyocyte marker</b>		
ANP	CAGACCAGAGCTAATCCCAT	GTCCAGCAAATTCTTGAAATCC
cTnT	TTACATCCAGAAGACAGAGCG	TCTCCCTCAGCTGATCTTCAT
MHC6	GAAGCACCAAGATGACCGATG	CTCTGACTTGCGGAGGTACT
ACTC1	ATGTGTGACGACGAGGAGAC	ACCCACCATAACTCCCTGGT

**Abbreviations:** GAPDH: Glyceraldehyde-3-phosphate dehydrogenase, Oct4: Octamer binding transcription factor 4, Sox2: Sex determining region Y-box 2, E-Cadherin: Epithelial cadherin, Klf4: Krüppel-like factor 4, c-Myc: Cancer myelocytomatosis, nsP: Non-structural protein, ANP: Atrial natriuretic peptide, cTnT: Cardiac troponin T, MHC6: Myosin heavy chain 6, ACTC:  $\alpha$ -actin, cardiac muscle.

## SUPPLEMENTARY FIGURES



**Supplementary Figure 1:** Pluripotency marker analysis of REC-iPSCs at passage 25 after initial picking. Flow cytometric measurement of TRA-1-60 and Nanog stained cells.

### VIDEO 1

**Characterization of contracting cardiomyocytes:** Recordings of 30 s (7 pictures/s) showing wide ranges of motion and directional synchronous contractions. Direction is indicated by arrows using Matlab application Motion GUI.

### VIDEO 2

**Calcium ion staining of contractile cardiomyocytes:** Fluorescent calcium indicators were used to visualize the intracellular calcium flux during contraction. Recordings of 30 s (7 pictures/s) showing Ca<sup>2+</sup> transients in the cardiomyocyte culture.