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**Supplemental Information**

**Anti-apoptotic Regulation Contributes to the Successful Nuclear Re-programming Using Cryopreserved Oocytes**

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**Title: Anti-apoptotic regulation contributes successful nuclear reprogramming using cryopreserved oocytes**

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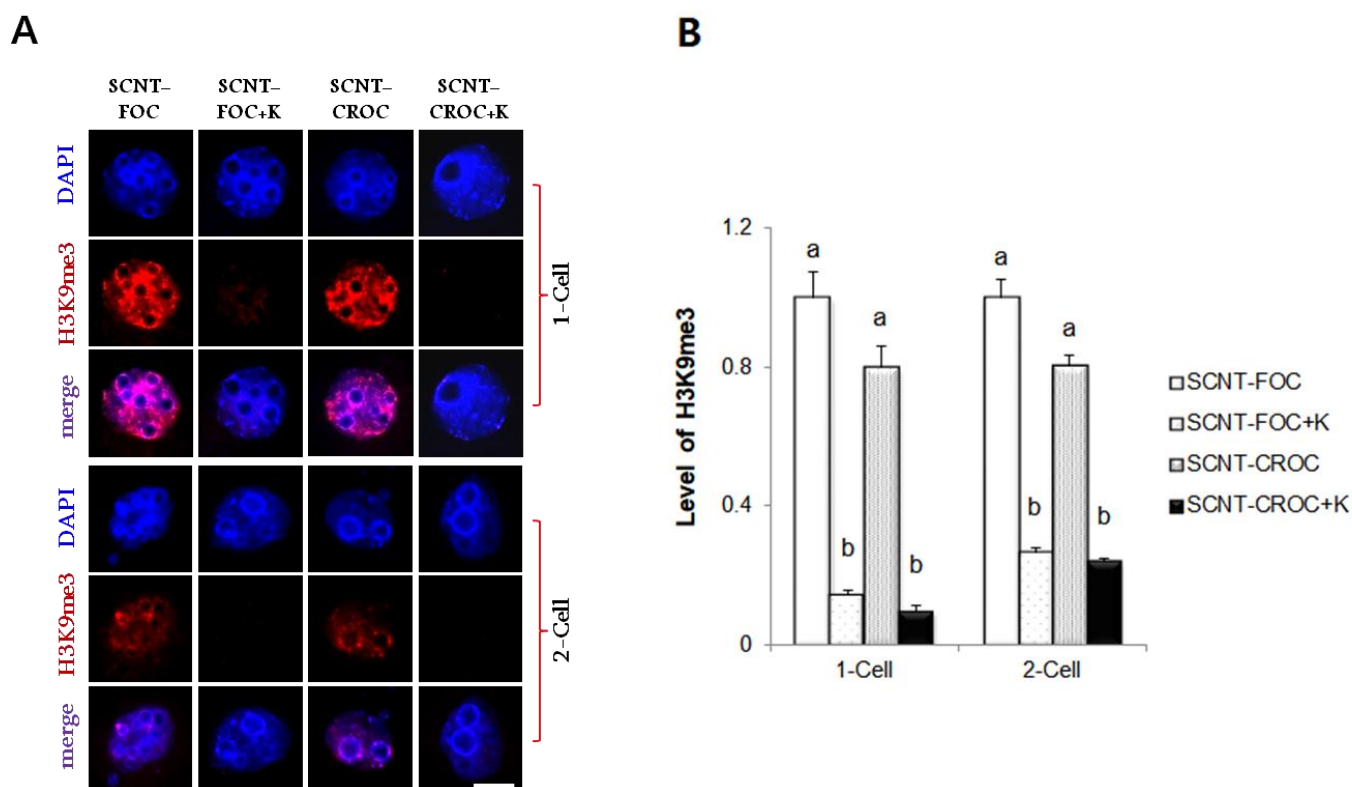
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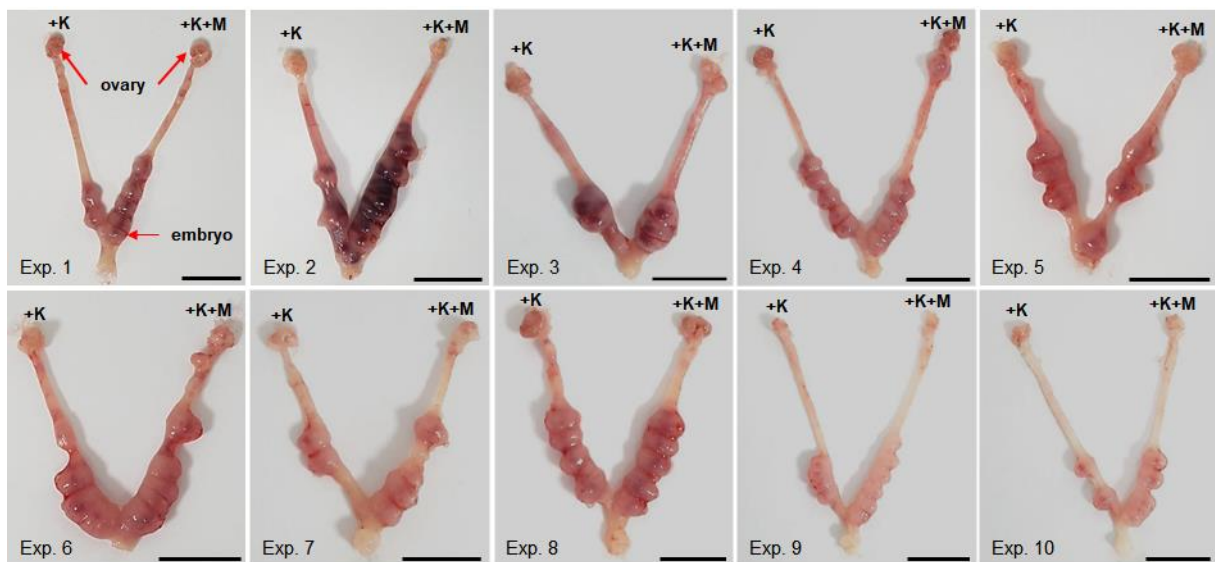
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**Running Head:** *Somatic cell nuclear transfer using cryopreserved oocytes*

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**FIGURE S1. Injection of *Kdm4a* mRNA removes H3K9me3 of SCNT embryos using fresh oocyte cytoplasm (SCNT-FOC) and cryopreserved oocyte cytoplasm (SCNT-CROC) at the 1-cell and 2-cell stage (related to Table 1). A) Representative nuclear images of 1-cell and 2-cell stage SCNT embryos stained with anti-H3K9me3 and DAPI. Shown in each panel is a nucleus of a single blastomere. Scale bar, 10  $\mu$ m. B) Bar graph showing reduced expression intensity of H3K9me3 between cloned embryos from SCNT-FOC and SCNT-CROC groups at the 1-cell and 2-cell stage.**



**FIGURE S2. Photographs of uteri with implantation sites at day 7 (related to Fig.4).** Allows indicate ovary and implantation site, respectively. In 9 out of 11 experiments, the number of implantation sites was increased in the SCNT-CROC+M+K group compared to those in the SCNT-CROC+K group.

## Reference of Table S1 and S2

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