A novel cytoplasmic adaptor for RAR and TR functions as a derepressor of RAR in the absence of retinoic acid*

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Running title: CART1 is a derepressor of RAR

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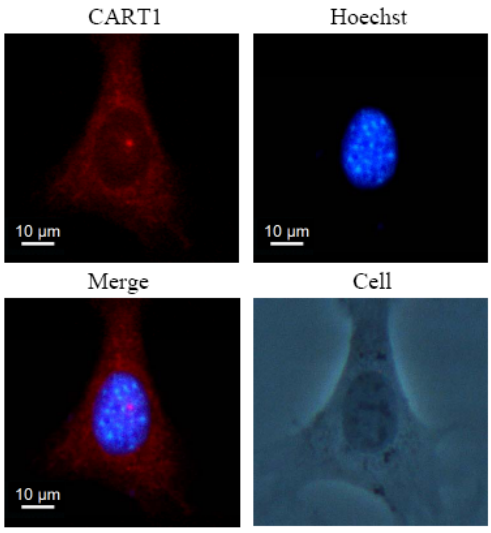
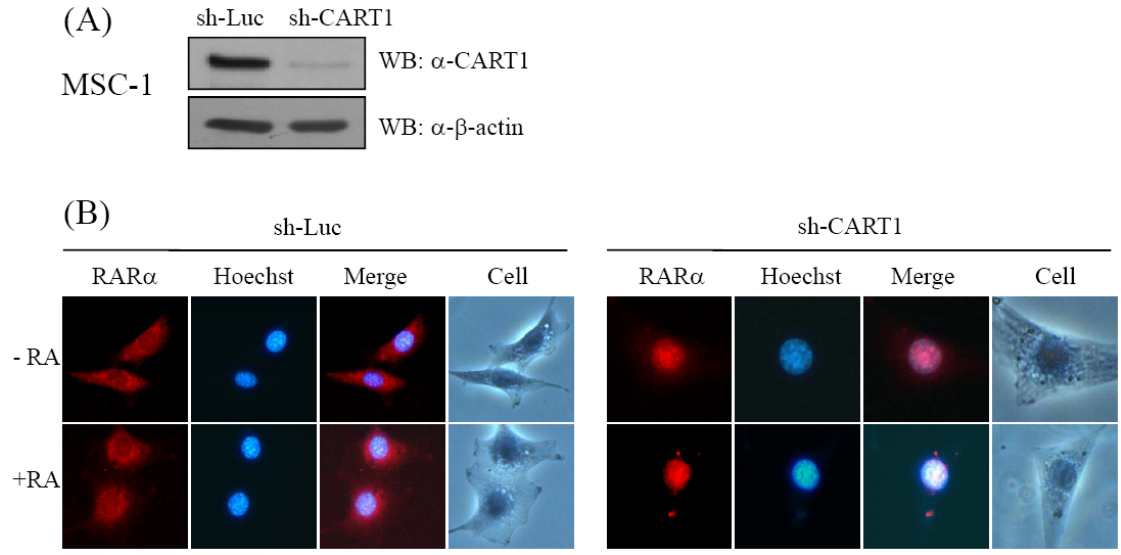
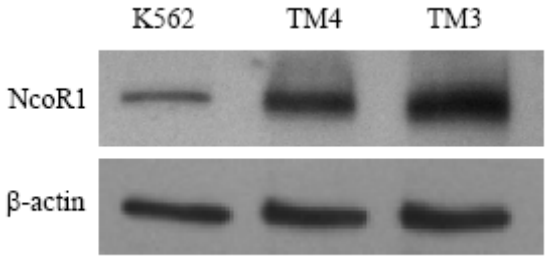


Supplemental Fig. 1. Subcellular location of endogenous CART1 in TM4 cells. TM4 cells were fixed and permeabilized. Cells were then stained with rabbit anti-CART1 polyclonal antibody and Texas red-conjugated anti-rabbit antibody, and observed by fluorescence microscopy. Hoechst was used to localize chromosomal DNA in the nucleus.



Supplemental Fig. 2. Effect of CART1 knockdown on subcellular location of RAR in MSC-1 mouse Sertoli cells. *(A)* Expression of CART1. MSC-1 cells were tranfected with either control small hairpin (sh) or CART1-specific sh RNA using Lipofectamine. CART1 expression was monitored by WB. β-actin was used as an internal control. *(B)* Subcellular location of endogenous RARα. MSC-1 cells were trasfected as indicated, fixed, and permeabilized. Cells were then stained with rabbit anti-RARα polyclonal antibody and Texas red-conjugated anti-rabbit antibody, and observed by fluorescence microscopy.



Supplemental Fig. 3. Expression of NcoR1. Cellular extracts were prepared from various cell lines as indicated and 80 μg lysates were subjected to WB using anti-NcoR1 antibody (1:250; Santa Cruz Biotechnology, Santa Cruz, CA; sc-1609). β-actin was used as an internal control.