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Effect of rtTA-M2 and rtTA on basal and inducible TRE-mediated transcription.

(a) Structures of expression units. CMVpro; cytomegalovirus promoter. (b) NIH3T3 cells were

transiently co-transfected with the pTRE-Luc reporter plasmid (10 ng), together with prtTA-

M2 (100 ng) or prtTA (100 ng). pCH110 (100 ng) was used as a control for transfection

efficiency. Empty vector (pcDNA3) was used to standardize the total amount of transfected

DNA (210 ng). Dox (10-1000 ng/ml) was added to the cell culture medium. * indicates p<0.05

compared with control cells transfected with pcDNA3 in the absence of Dox (Student's t test).

indicates p<0.05 compared with cells expressing rtTA in the presence of each dose of Dox

(Student's *t* test).

Article title: Development of the tetracycline-dependent transcription activator and repressor

co-expression system to tightly regulate and highly induce trangene expression

Journal name: Cytotechnology

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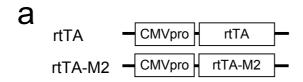
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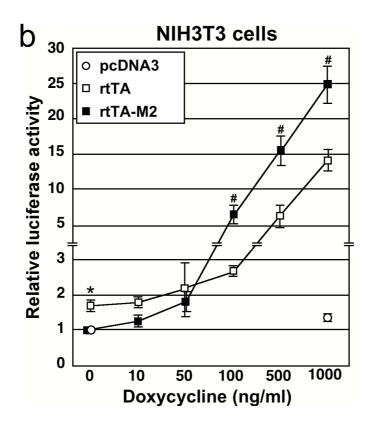
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