

autographic study using H³-actinomycin D (Schwartz BioResearch, New York: specific activity 8.4 Ci/m mole) also showed that actinomycin D entered to the nucleus of K3b/CR cells though in a reduced rate in comparison with the parental K3b cells. Therefore, persistence of RSV genome in K3b/CR cells treated with chromomycin or actinomycin D may be due chiefly to its specific mode of existence *i.e.* as provirus in the cells (21, 22) rather than due to the impermeability of cell membrane to the antibiotics.

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Erratum

In the paper by G. SIEGL, C. HALLAUER, and A. NOVAK: Parvoviruses as Contaminants of Permanent Human Cell Lines. IV. Multiplication of KBSH-Virus in KB-Cells, published in „Archiv für die gesamte Virusforschung“ **36**, 351—362 (1972), the legend for Fig. 5 on page 358 should be changed with the legend for Fig. 6 on page 359.