

Nucleotide sequence analysis of long terminal repeats of leukemogenic and non-leukemogenic MCF MuLVs

Theodore S.Theodore and Arifa S.Khan\*

Laboratory of Molecular Microbiology, National Institute of Allergy and Infectious Diseases, National Institutes of Health, Bethesda, MD 20892, USA
Submitted May 15, 1987

Accession no. Y00383

We sequenced the large terminal repeats (LTRs) present in infectious cloned DNA segments of leukemogenic, mink cell focus-forming MCF13 and non-leukemogenic, MCF111A murine leukemia viruses (MuLVs; 1). We compare these sequences with LTRs of AKR ecotropic (AKR ECO; 2), NZB xenotropic (NZB XENO; 3) and MCF247 (4) MuLV DNAs. MCF111A LTR is used as the "standard." Dots show identity; blank space, absence of bases; boxed nucleotides, regulatory signals; arrows, inverted repeats; underlines, regions of greatest nucleotide divergence; and brackets, enhancer region. Different nucleotides are shown. AKR ECO LTR was shortened by 99 bp by omission of the direct repeat. MCF111A LTR was identical except for one bp with AKR ECO LTR; MCF13 LTR had 94% base homology with NZB XENO LTR and 97% sequence homology with the LTR of leukemogenic MCF247 MuLV DNA.

Table with 5 columns: Sequence identifier (MCF 111A, AKR ECO, NZB XENO, MCF 247, MCF 13), Nucleotide sequence, and Position (100, 200, 300, 400, 500). Includes arrows and boxes highlighting specific regions.

\*To whom correspondence should be addressed

References 1.) Rowe, W.P., et al., (1980) Cold Spring Harbor Symp. Quant. Biol. 44, 1265-1268. 2.) van Beveren, C., et al. (1982) J. Virol. 41, 542-556. 3.) O'Neill, R.R., et al. (1985) J. Virol. 53, 100-106. 4.) Kelly, M., et al. (1983) J. Virol. 45, 291-298.