HL-60: Subcutaneous AML xenograft tumor model



Subcutaneous mouse tumor models

Subcutaneously implanted tumor cells represent a convenient means to test novel potential anticancer drugs *in vivo*. A large variety of human and murine cell lines derived from both, solid tumors or leukemias, covering a wide range of tumor geno- and phenotypes, have been adapted to grow in a murine host, and thus allow testing of a compound in the appropriate tumor model.

➤ HL-60 cells

Human HL-60 cells were isolated from a patient with AML (acute myeloid leukemia) and express FLT3 wild type.

A Hematoxylen-Eosin stained paraffin section of a subcutaneous HL-60 xenograft is shown on the right.

As routine quality controls, the cells are regularly checked for Mycoplasma contamination and authenticity (via STR DNA Typing).

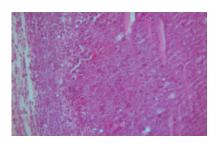


Figure 1: Hematoxylin-Eosin stained paraffin section of a subcutaneous HL-60 xenograft.

Expression of oncology relevant proteins

Expression data using western blotting and immunohistochemistry are available for a selection of protein kinases. For information, please inquire!

Tumor growth in vivo

HL-60 cells harvested from tissue culture flasks are implanted into the subcutaneous space of the left flank of the mice. Resulting tumors are monitored by calipering twice weekly.

Animal weights are measured three times weekly. Animal behaviour is monitored daily. All mice are maintained in separated isolated housing at constant temperature and humidity.

Accessory services: tumor wet weight and volume measurement at necropsy, blood sampling, flow cytometry, paraffin embedding of tumor tissue, histological & pathological analysis, cytokine determination, provision of tumor tissue for target validation.

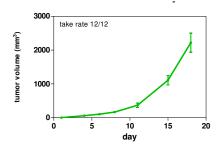


Figure 2: Tumor growth of HL-60 cells in a subcutaneous xenograft in vivo, tumor volume, mean values +/- SEM

Study example

In the study shown here, one group of mice bearing subcutaneous HL-60 xenografts was treated with Vincristin, the other group with vehicle only. Treatment started after randomisation when tumor volumes had reached a size of approximately 100-150 mm3.

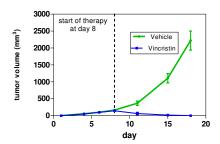


Figure 3: Effect of Vincristin on tumor growth of HL-60 cells in vivo. Tumor volume, mean values +/- SEM

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