

## In vivo Pharmacology: Xenograft (A549) Tumor Model

Species, strain, sex: Nude and athymic female Balb/c mice (CAnN.Cg-Fox1nu/Crl )  
 Number of animals /group: n=10  
 Pharmacological control: NA  
 Routes of administration: upon request  
 Treatment mode: upon request  
 Duration of dosing: 2-4 weeks

Host mice inoculated *s.c.* in the left axillary region with A549 lung adenocarcinoma sterile cell suspension. Tumors grew locally, encapsulated in a connective tissue capsule. Subsequently primary tumor growth monitored by periodic caliper measurements of superficial tumor's volume i.e. relative weight, once the tumor is sufficiently large, the mice are stratified by tumor size and randomly assigned to various cohorts and therapy can be started (usually after 1-3 weeks).

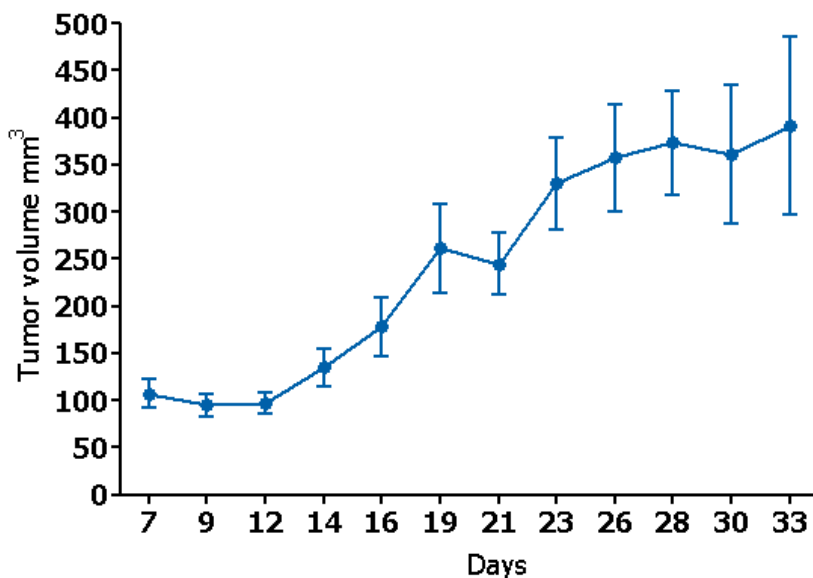
### Main read-outs:

BW  
 Relative and wet tumor Weight  
 Histopathology

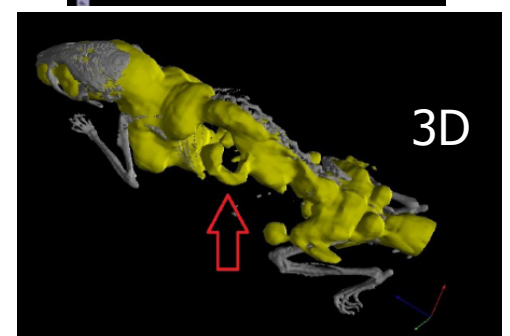
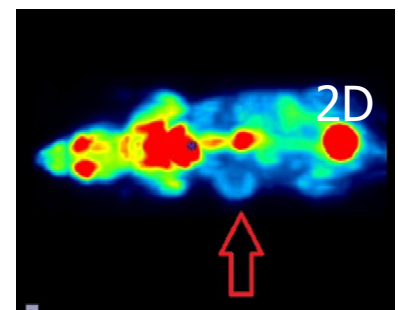
### Facultative read-outs:

Immunohistochemistry  
 Serum biomarkers  
 PET Scan imaging (PET/CT)

Tumor Growth Rate for A549 Cells Inoculated into Female Athymic Nude Mice  
 mean  $\pm$  SD; N=10 at start



### PET Scans on day 33



### References

Dasa Seveljevic-Jaran & Boska Hrvacic (2010): Reduction and refinement in a xenograft tumor efficacy study. FELASA symposium, Helsinki, Finland.