

Villin-Cre Apcfl/+ KRasG12D/+ Mouse

Catalogue number: 152932

Sub-type: Mouse

Images:

Contributor

Inventor: Owen Sansom

Institute: Cancer Research UK, Glasgow: The Beatson Institute

Images:

Tool details

***FOR RESEARCH USE ONLY**

Name: Villin-Cre Apcfl/+ KRasG12D/+ Mouse

Alternate name: GTPase KRasK-Ras 2Ki-Rasc-K-rasc-Ki-ras

Class:

Conjugate:

Description: Oncogenic mutations in the K-ras gene occur in 50% of human colorectal cancers. However, the precise role that K-ras oncogenes play in tumor formation is still unclear. To address this issue, we have conditionally expressed an oncogenic K-rasD12 allele in the small intestine of adult mice either alone or in the context of Apc deficiency.

Purpose:

Parental cell:

Organism:

Tissue:

Model: Conditional Knock-Out

Gender:

Isotype:

Reactivity:

Selectivity:

Host:

Immunogen:

Immunogen UNIPROT ID:

Sequence:

Growth properties:

Production details:

Formulation:

Recommended controls:

Bacterial resistance:

Selectable markers:

Additional notes:

Target details

Target: KRasG12D

Target alternate names:

Target background:

Molecular weight:

Ic50:

Applications

Application:

Application notes:

Handling

Format:

Concentration:

Passage number:

Growth medium:

Temperature:

Atmosphere:

Volume:

Storage medium:

Storage buffer:

Storage conditions:

Shipping conditions: Embryo/Spermatozoa- Dry Ice

Related tools

Related tools:

References

References: HCMV-infected cells maintain efficient nucleotide excision repair of the viral genome while abrogating repair of the host genome. ; O'Dowd et al. 2012. PLoS Pathog. 8(11):e1003038.

PMID: 23209410.

CancerTools.org