Datasheet for ABIN925589

10X TBS Fish Gel Concentrate (Azide and Mercury free)

2 Images

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Overview

Quantity:	100 mL

Product Details

Purification:	This product was aseptically filtered through a Millipore 0.22 micron filter into clean, pre-sterilized containers. The product was tested on trypticase soy agar for 24 hours, 48 hours and 72 hours and was found to be negative for bacteria.
Endotoxin Level:	Low Endotoxin : No

Application Details

Application Notes:	This product is a 10X concentrated stock solution. Prepare a 1X working solution by diluting 1 part 10X concentrate with 9 parts distilled-deionized water or equivalent. 10X TBS Fish Gel Concentrate consists of 1.0 M Tris hydrochloride, 1.5 M sodium chloride and Fish Gelatin at pH 7.5. A proprietary combination of stabilizers and preservatives are used that are azide and mercury free.
Comment:	Highly sensitive ELISA assays require minimal non-specific interactions. Non-specific binding results in elevated background levels and a decrease in signal-to-noise ratios. Blocking reagents act to minimize non-specific interactions of secondary reactants with each other and with the primary solid phase binding sites. Multiple formulations offer greater flexibility in experimental design.
Restrictions:	For Research Use only

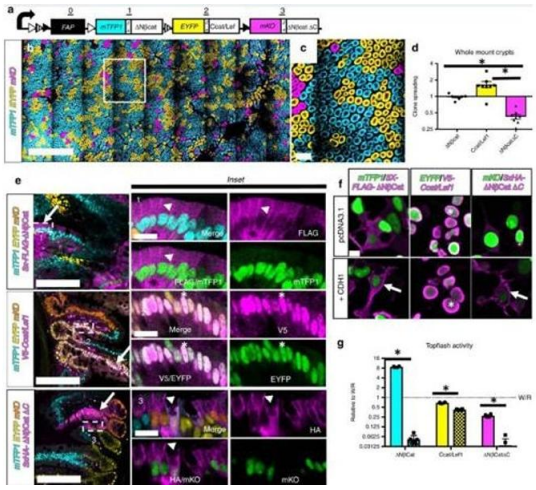
Handling

Format:	Liquid

Handling

Precaution of Use:	No special shipping conditions or precautions are required.
Handling Advice:	Protect from moisture and light.
Storage:	4 °C
Storage Comment:	Store container at 4 °C before opening.
Expiry Date:	Expiration date is six (6) months from date of opening.

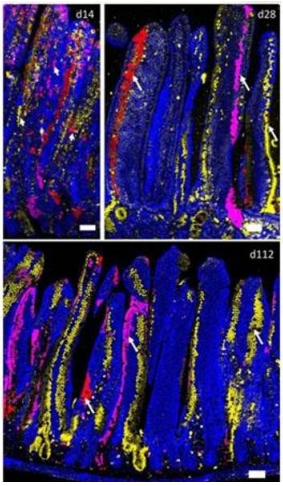
Images



Fluorescence Microscopy

Image 1. Widespread expansion of oncogenic clones during perinatal development. a Diagram of MCAT-Crainbow mice. b MCATVilCre small intestine (N=10 mice, 3-6 weeks of age) prepared as a wholemount and confocal imaged. c Inset in ""b"" at higher magnification. d MCATVilCre Crypts were color segmented, counted and normalized to the positional bias calculated in NCATVilCre mice. Asterisk denotes statistical significance by one-way ANOVA (mTFP1 vs. EYFP: p=0.003, mTFP1 vs. mKO: p=0.016, EYFP vs. mKO=3e-6). e Immunostaining for FLAG, V5, or HA epitopes (magenta) specific to each ßcat isoform in MCATVilCre small intestine vibratome slices and merged with fluorescent lineage markers (mTFP1: cyan, EYFP: yellow, and mKO: orange). Arrows denote isoform expression with cognate lineage reporter (FLAG and mTFP1, V5 and EYFP, and HA and mKO). Corresponding insets depict higher magnification images. Arrowheads denote membrane-localized ßcat, whereas asterisk denotes nuclear-localized ßcat. Epitope stains (magenta) are also presented as merged and as a single-channel image with its cognate fluorescent lineage reporter (green). f HEK cells were transiently transfected with MCAT isoforms, fixed, stained, and imaged for the indicated epitope (magenta) and fluorescent reporter (green). Cells were also cotransfected with epithelial cadherin (CDH1) as indicated.

Arrows denote sequestration of ßcat at the plasma membrane, and the asterisk denotes nuclear ßcat. g Wnt signalling activity for each oncogene in the absence of CDH1 (solid bar) or in the presence of overexpressed CDH1 (hatched bar) (N=6 wells per condition and independently repeated in four experiments). TOP FLASH activity was normalized to WNT/RSPO-stimulated control cells (dashed line). Asterisk denotes statistical significance by two-way ANOVA and Bonferroni's multiple comparisons test (cyan<1e-6, yellow=0.01, magenta=0.02). (SEM included for each graph). Scale Bars=1mm in b, 100µm in c/e, 15µm in e: insets 1-3, and 10µm in f. Cells were fixed at room temperature for 15 min in 4 % PFA, washed once with PBS, and permeabilized/blocked in FISHX (0.25 % Triton-X diluted in 1 % Fish Gelatin (p/n MB-067-0100)) for 20 min at room temperature. Fig. 3. PMID: 31792216.



Fluorescence Microscopy

Image 2. LGR5-rainbow lineage tracing. LBOW mice were crossed to ROSA-CreER/T2 mice and intraperitoneally injected with 200 mg/kg tamoxifen every other day for a total of three injections. Mice (n = 4-6/time point) were chased for 14, 28, 56, 112, 224, or 365 days. The small intestine was harvested, sectioned with a Vibratome, fixed and permeabilized in FISHX comprising fish gelatin extract (p/n MB-067-0100) and 0.2 % Triton-X-100 for 30 min, stained for EYFP and mCherry/E2-Crimson (p/n 600-401-379), and processed. ections from each mouse intestine were tile-imaged by confocal microscopy. A subset of each tiled imaged that is representative for days 14 (d14), 28 (d28), and 112 (d112) is presented. (mKO1, blue, EYFP, yellow, mCherry, red, E2-Crimson, magenta) (scale bars, 50 µm). Small arrows depict small patches of clones, whereas large arrows depict full-length large clones. Figure 9. PMID: 28275053.