${\bf Package~`flowFitExampleData'}$

April 14, 2020

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Type Package				
Title Example data for the flowFit package				
Version 1.22.0 Date 2013-03-04 Author davide Rambaldi Maintainer Davide Rambaldi <davide.rambaldi@gmail.com></davide.rambaldi@gmail.com>				
			Description Two dataset that can be used to run examples from the flowFit vignette and examples	
			License Artistic-2.0	
			biocViews FlowCytometryData	
Depends R (>= 2.12.2), flowCore				
Imports methods				
git_url https://git.bioconductor.org/packages/flowFitExampleData				
git_branch RELEASE_3_10				
git_last_commit 9817a57				
git_last_commit_date 2019-10-29				
Date/Publication 2020-04-14				
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PKH26data genFitting: example data with PKH26 stain				
Description				
example data with PKH26 dye. Two samples: NONPROL (parent population) and PROL (prolating population).	ifer-			
Usage				
data(PKH26data)				

2 QuahAndParish

Format

The format is an object of class flowSet with 2 flowFrame

QuahAndParish	Example dataset from: New and improved methods for measuring
	lymphocyte proliferation in vitro and in vivo using CFSE-like fluores-
	cent dyes (Benjamin J C Quah and Christopher R Parish, 2012)

Description

Detection of lymphocyte division by carboxyfluorescein diacetate succinimidyl ester (CFSE), Cell Trace Violet (CTV) and Cell Proliferation Dye eFluor 670 (CPD) in vitro.

Usage

data(QuahAndParish)

Format

The format is an object of class flowSet with 4 objects of class flowFrame

- Fig 2a All CD4 T Nonstim.fcs Control sample including non-activated cells (non-dividing) labelled with CFSE, CPD and CTV
- 2. Fig 2a CFSE CD4 T Stim.fcs CD4 T cells stained with CFSE
- 3. Fig 2a CPD CD4 T Stim.fcs CD4 T cells stained with CPD
- 4. Fig 2a CTV CD4 T Stim.fcs CD4 T cells stained with CTV

The phenodata lists:

Filename The filename

SampleType The sample type (Nonstim or Stim)

Stain Stain type
CellType Cell type

Details

This QuahAndParish dataset represents the measurements of CD4 T cells division by CFSE, CTV and CPD in vitro. Spleen cells from B6 mice were labelled with $10\mu M$ CFSE, CTV and/or CPD and cultured for 4 days in the presence of a range of polyclonal stimuli that activate T and B cells. Viable CD4+ cells were discriminated using specific antibody staining. The dataset represent the measurements used in figure 2a (CD4+ population) from the paper: New and improved methods for measuring lymphocyte proliferation in vitro and in vivo using CFSE-like fluorescent dyes (Benjamin J C Quah and Christopher R Parish, 2012).

References

1. Benjamin J.C. Quah, Christopher R. Parish, New and improved methods for measuring lymphocyte proliferation in vitro and in vivo using CFSE-like fluorescent dyes, Journal of Immunological Methods, Volume 379, Issues 1-2, 31 May 2012, Pages 1-14, ISSN 0022-1759, 10.1016/j.jim.2012.02.012.

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