

7-AAD

Double-stranded nucleic acids stain that doesn't readily pass through intact cell membranes but can penetrate cell membranes of dying or dead cells

## **Product Description**

Name :	7-Aminoactinomycin D (7-AAD)	H <sub>2</sub> N $\downarrow$	
<b>Catalog Number :</b>	FP-132303, 1mg	U	
	FP-1J7621, 1 ml in DMSO:water (1:1)	L-Pro -D-ValL-Thr -C	C-L-Thr-D-Val-L
Structure :	$C_{62}H_{87}N_{13}O_{16}$	Sar-L-MeVal-O O	O O-L-MeVal-
Molecular Weight :	MW= 1270.45		
Solubility:	DMSO, DMF and CH <sub>3</sub> OH		
Absorption / Emission :	$\lambda_{exc} = (CH_3OH) = 546/648 \text{ nm}$		

Storage:-20°CProtect from light and moisture

## Introduction

7-Aminoactinomycin D (7-AAD) is a fluorescent chemical compound with a strong affinity for DNA. It is used as a fluorescent marker for DNA in fluorescence microscopy and flow cytometry. It intercalates in double-stranded DNA, with a high affinity for GC-rich regions, making it useful for chromosome banding studies.

7-AAD is compatible with most blue and green fluorophores – and even many red fluorophores – in multicolour applications. 7-AAD/DNA complexes can be excited at 488 nm with an argon-ion laser, and has a large Stokes shift with an emission maxima of 647 nm.

7-AAD does not readily pass through intact cell membranes; if it is to be used as a stain for imaging DNA fluorescence, the cell membrane must be permeabilized or disrupted. 7-AAD is also used as a cell viability stain. Cells with compromised membranes will stain with 7-AAD, while live cells with intact cell membranes will remain dark.

### **Directions for use**

### Guidelines for use in flow cytometry

#### **Stock Solution**

For long-term storage, store unopened vials of 7-AAD in the freezer. Dissolve 1 mg of 7-AAD powder by adding 50 microliters of absolute methanol directly to the vial. Mix well and add 950 microliters of 1 X PBS with Ca2+ and Mg2+ to achieve a concentration of 1 mg/ml. Store solution tightly closed and protected from light at 4°C. This solution is stable for several months.



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#### FT-132303

#### Protocol



Stain your cells as outlined in the protocol for single color or dual-color staining with FITC and/or PE-labeled monoclonal antibodies.

After the last washing step resuspend your cells as usual in 1 ml of buffer for analysis. If you want to assess viability of your samples add 1-2 microliters of the 7-AAD stock solution to each tube and mix well. Keep the samples in this solution at 4°C protected from light for approximately 20 minutes or until analysis on the flow cytometer.

NOTE: This method can now be used in combination with formaldehyde fixation of samples. Samples are first stained with 7-AAD, then fixed in 1% formaldehyde that contains 2-5 microliters/ml of actinomycin D (Fetterhoff). 7-AAD can be used for dead cell exclusion on samples that are stained with PE (phycoerythrin)-conjugated antibodies, because the emission spectra of 7-AAD and PE can be easily separated on the flow cytometer.

Other protocol may be found in the literature.

### References

- **Boomershine CS** *et al.*,Autoimmune pancreatitis results from loss of TGFβ signalling in S100A4-positive dendritic cells, *Gut*, 58: 1267 1274 (2009) <u>Article</u>
- Fetterhoff TJ *et al.* Fluorescent detection of non-viable cells in fixed cell preparations. *Cytometry* 14 (Suppl. 6):27 (1993)
- Torsten K. *et al.*,Inducible expression of EVI1 in human myeloid cells causes phenotypes consistent with its role in myelodysplastic syndromes, *J. Leukoc. Biol.*, 10.1189 (2009) <u>Article</u>

## Technical and scientific information

### **Related products**

• Annexin V-FP488, <u>FP-BH9390</u>

• Propidium iodide, FP-36774A

• Annexin V, R-PE, FP-AH191A

# **Ordering information**

<u>Catalog size quantities and prices may be found at www.interchim.com/</u> Please inquire for higher quantities (availability, shipment conditions).

For any information, please ask : FluoProbes<sup>®</sup> / Interchim; Hotline : +33(0)4 70 03 73 06

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