Indian Institute of Technology Bombay



Date:

Flow Cytometry facility

Requisition form for using FACS facility (facs@iitb.ac.in)

1) Name of the User*:
2) Name of the Organization*:
3) Name of the HOD/ Principal Investigator*:
4) User's Email ID:
5) Tel No*:

	Sample Information
Experiment name *	Others
Experiment Type *	• Acquisition/Sample analysis O Sorting with acquisition O Data analysis (using FlowJo software)
Number of controls *	
Number of test samples *	
Size of the cells/particles *	
Number of parameters to be analyzed *	
Name of Fluorophores	
Preferred date for the slot	

Acquisition/Sample analysis						
Laser & Filter selection :						
Blue (488nm Excitation) SC SSC	FITC (515-545nm)	DE (578-587nm)	OPE-Texas Red (600-620nm)	PerCp (667-702nm)	PerCP-Cy5.5 (675-715nm)	□ PE- Cy7/PerCp–Cy 7 (750-810nm)
UV (355nm)	DAPI (425-475nm)		□ Indo-1 Blue (515-545nm)		SP Red (645-695nm)	
Yellow Green (561 nm)	Ds Red/PE-561 (577-592nm)		mcherry/PE-Texas Red-561 (600- 620nm)		DPE -Cy5 (655- 685nm)	DE -Cy7-561 (750-810nm)
Red (633/637nm)	APC (655-670nm)		O APC -Alexa 700 (707-752nm)		APC-Cy7 (750-810nm)	
Violet (405nm)	Pacific Blue (425-475nm)		AmCyan (500-550nm)		Qdot 655 (655-685nm)	
If not sure about selection of filters & laser then list down the Excitation & Emission spectra of the dye/fluorophore in the additional information box provided.				/		
Additional information : (Constraints/Preferences/ etc.)						
I understand that for any experiment, if more than one filter is selected, compensation control for every fluorochrome will be needed for sample analysis					nple analysis	

along with the unstained control. The samples will be analyzed according to the choices recorded in the form.

Please fill below form only in case 'Sorting with acquisition' is opted.

Sorting with acquisition						
Number of samples to be sorted *						
Population to be sorted *						
Device to be used for sorting *	GFACS tube (5ml)	□ 15ml Falcon tube	96-well plate			
	24-well plate	□ 6-well plate	🗆 Agar plate			
Number of collection devices (tubes/plated) used for sorting *						
Type of sorting	O Sterile (Sterile sorting is in trial mode)	O Unsterile				
Additional information : (Constraints/Preferences/ etc.)						

□ I understand that for any experiment, if more than one filter is selected, compensation control for every fluorochrome will be needed for sample analysis along with the unstained control. The samples will be analyzed according to the choices recorded in the form.

Fields marked with * are mandatory fields.

- ✓ KINDLY ENSURE THAT THE GIVEN SAMPLE IS NOT INFECTIOUS OR POISONOUS OR TOXIC IN ANY WAY
- ✓ Whenever the results are used in the publications, appropriate acknowledgment of usage of IIT Bombay's Flow
 Cytometry central facility & IRCC must be mentioned. The details can be forwarded to <u>facs@iitb.ac.in</u>

Important information:

- 1. FACS1 refers to FACS machine located in CRNTS department while FACS2 refers to FACS machine located in BSBE department.
- 2. Acquisition/Sample analysis' refers to analyzing samples on Flow Cytometer and 'Sorting with acquisition' refers to analyzing and separately collecting desired population.
- 3. Minimum volume of sample required for acquisition is 500ul and the concentration of cells/particles required is 1x10⁶ cells or particles per ml.
- 4. Every sample for analysis should be accompanied with an Unstained control
- 5. Gating of population is experiment design specific and should be suggested by the user.

- 6. Gating can also be done offline by the user.
- 7. The samples will be analyzed on the channels using the lasers chosen by the user in the form. The user is therefore requested to pay utmost attention while selecting the lasers & channels.
- 8. Instrument is equipped to normally handle cells/particles in the size range 0.2 to 25 microns. While instrument cannot "strictly" handle cells/particles above 25 microns, it may be possible to analyze samples of size less than 0.2 microns. Such samples, that is, those containing particles less than 0.2 microns will be handled on a case-by-case basis and user requesting analyzes of such samples must discuss with the operator at facs@iitb.ac.in prior to submitting a request.
- 9. <u>Please refer to https://rnd.iitb.ac.in/research-facility/fluorescence-activated-cell-sorting-flow-cytometer-facility</u> for Instructions for sample preparation/submission
- 10. UV laser of FACS1 is non functional. However, for it is working for FACS2.
- 11. Mixed samples (e.g. blood samples or co culture samples) cannot be resolved efficiently on the basis of side scatter properties on FACS1. However, it can be resolved efficiently on FACS2.

I have read above mentioned Important information

For IITB use only

Date of sample receipt:	.Date of analysis:
Flow Cytometry facility machine name:	
Name of the Operator:	.Signature of Operator:
Registration number:	
Remarks:	