

Downstream BioProcessing Questionnaire

Please answer the following questions as completely as possible.

ı. Cu	stomer Information	
Cont	act Person	
Desi	gnation	
Department		
Company Name		
Contact Number		
Email Address		
II. Ex	cperimental Details	
	eneral Details	
1.	The target molecule to be purified	
2.	Size or M _r of target molecule	
3.	NaCl/pH/T ranges within which the target molecule is stable	□ NaCl: □ pH: □ T:
4.	Is the standard (purified target molecule) available?	□ Yes □ No
5.	Typical concentration of the target molecule (titer in the case of virus) in the sample that will be provided for the method/process development	t
6.	Main impurities in the sample (Specify concentrations of each)	 □ DNA □ Proteins □ Lipids □ Endotoxin □ Others:
7.	Sample volume/total virus titer is available for the method development	
8.	Current detection/ quantification method for the target molecule	 □ DNA/protein assays □ SDS PAGE □ qPCR □ ELISA □ Western blot □ Cell-based assays



		□ Others:	
9.	Desired concentration of the target molecule in the final product	Concentration: Unit of measure:	
10.	Purity requirements of the final product (Host cell protein/DNA concentration, endotoxin content)		
11.	Final formulation buffer		
12.	Other requirements regarding the method/process to be developed		
13.	Please describe upstream and downstream process detail or block scheme.		
b. Virus Products (Skip of not applicable)			
14.	Virus condition needed	□ Live□ Attenuated□ Killed	
15.	Virus Type	□ Wild type strain□ GMO	
16.	For GMO virus, please provide the detailed data/document descriabing preparation of the GMO. (e.g., constructs, vectors, expression system used)		
17.	What is the biosafety level of this virus? If safety assessment study is available, please provide the documentation.		