

The BioPro KS-y capsule filters are designed for protecting your critical and high value downstream systems.

These help in significant reduction of bioburden and complete removal of particulate contamination and are ideal for applications which do not require sterilization but where reduction in bio load in the process fluid is the objective.

These improve the process efficiency by reducing filter sizing and prolonging life of expensive sterilizing filters.

Special Features

- Validated for high bio-burden reduction
- High flow rates
- High throughput
- Low protein binding
- No media migrating
- Biologically inert
- Easy installation

Applications

- Clarification of cell harvest
- Buffer filtration
- In process protein filtration
- Prefiltration to sterile filtration
- Prefiltration to virus filtration

Ordering Information

Type	Size		Pore Size		Inlet/outlet		Radiation Sterilizable		Inline/T-Line		Sterility		Pack size		
	Code	Code	Code	Code	Code	Code	Code	Code	code	code	code	code	code	code	
BioPro KS	LBKS	5"	53	0.1µm	36	Single Step ½" hose barb	Q	Yes	R	Inline	X	Non Sterile	1	1	01
		10"	54	0.2µm	01	1½" Sanitary Flange	E	No**	X	T-Line***	T	Gamma Sterile	3		
		20"	55	0.45µm	02	3/8" Hose Barb	I								
		30"	56			1" Hose Barb*	Z								

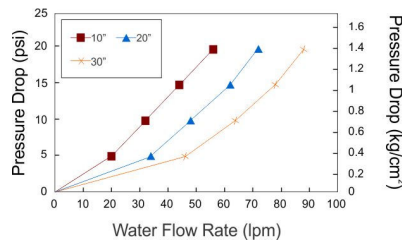
Example :

LBKS	56	01	QQ	R	X	1	01
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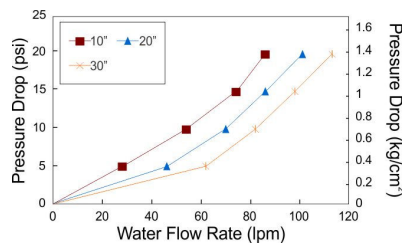


Typical Water Flow Rates

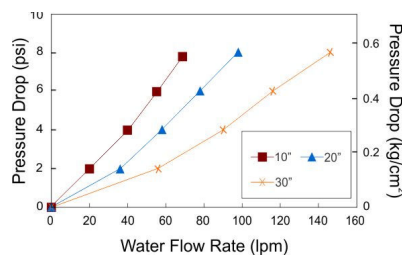
0.1µm BioPro KS-y



0.2µm BioPro KS-y



0.45µm BioPro KS-y



Specifications

Integrity Test (Bubble Point) Specifications (water wetted)

0.1µm: ≥ 40psi, 0.2µm: ≥ 30psi

Bacterial Retention

0.1µm: LRV> 6 for *B.diminuta* ATCC 19146 per cm² of filter area

0.2µm: LRV> 5 for *B.diminuta* ATCC 19146 per cm² of filter area

Material of Construction

Housing – Polypropylene

Filter – Polyethersulfone

Drainage Layer-Polyester

Maximum Differential Pressure

≤ 4 Kg/cm² @ 30° C

Maximum Operating Temperature

80° C @ < 2 Kg/cm²

Sterilization

Irradiation	Gamma irradiatable upto 50 kGy
Autoclave	Autoclavable at 125° C for 30 minutes, 1 cycle after gamma irradiation. Cannot be in-line steam sterilized.

Oxidizable Matter

Passes test as per USP <1231>

Extractables

Passes NVR test as per USP <661>

Bacterial Endotoxin

Aqueous extracts exhibit < 0.25 EU/ml as established by LAL Test as per USP <85>

Fiber Release:

Complies with USFDA CFR Title 21, 210.3 (b) (6)

Particle Release:

The filtrate complies with USP <788> test for particulate matter in injections

Biosafety

Toxicity: Passes Bioreactivity test, *In-vivo*, as per USP <88> for Class VI plastics

Cytotoxicity: Passes Biological Reactivity Tests, In Vitro, USP <87> for Cytotoxicity

Indirect Food Additives:

Passes as per USFDA 21 CFR 177.1520