

3M™ Emphaze™ AEX Hybrid Purifier



One step to increase efficiency

The 3M™ Emphaze™ AEX Hybrid Purifier is a synthetic, single-use chromatographic purifier used for biopharmaceutical clarification. It delivers consistent, high-purity clarified process fluid by reducing negatively charged DNA, HCP, endotoxin, and cell debris through a chromatographic mechanism.

Integrating 3M Emphaze AEX Hybrid Purifier in the clarification stage can improve product purity and yield early in the process and may reduce the total cost of ownership for the mAb purification process.

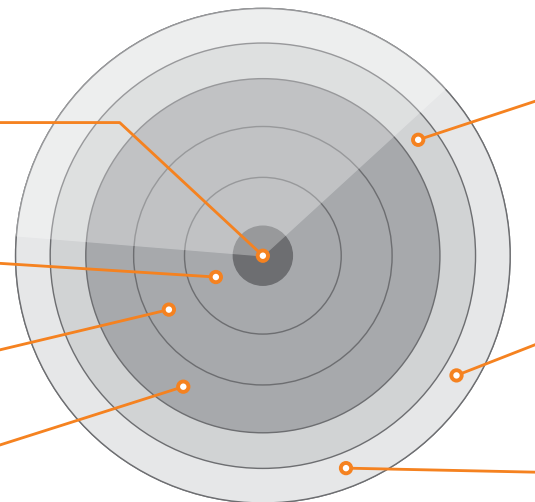
One product performing multiple actions

DNA reduction
4 logs

HCP reduction
20-40%

Sanitizable
with **NaOH**

Turbidity reduction
<100 → 5 NTU



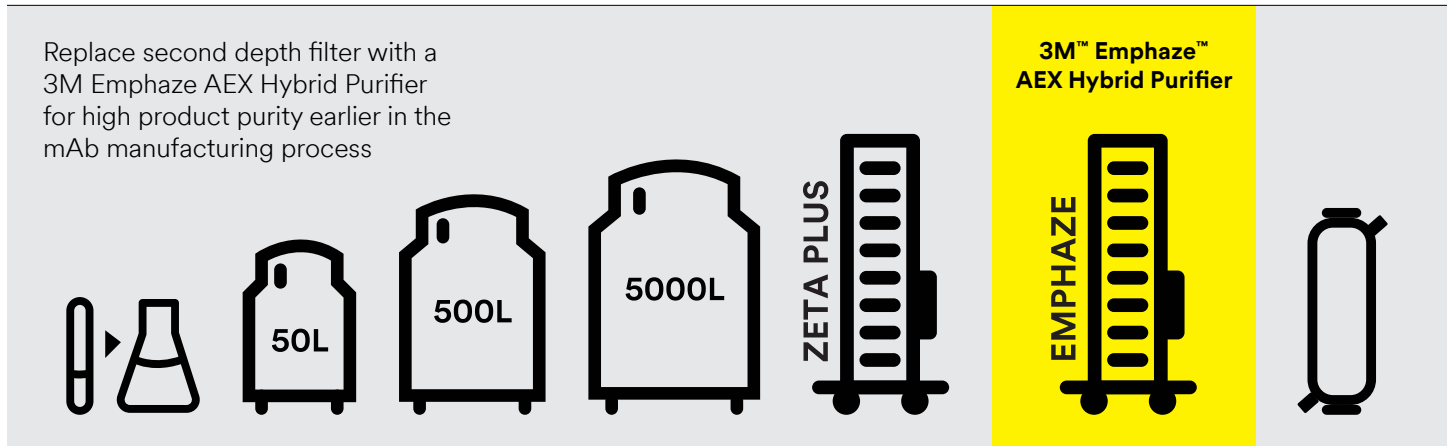
Protein A Column Protection by reduced fouling and shorter regeneration times

Excellent mAb Recovery
> 95% mAb

Sterilizable by
Autoclaving

3M™ Emphaze™ AEX Hybrid Purifier for Monoclonal Antibody Processing

In a representative monoclonal antibody (mAb) manufacturing process, when used in combination at the clarification stage with a depth filter and membrane filters, the 3M Emphaze AEX Hybrid Purifier increases process efficiency and protein purity post Protein A.



Impact of 3M™ Emphaze™ AEX Hybrid Purifier on Clarification step and Protein A performance

- ▶ 10X less HCP post-Protein A
- ▶ 1,000X DNA reduction post-Protein A
- ▶ 10X less residual contaminants on Protein A column
- ▶ Significant reduction in filter surface area of sterilizing filters
- ▶ Consistent output turbidity (<5 NTU)
- ▶ Nominal 20-40% HCP reduction pre-Protein A

Figure 1: HCP concentrations in CHO harvest or centrate (1), clarified fluid (2), protein A eluate (3), and clean-in-place fluid (4) when CHO harvest/centrate was clarified using either 3M Depth Filtration Clarification (grey) or 3M™ Emphaze™ AEX Hybrid Purifier (orange).

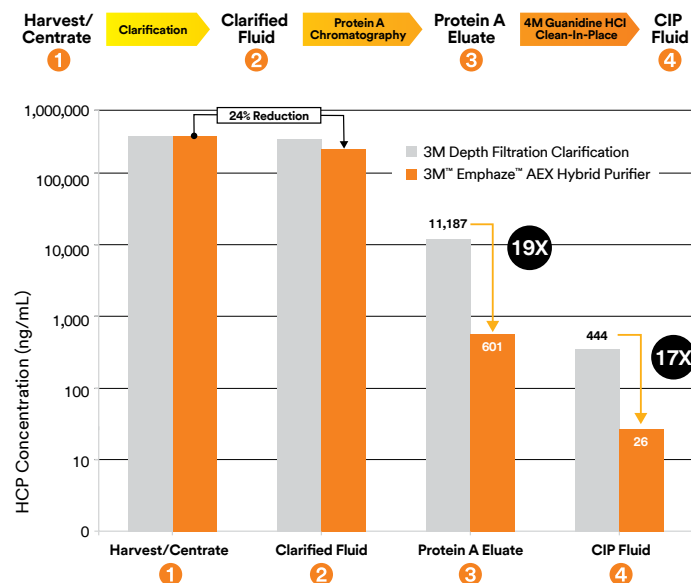
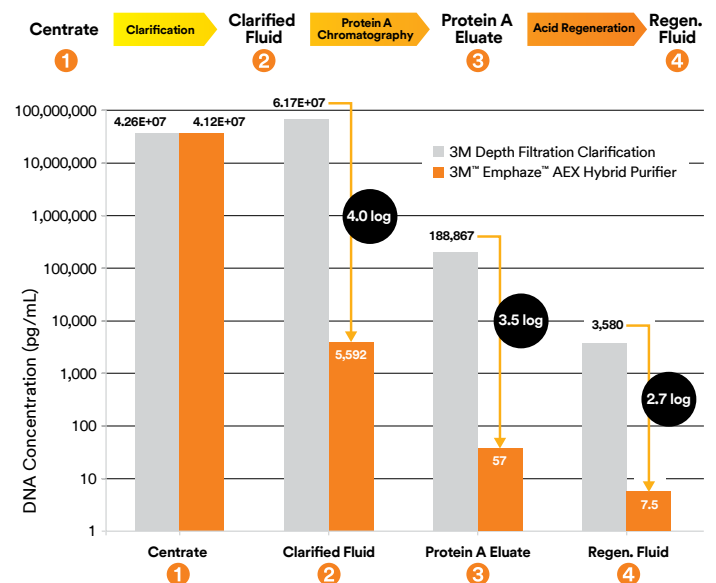


Figure 2: DNA concentrations in CHO centrate (1), clarified fluid (2), protein A eluate (3), and acid regeneration fluid (4) when CHO centrate was clarified using either 3M Depth Filtration Clarification (grey) or 3M™ Emphaze™ AEX Hybrid Purifier (orange).



Reference: 3M Tech Note 01 - Performance of 3M™ Emphaze™ AEX Hybrid Purifier in monoclonal antibody (mAb) purification process.

Excellent contaminant reduction coupled with Protein A chromatography

Figure 3: HCP concentration in Protein A eluate over multiple chromatography cycles

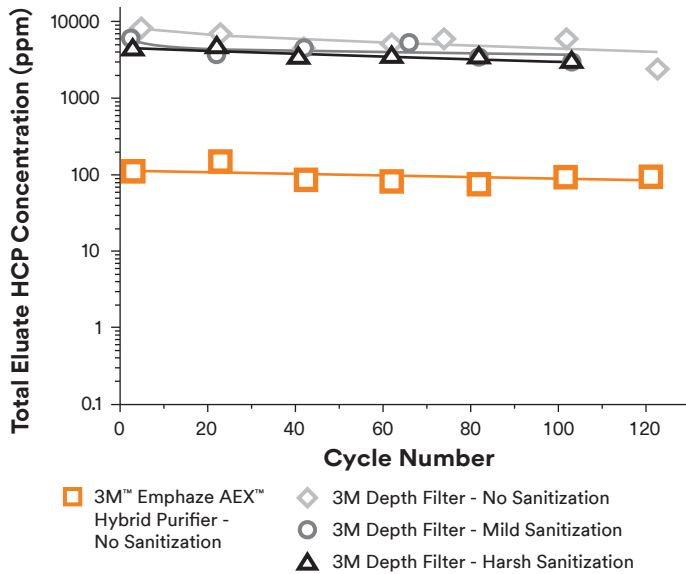
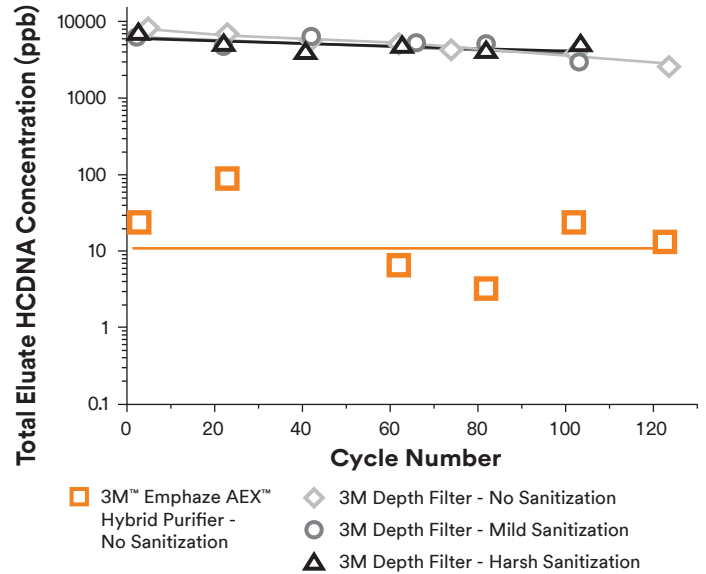
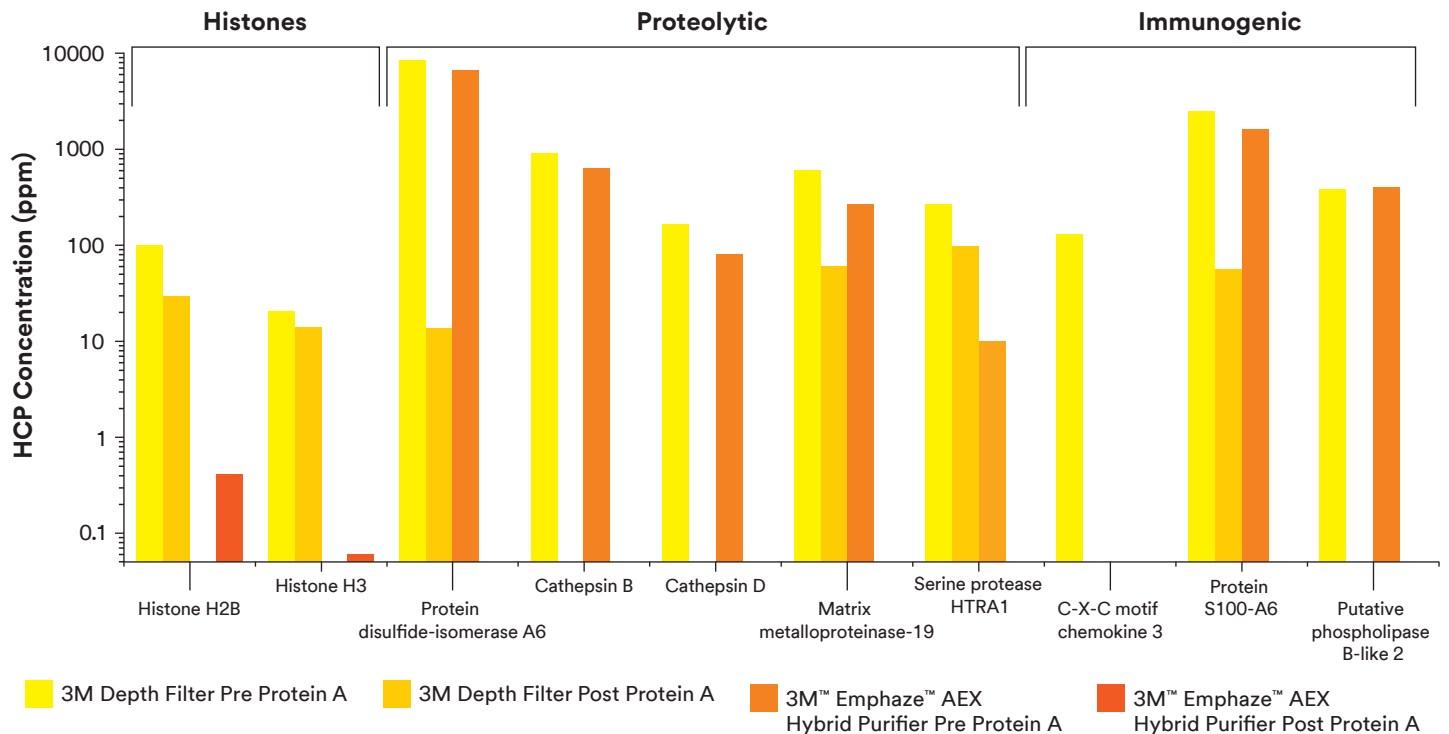


Figure 4: DNA concentration in Protein A eluate over multiple chromatography cycles



Problematic host cell protein reduction pre- and post-Protein A

Figure 5: Removal of selected problematic HCPs during primary clarification through Protein A chromatography



Reference: Modified from Gilgunn et al. 2019. Identification and tracking of problematic host cell proteins removed by synthetic, highly functionalized nonwoven media in downstream bioprocessing of monoclonal antibodies. Journal of Chromatography A. DOI: 10.1016/j.chroma.2019.02.056

Bacterial Lysate Processing

In a representative bacterial lysate process, a decrease of soluble contaminants and increase of process robustness may be achieved by using 3M™ Emphaze™ AEX Hybrid Purifier.

High Performance Clarification Process

- ▶ Endotoxin reduction
- ▶ DNA reduction
- ▶ HCP reduction

Impact Downstream of Clarification

- ▶ Reduced impurities may lead to chromatography column protection (e.g. IMAC, CEX)

Figure 6: Representative process for clarification and purification of products from bacterial lysate.

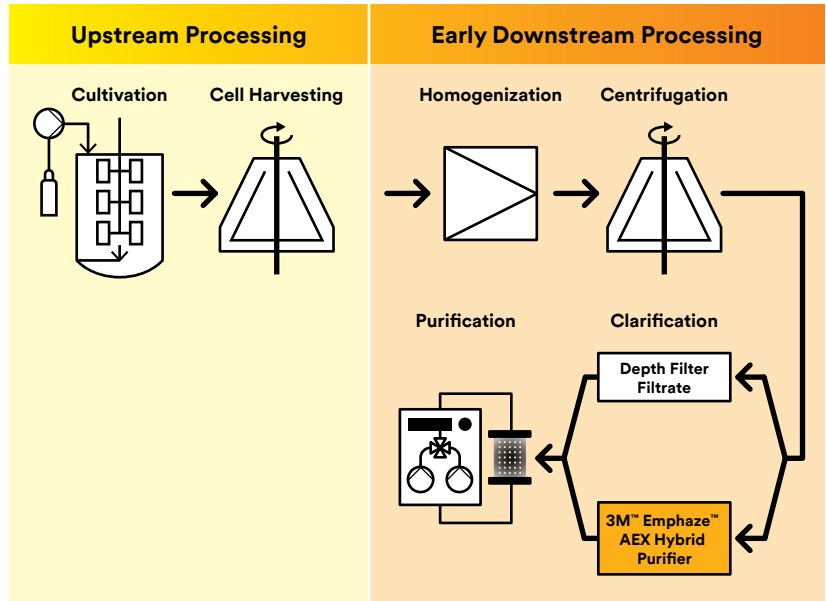
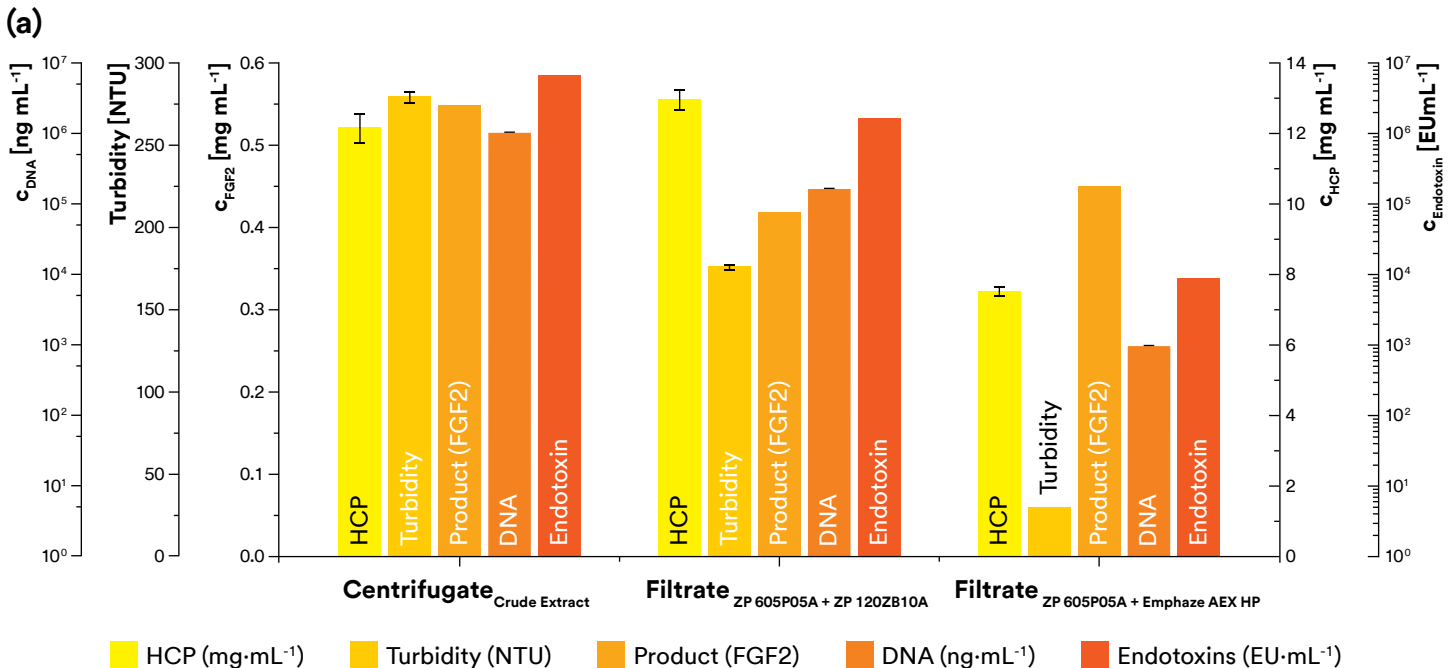


Figure 7: Proof-of-concept: comparison of a conventional depth filtration and an adsorptive filtration. (a) Analyte/impurity concentration comparison of the crude pool against depth filter filtrate and an adsorptively clarified filtrate.



Reference: Modified from Metzger et al. 2019. Adsorptive filtration: A case study for early impurity reduction in an *Escherichia coli* production process. Biotechnology Progress. DOI: 10.1002/btpr.2948

Laboratory, pilot, and production capsules

Product Selection/Specification

(NOTE: R after product and model name indicates the sterilization/sanitization compatible products)



| Product Name | BV0.3R | BV1R | BV8R | | BV60R | BV120R | BV360R | BV800R | BV5600R | |
|--|---|--|---|---------------------------------|--|---------------|-----------------------------------|---|------------------------------------|---------------|
| Model Name | EMP101AEX020R | EMP201AEX020R | EMP301AEX020R | EMP303AEX020R | EMP503AEX020R | EMP513AEX020R | EMP533AEX020R | EMP710AEX020R | EMP770AEX020R | |
| Height x Diameter | 4.8 × 4.3 cm (1.9 × 1.7 in.) | 5.9 × 4.3 cm (2.3 × 1.7 in.) | 4.5 × 7.7 cm (1.7 × 3.0 in.) | 8.8 × 7.7 cm (3.5 × 3.0 in.) | 10.3 × 21.6 cm (4.1 × 6.5 in.) | | 15.2 × 21.6 cm (6.0 × 6.5 in.) | 5.7 × 45.2 cm (2.2 × 17.8 in.) | 20.3 × 45.2 cm (8.0 × 17.8 in.) | |
| Dry Weight | 9.0 g | 14.5 g | 71 g | 77 g | 1.0 kg | 1.1 kg | 1.6 kg | 3.4 kg | 9.5 kg | |
| Weight Wet Post Blow Down | 9.5 g | 16.0 g | 80 g | 85 g | 1.1 kg | 1.2 kg | 2.1 kg | 4.1 kg | 14.2 kg | |
| Fill Volume¹ | 2.0 mL | 5.5 mL | 13 mL | 16 mL | 0.55 L | 0.55 L | 1.4 L | 3.4 L | 10.6 L | |
| Hold up Volume Post Blow Down² | 0.5 mL | 1.5 mL | 9 mL | 9 mL | 0.10 L | 0.15 L | 0.46 L | 0.70 L | 4.7 L | |
| Capsule Material | Polypropylene | | Polypropylene, Glass Filled Polypropylene | | Polypropylene, Glass Filled Polypropylene, Polysulfone, Fluorocarbon | | | Polypropylene, Glass Filled Polypropylene, Glass Filled Polyphenylene Oxide/Polystyrene, Silicone | | |
| Autoclave Cycle | Sterilization Pre-Use | 121°C, 30 min | 121°C, 30 min | 121°C, 30 min | | 121°C, 30 min | 121°C, 30 min | 121°C, 40 min | 121°C, 40 min | 121°C, 40 min |
| | Sterilization Post Use | 121°C, 40 min | 121°C, 40 min | 121°C, 40 min | | 121°C, 40 min | 121°C, 40 min | 121°C, 40 min | 121°C, 40 min | 121°C, 40 min |
| Alkaline Resistance | Pre-Use | 1M NaOH soak for 1 hour at ambient temperatures followed by gravity drain to remove excess base; DO NOT BLOW DOWN | | | | | | | | |
| | Post Use | Capsule soak for 1 hour with 1M NaOH or 5% NaClO (bleach) | | | | | | | | |
| Inlet/Outlet Connections | Luer-Lok | | | 3/4 in. Sanitary Connectors | | | | 1-1/2 in. Sanitary Connectors | | |
| Maximum Inlet Pressure³ | 3.4 bar | | 2.8 bar | | 3.1 bar | | | 3.4 bar | | |
| Maximum Differential Pressure | 2.4 bar | | 2.4 bar | | 2.4 bar | 2.4 bar | 2.4 bar | 2.4 bar | 2.4 bar | |
| Maximum Temperature | 40°C (104°F) | 40°C (104°F) | 40°C (104°F) | 40°C (104°F) | 40°C (104°F) | 40°C (104°F) | 40°C (104°F) | 40°C (104°F) | 40°C (104°F) | |
| Required Preconditioning Flush Volume⁴ | 6.5 mL | 22 mL | 130 mL | | 0.9 L | 1.8 L | 5.5 L | 12 L | 85 L | |
| Recommended Use Flow Rate | 0.4 mL/min | 1.4 mL/min | 8 mL/min | | 50 mL/min | 100 mL/min | 300 mL/min | 680 mL/min | 4700 mL/min | |
| Storage Conditions | Controlled indoor temperatures: 0–30°C (32–86°F) in original sealed packaging | | | | | | | | | |
| Shelf Life | Up to 2 years from date of manufacture @ 30°C maximum storage | | | | | | | | | |

A full support package is available for the 3M™ Emphaze™ AEX Hybrid Purifier. This package includes Installation and Operation Instructions, Certificate of Quality or Certificate of Lot Conformance, and a Regulatory Support File.

1. Capsule Fill Volume is defined as the volume of liquid that is required to fill the capsule.
2. Post Blow-Down Hold-Up Volume is defined as the volume of the residual liquid after air/gas blow down.
3. Do not use this product for continuous service with compressed gasses. The use of compressed gas is permissible for integrity testing and blow down purposes.
4. A Preconditioning Flush is required for the product to be compliant with USP Biological Reactivity Tests, including USP <87> and <88> Class VI 70C minimum. The flush solution can be a buffer or 25–150mM sodium chloride solution. Refer to Installation and Operation Instructions for complete instructions on how to perform the preconditioning flush.

Additional References and Literature

- ▶ 3M Tech Note 01 - Performance of 3M™ Emphaze™ AEX Hybrid Purifier in monoclonal antibody (mAb) process in mammalian cell cultures
- ▶ 3M Application Note 02 - Endotoxin removal by 3M™ Emphaze™ AEX Hybrid Purifier
- ▶ 3M Tech Note 01 - Identification and tracking of problematic host cell proteins when using 3M™ Emphaze™ AEX Hybrid Purifier in mAb manufacturing processes
- ▶ 3M Application Note 01 - Quantification of 3M™ Emphaze™ AEX Hybrid Purifier value in mAb manufacturing process

- ▶ Maurer et al. 2014. Advanced clarification of cell culture supernatant by 3M™ Emphaze™ AEX Hybrid Purifier for fast and economic bioprocessing of recombinant proteins. DOI: 10.1016/j.nbt.2014.05.1897
- ▶ Singh et al. 2017. Development of adsorptive hybrid filters to enable two-step purification of biologics. MAbs. DOI: 10.1080/19420862.2016.1267091
- ▶ El-Sabbahy et al. 2018. The effect of feed quality due to clarification strategy on the design and performance of protein A periodic counter-current chromatography. Biotechnology Progress. DOI: 10.1002/btpr.2709
- ▶ Koehler et al. 2019. Enhancing Protein A performance in mAb processing: A method to reduce and rapidly evaluate host cell DNA levels during primary clarification. Biotechnology Progress. DOI: 10.1002/btpr.2882
- ▶ Gilgunn et al. 2019. Identification and tracking of problematic host cell proteins removed by a synthetic, highly functionalized nonwoven media in downstream bioprocessing of monoclonal antibodies. Journal of Chromatography A. DOI: 10.1016/j.chroma.2019.02.056
- ▶ Metzger et al. 2019. Adsorptive filtration: A case study for early impurity reduction in an Escherichia coli production process. Biotechnology Progress. DOI: 10.1002/btpr.2948
- ▶ Pinto et al. 2020. Impact of micro and macroporous TFF membranes on product sieving and chromatography loading for perfusion cell culture. Biotechnol Bioeng. DOI: 10.1002/bit.27192
- ▶ Van de Velde et al. 2020. Chromatographic clarification overcomes chromatin-mediated hitch-hiking interactions on Protein A capture column. Biotech Bioeng. DOI: 10.1002/bit.27513
- ▶ Metzger et al. 2020. IGF1 inclusion bodies: A QbD based process approach for efficient USP as well as early DSP unit operations. Journal of Biotechnology. DOI: 10.1016/j.jbiotec.2020.02.014
- ▶ Meingast et al. 2021. Physicochemical properties of enveloped viruses and arginine dictate inactivation. Biotechnology Journal. DOI: 10.1002/biot.202000342
- ▶ Holstein et al. 2021. Control of leached beta-glucan levels from depth filters by an improved depth filtration flush strategy. Biotechnology Progress. DOI: 10.1002/btpr.3086

For more information about the 3M™ Emphaze™ AEX Hybrid Purifier, contact your local sales rep by calling 1-800-243-6894, option 4, or visiting us at 3M.com/Emphaze

Intended Use Statement: Single-use filter products are intended for use in biopharmaceutical processing applications of aqueous based pharmaceuticals (drugs) and vaccines in accordance with the product instructions and specifications, and cGMP requirements, where applicable.

Since there are many factors that can affect a product's use, the customer and user remain responsible for determining whether the 3M product is suitable and appropriate for the user's specific application, including user conducting an appropriate risk assessment and evaluating the 3M product in user's application.

Restrictions on Use: 3M advises against the use of these 3M products in any application other than the stated intended use(s), since other applications have not been evaluated by 3M and may result in an unsafe or unintended condition. Do not use in any manner whereby the 3M product, or any leachable from the 3M product, may become part of or remains in a medical device that is regulated by any agency, and/or globally exemplary agencies, including but not limited to: a) FDA, b) European Medical Device Regulation (MDR), c) Japan Pharmaceuticals and Medical Devices Agency (PMDA) or in applications involving permanent implantation into the body; Life-sustaining medical applications; Applications requiring food contact compliance.

Product Selection and Use: Many factors beyond 3M's control and uniquely within user's knowledge and control can affect the use and performance of a 3M product in a particular application. As a result, end-user is solely responsible for evaluating the product and determining whether it is appropriate and suitable for end-user's application, including completing a risk assessment that considers the product leachable characteristics and its impact on drug safety, conducting a workplace hazard assessment and reviewing all applicable regulations and standards. Failure to properly evaluate, select, and use a 3M product and appropriate safety products, or to meet all applicable safety regulations, may result in injury, sickness, death, and/or harm to property.

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