



## PureFlo® PE Capsules

PureFlo® PE capsules are gamma-irradiatable, easy-to-use filtration units, ideally suited for integration into single-use sterile assemblies. The hydrophobic polyethylene (PE) membrane demonstrates exceptional flow performance in venting and compressed gas applications.

### Superior Flow Performance

PureFlo® PE capsules incorporate a sterilizing grade 0.2 µm membrane that exhibits a lower pressure drop compared to other hydrophobic membranes such as PVDF. The superior flow performance results in a light-weight, compact filter designed to promote optimal process efficiency.

### Engineered Filtration Solutions

To meet the needs of an innovative and rapidly growing industry, and in addition to the standard product offering, Saint-Gobain offers a collaborative service with a dedicated team of engineers and subject matter experts to develop specialized filtration solutions that are tailored to achieve optimal performance. Please contact us for more information.

### Features / Benefits



Inherently  
Hydrophobic



Gamma-  
Irradiatable



High Flow  
Rate



EMA/410/01 Rev. 3  
Compliant



Sterilizing  
Grade



100% Integrity  
Tested

### Typical Applications

- Bioreactor Vent
- Moisture Barrier
- Compressed Gas
- Fermentation Tank Vent
- Bottle or Carboy Vent

# Technical Summary

| Materials                 |                   |
|---------------------------|-------------------|
| Membrane                  | Polyethylene (PE) |
| Support Material          | Polyethylene (PE) |
| Molded Components (shell) | Polyethylene (PE) |
| O-rings (when present)    | Silicone          |

| Membrane Configurations |             |                                                          |
|-------------------------|-------------|----------------------------------------------------------|
| Membrane                | Description | Minimum Bubble Point (at 22 °C)                          |
| UEO20                   | 0.2 µm      | 1.2 bar   17.4 psi (in 60% isopropyl alcohol, 40% water) |

| Measurements (nominal) |                       |                                   |                       |
|------------------------|-----------------------|-----------------------------------|-----------------------|
| Size                   | Outside Diameter (mm) | Body Length Without Fittings (mm) | Filtration Area (cm²) |
| Junior (JKP)           | 41                    | 4.1                               | 130                   |
| Mini (MKP)             | 59                    | 60                                | 400                   |
| Mid (SKV)              | 71.5                  | 52 - 257                          | 550 - 3800            |
| Full (L)               | 90                    | 279 - 1007                        | 5300 - 21000          |

| Recommended Operating Conditions (for untreated/non-sterile product) |                     |
|----------------------------------------------------------------------|---------------------|
| Maximum Temperature                                                  | 60 °C               |
| Maximum Inlet Pressure (at 22 °C)                                    | 4.1 bar   59.5 psi  |
| Maximum Forward Differential Pressure (at 22 °C)                     | 5 bar   72.5 psi    |
| Maximum Reverse Differential Pressure (at 22 °C)                     | 2.1 bar   30.45 psi |

| Sterilization  |                         |
|----------------|-------------------------|
| Gamma          | Up to 50 kGy            |
| Ethylene Oxide | Testing is recommended. |

## Shelf Life

Non-sterile products have a shelf life of 3 years after the date of manufacture.

## Traceability

For traceability and easy identification:

- A Certificate of Conformance is provided in each box.
- Each filter bag and box are labeled with the product part number, lot number, and identifying characteristics.
- Each Junior, Mini, Mid, and Full size capsule is engraved and labeled with the product part number, lot number, and identifying characteristics.

## Regulatory & Stewardship

Saint-Gobain's PureFlo® PE Junior, Mini, Mid and Full Size Capsules were designed, manufactured, and qualified with an ISO 13485 certified Quality Management System. For more information, please refer to the product Validation Guide and Regulatory Information Overview (RIO).

| Category                  | Standard or Reference Test                                                                                         |
|---------------------------|--------------------------------------------------------------------------------------------------------------------|
| Cleanliness               | USP <85> Bacterial Endotoxin Test                                                                                  |
| Biocompatibility          | USP <88> Class VI, and/or USP <87>, and/or ISO 10993-5<br>USP 21 CFR 177 FDA Indirect Food Additive                |
| Bacterial Retention       | Quantitative retention of 10 <sup>7</sup> CFU/cm <sup>2</sup> <i>Brevundimonas diminuta</i> per ASTM methodology   |
| Animal-derived Components | EMA/410/01 Rev. 3                                                                                                  |
| RoHS                      | Restriction of Hazardous Substances (RoHS 3) Directive 2015/863                                                    |
| REACH                     | REACH (Registration, Evaluation, Authorisation and Restriction of Chemicals) Article 57, Regulation No. 1907/2006) |

## Lot Release Criteria

Each PureFlo® PE Junior, Mini, Mid and Full Size Capsules product lot is sampled and tested for conformity to the following criteria:

| Criteria                 | Description                                                                                                                                                                        |
|--------------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Integrity (membrane)     | Each filter (100%) in a lot is integrity tested during the manufacturing process.                                                                                                  |
| Integrity (housing)      | A representative sample of the lot is tested to meet published burst pressure values.                                                                                              |
| USP Bacterial Endotoxins | A representative sample from the lot is tested to confirm that an aqueous extraction of the product contains <0.25 EU/mL as determined by the Limulus Amebocyte Lysate (LAL) Test. |

# PureFlo® PE

## Junior Capsules

Saint-Gobain  
Life Sciences –  
Bioprocess Solutions



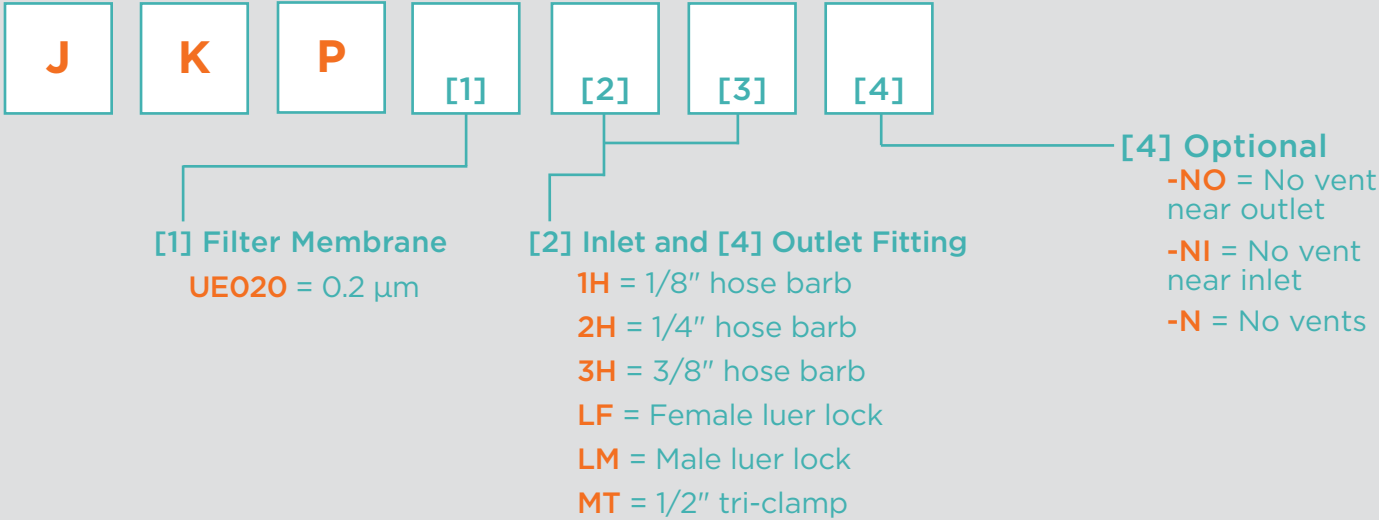
**PRODUCT TECH SNAPSHOT:** (Values are nominal. Please see Technical Summary for additional information)

| Body Diameter (mm) | Body Length Without Fittings (mm) | Filtration Area (cm <sup>2</sup> ) | Typical Flow Performance*        |
|--------------------|-----------------------------------|------------------------------------|----------------------------------|
| 41                 | 41                                | 130                                | JKPUE0202H2H – 0.2 psid at 4 LPM |

1 bar =14.5038 psi

## Ordering Guide

PART NUMBER:



### Customer Service

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Gaithersburg, MD 20878  
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Don't see what you're looking for?  
Please let us know! We're here to help.

\* IMPORTANT: The flow performance of a filter capsule is dependent on several variables including the process fluid characteristics, operating conditions, filter surface area, pore size and fittings. The above values are approximate for example part numbers and provided for reference only. Actual values may vary. For application guidance and flow performance information for a specific part number please contact your local Saint-Gobain representative.

# PureFlo® PES Capsules

Saint-Gobain  
Life Sciences –  
Bioprocess Solutions



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**WARRANTY:** For a period of 12 months from the date of first sale, Saint-Gobain Life Sciences warrants this product to be free of defects in materials and workmanship. Our only obligation will be to replace any portion proving defective, or at our option, to refund the purchase price thereof.

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