

Supplemental Figure 1. EX-PRESS minishunt, Model P-50. Image courtesy of Fort Worth, Alcon Laboratories, Inc. Used with written approval from Alcon.





EXPRESS®

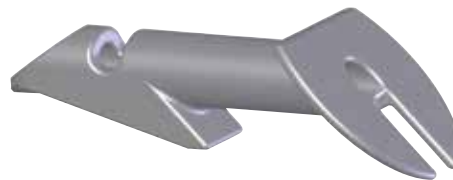
Glaucoma Filtration Device

EX-PRESS® Glaucoma Filtration Device Ordering Information

EX-PRESS® Glaucoma Filtration Device P Model

Vertical Channel for Posterior Flow

| | |
|---------------------|-----------------------|
| External lumen | 400µm |
| Internal lumen size | 50µm |
| Device length | 2.64 mm |
| Tip shape | Decreased bevel angle |
| Back plate shape | Vertical channel |
| SKU | P-50 PL-24053 |



Preloaded and
Ready to Use

To order, please contact
Alcon customer service at
1-800-TO-ALCON
(1-800-862-5266)

CAUTION: Federal (USA) law restricts this device to sale by, or on the order of, a physician.

INDICATION: The EX-PRESS® Glaucoma Filtration Device is intended to reduce intraocular pressure in glaucoma patients where medical and conventional surgical treatments have failed.

GUIDANCE REGARDING THE SELECTION OF THE

APPROPRIATE VERSION: Prior clinical studies were not designed to compare between the various versions of the EX-PRESS® Glaucoma Filtration Device.

The selection of the appropriate version is according to the doctor's discretion.

CONTRAINDICATIONS: The use of this device is contraindicated if one or more of the following conditions exist:

- Presence of ocular disease such as uveitis, ocular infection, severe dry eye, severe blepharitis.
- Pre-existing ocular or systemic pathology that, in the opinion of the surgeon, is likely to cause postoperative complications following implantation of the device.
- Patients diagnosed with angle closure glaucoma.

WARNINGS/PRECAUTIONS:

- The surgeon should be familiar with the instructions for use.
- The integrity of the package should be examined prior to use

and the device should not be used if the package is damaged and sterility is compromised.

- This device is for single use only.
- MRI of the head is permitted, however not recommended, in the first two weeks post implantation.

ATTENTION: Reference the Directions for Use labeling for a complete listing of indications, warnings, precautions, complications and adverse events.