

THIS IS UTILITY

This is

vektor™

ARTICULATING ILLUMINATED LASER PROBE

Allows surgeon flexibility with adjustable articulation¹

Enables unassisted scleral depression¹

Expands surgical range to help avoid the lens when reaching to the far side of the periphery¹

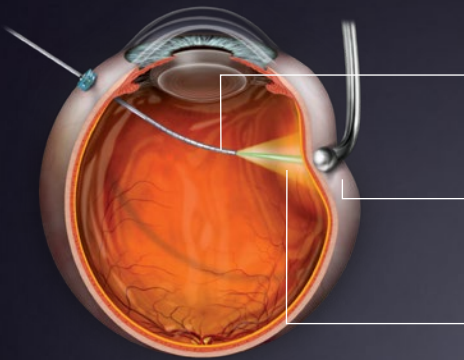


THIS IS UTILITY



Engineered for Versatility

Provides more functionality and control for photocoagulation in the retinal periphery²:



Continuously adjustable tip angle enables greater surgeon control

Design enables unassisted scleral depression

Illumination assists surgical visualization



Engineered for Integration With The CONSTELLATION® Vision System



HOW TO ORDER

To learn more about the VEKTOR™ Articulating Illuminated Laser Probe from Alcon, contact your local sales representative.

Product	Description
8065752556	27GA Articulating Illuminated Laser Probe w/RFID
8065752555	25GA Articulating Illuminated Laser Probe w/RFID
8065752554	23GA Articulating Illuminated Laser Probe w/RFID

PUREPOINT® LASER IMPORTANT PRODUCT INFORMATION

Caution: Federal (USA) law restricts this device to sale by, or on the order of, a physician. **Indications for Use:** The PUREPOINT® Laser is indicated for use in photocoagulation of both anterior and posterior segments of the eye including: • Retinal photocoagulation, panretinal photocoagulation and intravitreal endophotocoagulation of vascular and structural abnormalities of the retina and choroid including: Proliferative and nonproliferative retinopathy (including diabetic); choroidal neovascularization secondary to age-related macular degeneration; retinal tears and detachments; macular edema, retinopathy of prematurity; choroidal neovascularization; leaking microaneurysms. • Iridotomy/Iridectomy for treatment of chronic/primary open angle glaucoma, acute angle closure glaucoma and refractory glaucoma. • Trabeculoplasty for treatment of chronic/primary open angle glaucoma and refractory glaucoma. • And other laser treatments including: internal sclerostomy; lattice degeneration; central and branch retinal vein occlusion; suturelysis; vascular and pigment skin lesions. **Contraindications:** Patients with a condition that prevents visualization of target tissue (cloudy cornea, or extreme haze of the aqueous humor of the anterior chamber of vitreous humor) are poor candidates for LIO delivered laser treatments. **Warnings and Precautions:** • The disposables used in conjunction with ALCON® instrument products constitute a complete surgical system. Use of disposables and handpieces other than those manufactured by Alcon may affect system performance and create potential hazards. • Attach only Alcon supplied consumables to console and cassette luer fittings. Do not connect consumables to the patient's intravenous connections. • Mismatch of consumable components and use of settings not specifically adjusted for a particular combination of consumable components may create a patient hazard. • Back scattered radiation is of low intensity and is not harmful when viewed through a protective filter. All personnel in the treatment room must wear protective eyewear, OD4 or above at 532nm, when the system is in Standby/Ready mode as well as during treatment. The doctor protection filter is an OD greater than 4 at 532nm. **Attention:** Reference the Directions for Use for a complete listing of indications, warnings, and precautions.

References: 1. Alcon data on file. Alcon Laboratories, Inc; February 2017. 2. Auld JR, Farley MH, inventors; Alcon Research Ltd, assignee. Multi-fiber flexible surgical probe. US published patent CA 2787024 A1. August 25, 2011.