Online Resource 3 Eligibility criteria

Key Inclusion Criteria

 \geq 18 years of age

Diagnosis of OAG in the study eye, with >1 dB and <17 dB of mean deviation visual field loss on Humphrey perimetry (24-2 full threshold or SITA–standard program)

Diagnosis of OAG or ocular hypertension in the fellow eye that, in the investigator's opinion, could be treated adequately with topical bimatoprost 0.03% monotherapy

History of a minimum of approximately 20% IOP lowering in response to a topical PGA ocular hypotensive medication

Iridocorneal angle in the study eye ≥ 3 inferiorly (Shaffer grade), based on clinical gonioscopy, and of sufficient size to fit an implant without corneal endothelial cell touch on optical coherence tomography, as determined by an independent reading center

Post-washout IOP between 22 and 36 mm Hg (inclusively) in both eyes at hour 0 (8 AM \pm 1 hour) at the baseline visit

Key Exclusion Criteria

History of posterior capsule tear during cataract surgery

Any ocular surface finding with severity greater than trace on biomicroscopic examination at baseline

History of conjunctival hyperemia with severity greater than mild, due to bimatoprost or other PGA use History of narrow-angle or closed-angle glaucoma

Central corneal thickness <470 μ m or >630 μ m, or a difference between eyes >70 μ m

Central corneal endothelial cell count <2000 cells/mm² by specular microscopy

IOP intraocular pressure, *OAG* open-angle glaucoma, *PGA* prostaglandin analog, *SITA* Swedish interactive thresholding algorithm