

BRANCH RETINAL VEIN OCCLUSION STUDY

STUDY ARMS:

- 1. Standard therapy (Lucentis)**
- 2. Study therapy (peripheral laser + Lucentis)**

Study duration: 12 months

Inclusion Criteria

Subjects will be eligible if the following criteria are met:

- Ability to provide written informed consent and comply with study assessments for the full duration of the study**
- Age ≥ 18 years**

Patient-related considerations:

- Patients with branch retinal vein occlusion**

Disease-related considerations:

- Study eye with macular edema with no previous laser treatment and no pharmacologic injection in last 3 months.**
- Study eye with best corrected visual acuity between 20/40 (≤ 73 letters ETDRS chart) and 20/320 (≥ 19 letters on ETDRS chart)**

Other considerations:

- Patients able to complete all study visits**
- Female patients must be using two forms of contraception**