Summary GLC 04-10

Comparison of the incidence and severity of conjunctival hyperemia associated with topical use of bimatoprost 0.01% and latanoprost 0.005% in glaucoma or ocular hypertensive patients

The study was approved on May 27, 2010, the first patient was enrolled on June 01, 2010 and was terminated early on September 15, 2011.

We stopped the study early due to difficulties in enrolling patients willing to submit to study procedures; 14 patients were enrolled in the study.
The consequences of early termination for the evaluation of the results of this study only marginally impacted on the evaluation of the risks and benefits of the investigational drug since there are numerous reports that have evaluated hyperemia as well as the effectiveness of the treatment.

VIA LIVENZA, 3 – 00198 ROMA
P.I. 01891741009 – COD. FISC. 07864670588 – UFFICIO TERRITORIALE GOVERNO DI ROMA, REG. PERS. GIUR. N. 2148

TEL. +39/068552055 – 0685356727 – FAX 0684242333
e-mail: info@fondazionebietti.it https://www.fondazionebietti.it/