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Novartis Pharma AG
CH-4002, Basel, Switzerland

Reference: EudraCT 2011-004160-30/ Novartis Protocol ID CFTY720D2205

A multicenter, randomized, active-controlled study to assess the safety, tolerability, and efficacy of FTY720 in patients with acute, noninfectious intermediate, posterior and pan uveitis

Trial CFTY720D2205 was cancelled with no patient enrollment and as such, no results will be reported.