FDA 510(k) Premarket Notification

K112154 FEMTO LDV

CLEARED

Mar 16, 2012
233 days to decision

K112154 · Product code: **HQF** · Ophthalmic
Source: https://www.510kdatabase.net/k112154/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	Jul 27, 2011
Decision date	Mar 16, 2012
Days to decision	233 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Sie Ag,Surgical Instument Engineering**
Location	Greenwood Village, CO, US
Contact	Kevin Walls
510(k) history	1 submissions · 1 cleared · 2012-2012