FDA 510(k) Premarket Notification

K121738 MYVISIONTRACK(TM)

CLEARED

Feb 22, 2013
254 days to decision

K121738 · Product code: **HOQ** · Ophthalmic
Source: https://www.510kdatabase.net/k121738/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Grid, Amsler (HOQ)
Date received	Jun 13, 2012
Decision date	Feb 22, 2013
Days to decision	254 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Vital Art and Science Incorporated**
Location	Richardson, TX, US
Contact	MICHAEL BARTLETT
510(k) history	2 submissions · 2 cleared · 2013-2015