FDA 510(k) Premarket Notification

K101838 VTS1000

CLEARED

Feb 18, 2011
232 days to decision

K101838 · Product code: **KZA** · General Hospital
Source: https://www.510kdatabase.net/k101838/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Vein Location, Liquid Crystal (KZA)
Date received	Jul 1, 2010
Decision date	Feb 18, 2011
Days to decision	232 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Vuetek Scientific, LLC**
Location	Marietta, GA, US
Contact	PAUL SUMNER
510(k) history	1 submissions · 1 cleared · 2011-2011