FDA 510(k) Premarket Notification

K173381 Nexxis OR

CLEARED

Feb 22, 2018
115 days to decision

K173381 · Product code: **DXJ** · Cardiovascular
Source: https://www.510kdatabase.net/k173381/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Display, Cathode-ray Tube, Medical (DXJ)
Date received	Oct 30, 2017
Decision date	Feb 22, 2018
Days to decision	115 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Barco N.V.**
Location	La Jolla, CA, US
Contact	Eric Caus
Website	http://www.barco.com
510(k) history	60 submissions · 60 cleared · 1998-2024