FDA 510(k) Premarket Notification

K093134 CERNER CAREAWARE IBUS

CLEARED

Nov 27, 2009
53 days to decision

K093134 · Product code: **DRG** · Cardiovascular
Source: https://www.510kdatabase.net/k093134/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG)
Date received	Oct 5, 2009
Decision date	Nov 27, 2009
Days to decision	53 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Cerner Corp.**
Location	Kansas City, MO, US
Contact	SHELLEY S LOOBY
510(k) history	7 submissions · 7 cleared · 1996-2012