FDA 510(k) Premarket Notification

K122259 BLOOD PRESSURE MONITOR WITH STETHOSCOPE

CLEARED

Sep 24, 2012
59 days to decision

K122259 · Product code: **DXQ** · Cardiovascular
Source: https://www.510kdatabase.net/k122259/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Blood Pressure Cuff (DXQ)
Date received	Jul 27, 2012
Decision date	Sep 24, 2012
Days to decision	59 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Hangzhou Reli-On Co., Ltd.**
Location	Echo, OR, US
Contact	CHARILE MACK
510(k) history	1 submissions · 1 cleared · 2012-2012