FDA 510(k) Premarket Notification

CLEARED

K062302 MILLICORE AB DIGIVENT CHEST DRAINAGE SYSTEM

Oct 6, 2006
59 days to decision

K062302 · Product code: **KDQ** · General Hospital
Source: https://www.510kdatabase.net/k062302/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bottle, Collection, Vacuum (KDQ)
Date received	Aug 8, 2006
Decision date	Oct 6, 2006
Days to decision	59 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Millicore AB**
Location	North Attleboro, MA, US
Contact	ROSINA ROBINSON
510(k) history	1 submissions · 1 cleared · 2006-2006

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k062302/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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