FDA 510(k) Premarket Notification

K830780 KOALA C99 I.V. CONTROLLER

CLEARED

Apr 8, 1983
28 days to decision

K830780 · Product code: **LDR** · General Hospital
Source: https://www.510kdatabase.net/k830780/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Controller, Infusion, Intravascular, Electronic (LDR)
Date received	Mar 11, 1983
Decision date	Apr 8, 1983
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Chesebrough-Pond'S U.S.A. Co.**
Location	Mchenry, IL, US
510(k) history	33 submissions · 30 cleared · 1976-1987