FDA 510(k) Premarket Notification

K832907 INVASIVE MONITORING KITS

CLEARED

Jan 10, 1984
134 days to decision

K832907 · Product code: **DRS** · Cardiovascular
Source: https://www.510kdatabase.net/k832907/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Aug 29, 1983
Decision date	Jan 10, 1984
Days to decision	134 days
Third-party review	No

APPLICANT

Company	**American Pharmaseal Div. Ahsc**
Location	Mchenry, IL, US
510(k) history	64 submissions · 61 cleared · 1980-1987