FDA 510(k) Premarket Notification

K902381 PENLON BREATHING SYSTEM PRESSURE MONITOR

CLEARED

Aug 17, 1990
79 days to decision

K902381 · Product code: **CAP** · Toxicology
Source: https://www.510kdatabase.net/k902381/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Airway Pressure (includes Gauge And/or Alarm) (CAP)
Date received	May 30, 1990
Decision date	Aug 17, 1990
Days to decision	79 days
Third-party review	No

APPLICANT

Company	**Penlon, Inc.**
Location	Mchenry, IL, US
Contact	JIM GUNNERSON
510(k) history	10 submissions · 10 cleared · 1980-1994