

**K902381 PENLON BREATHING SYSTEM PRESSURE MONITOR**Aug 17, 1990  
79 days to decisionK902381 · Product code: **CAP** · Toxicology  
Source: <https://www.510kdatabase.net/k902381/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k902381/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 26, 2026