

K812041 VENTI-MONITOR MODEL 101Sep 25, 1981
66 days to decisionK812041 · Product code: **CAP** · AnesthesiologySource: <https://www.510kdatabase.net/k812041/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Airway Pressure (includes Gauge And/or Alarm) (CAP)
Date received	Jul 21, 1981
Decision date	Sep 25, 1981
Days to decision	66 days
Third-party review	No

APPLICANT

Company	Medix Medical Electronics (U.S.A.), Inc.
Location	Mchenry, IL, US
510(k) history	14 submissions · 14 cleared · 1981-1986

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Device record: <https://www.510kdatabase.net/k812041/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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