

K811781 TENSIO METERJul 10, 1981
17 days to decisionK811781 · Product code: **DXN** · CardiovascularSource: <https://www.510kdatabase.net/k811781/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Jun 23, 1981
Decision date	Jul 10, 1981
Days to decision	17 days
Third-party review	No

APPLICANT

Company	Vivitek, Inc.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981

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Device record: <https://www.510kdatabase.net/k811781/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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