

K832171 SELF CHECK DIGITAL BLOOD PRESS. MONITOROct 4, 1983
90 days to decisionK832171 · Product code: **DXN** · CardiovascularSource: <https://www.510kdatabase.net/k832171/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Jul 6, 1983
Decision date	Oct 4, 1983
Days to decision	90 days
Third-party review	No

APPLICANT

Company	Ritter Co.
Location	Mchenry, IL, US
510(k) history	10 submissions · 10 cleared · 1976-1983

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Device record: <https://www.510kdatabase.net/k832171/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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