

K813020 BOMED-NCCOMDec 14, 1981
48 days to decisionK813020 · Product code: **DXG** · Cardiovascular
Source: <https://www.510kdatabase.net/k813020/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Diagnostic, Pre-programmed, Single-function (DXG)
Date received	Oct 27, 1981
Decision date	Dec 14, 1981
Days to decision	48 days
Third-party review	No

APPLICANT

Company	Bomed Medical Mfr., Ltd.
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1981-1986

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Device record: <https://www.510kdatabase.net/k813020/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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