

K802778 LIQUID CRYSTAL THERM. 8 EVENT MONITORJan 9, 1981
65 days to decisionK802778 · Product code: **KPD** · General Hospital
Source: <https://www.510kdatabase.net/k802778/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Strip, Temperature, Forehead, Liquid Crystal (KPD)
Date received	Nov 5, 1980
Decision date	Jan 9, 1981
Days to decision	65 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k802778/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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