

K771539 STEPPEROct 7, 1977
57 days to decisionK771539 · Product code: **KYX** · General Hospital
Source: <https://www.510kdatabase.net/k771539/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dispenser, Liquid Medication (KYX)
Date received	Aug 11, 1977
Decision date	Oct 7, 1977
Days to decision	57 days
Third-party review	No

APPLICANT

Company	Tridak
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1977-1977

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Device record: <https://www.510kdatabase.net/k771539/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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