

**K802245 7 TRICYCLIC ANTIDE PRESSANTS**Oct 10, 1980  
24 days to decisionK802245 · Product code: **LFI** · ToxicologySource: <https://www.510kdatabase.net/k802245/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	High Pressure Liquid Chromatography, Tricyclic Antidepressant Drugs (LFI)
Date received	Sep 16, 1980
Decision date	Oct 10, 1980
Days to decision	24 days
Third-party review	No

**APPLICANT**

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Company	<b>Technicon Instruments Corp.</b>
Location	Mchenry, IL, US
510(k) history	157 submissions · 156 cleared · 1976-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k802245/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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