

**K801259 NEEDLE COUNTER-SURE COUNT/100 COUNT**Jun 9, 1980  
12 days to decisionK801259 · Product code: **MMK** · General Hospital  
Source: <https://www.510kdatabase.net/k801259/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, Sharps (MMK)
Date received	May 28, 1980
Decision date	Jun 9, 1980
Days to decision	12 days
Third-party review	No

**APPLICANT**

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Company	<b>Kleen Test Products</b>
Location	Mchenry, IL, US
510(k) history	17 submissions · 17 cleared · 1976-1989

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Device record: <https://www.510kdatabase.net/k801259/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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