

**K781581 SURGICAL LIGHT 22 SERIES**Oct 17, 1978  
32 days to decisionK781581 · Product code: **FQP** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k781581/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lamp, Operating-room (FQP)
Date received	Sep 15, 1978
Decision date	Oct 17, 1978
Days to decision	32 days
Third-party review	No

**APPLICANT**

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Company	<b>American Sterilizer Co.</b>
Location	Mchenry, IL, US
510(k) history	36 submissions · 35 cleared · 1977-1994

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

Device record: <https://www.510kdatabase.net/k781581/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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