FDA 510(k) Premarket Notification

K781581 SURGICAL LIGHT 22 SERIES

CLEARED

Oct 17, 1978
32 days to decision

K781581 · Product code: **FQP** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k781581/

SUBMISSION DETAILS

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

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