FDA 510(k) Premarket Notification

CLEARED

K883556 LATEX SURGICAL GLOVE (NON STERILE)

Feb 3, 1989
168 days to decision

K883556 · Product code: **KGO** · General Hospital
Source: https://www.510kdatabase.net/k883556/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Surgeon's Gloves (KGO)
Date received	Aug 19, 1988
Decision date	Feb 3, 1989
Days to decision	168 days
Third-party review	No

APPLICANT

Company	**Makintech Corp.**
Location	Cleveland, OH, US
Contact	MAK, PHD
510(k) history	2 submissions · 2 cleared · 1989-1989

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k883556/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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