FDA 510(k) Premarket Notification

K912105 EYE TRAY

CLEARED

Aug 29, 1991
108 days to decision

K912105 · Product code: **LRP** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k912105/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tray, Surgical (LRP)
Date received	May 13, 1991
Decision date	Aug 29, 1991
Days to decision	108 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Medical Device Inspection Co., Inc.**
Location	Great Neck, NY, US
Contact	ALAN P SCHWARTZ
510(k) history	30 submissions · 26 cleared · 1990-1995