FDA 510(k) Premarket Notification

K890721 SHAPE-ACTIO METAL TOXICOLOGY CONTROL

CLEARED

Apr 10, 1989
56 days to decision

K890721 · Product code: **DIE** · Toxicology
Source: https://www.510kdatabase.net/k890721/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Heavy Metals Control Materials (DIE)
Date received	Feb 13, 1989
Decision date	Apr 10, 1989
Days to decision	56 days
Third-party review	No

APPLICANT

Company	**Actio, Inc.**
Location	Oakland, CA, US
Contact	LEWIN, PH.D.
510(k) history	4 submissions · 4 cleared · 1989-1990