

K890721 SHAPE-ACTIO METAL TOXICOLOGY CONTROLApr 10, 1989
56 days to decisionK890721 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k890721/>, Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 29, 2026