

K970574 ACUPUNCTURE NEEDLESApr 24, 1997
70 days to decisionK970574 · Product code: **MQX** · General Hospital
Source: <https://www.510kdatabase.net/k970574/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Acupuncture, Single Use (MQX)
Date received	Feb 13, 1997
Decision date	Apr 24, 1997
Days to decision	70 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	C.A.I. Industries Corp.
Location	South El Monte, CA, US
Contact	LU RAO
510(k) history	3 submissions · 3 cleared · 1997-2004

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k970574/> 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