FDA 510(k) Premarket Notification

K900440 AIA-PACK BMG

CLEARED

Feb 26, 1990
28 days to decision

K900440 · Product code: **JZG** · Immunology
Source: https://www.510kdatabase.net/k900440/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Test, Beta-2-microglobulin Immunological (JZG)
Date received	Jan 29, 1990
Decision date	Feb 26, 1990
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Tosoh Medics, Inc.**
Location	Washington, DC, US
Contact	HOWARD M HOLSTEIN
510(k) history	41 submissions · 41 cleared · 1990-2002

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k900440/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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