

**K882393 DIGITAL BLOOD PRESSURE MONITOR MODEL  
HEM-703CP**Aug 30, 1988  
81 days to decisionK882393 · Product code: **DXN** · Cardiovascular  
Source: <https://www.510kdatabase.net/k882393/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Jun 10, 1988
Decision date	Aug 30, 1988
Days to decision	81 days
Third-party review	No

**APPLICANT**

---

Company	<b>Omron Electronics, Inc.</b>
Location	Schaumburg, IL, US
Contact	OISHI
510(k) history	10 submissions · 10 cleared · 1987-1990

---

**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k882393/> 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