

K083681 BIOLAND BLOOD PRESSURE MONITORJan 16, 2009
35 days to decisionK083681 · Product code: **DXN** · Cardiovascular
Source: <https://www.510kdatabase.net/k083681/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | System, Measurement, Blood-pressure, Non-invasive (DXN) |
| Date received | Dec 12, 2008 |
| Decision date | Jan 16, 2009 |
| Days to decision | 35 days |
| Third-party review | Yes |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Bioland Technology, Ltd. |
| Location | North Point, HK |
| Contact | DAVID LEE |
| 510(k) history | 8 submissions · 8 cleared · 2009-2022 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k083681/> 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 May 27, 2026