

K810662 LEECO LH-QUANT DIAGNOSTIC KITMar 31, 1981
20 days to decisionK810662 · Product code: **CEP** · Chemistry
Source: <https://www.510kdatabase.net/k810662/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Radioimmunoassay, Luteinizing Hormone (CEP) |
| Date received | Mar 11, 1981 |
| Decision date | Mar 31, 1981 |
| Days to decision | 20 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Leeco Diagnostics, Inc. |
| Location | Mchenry, IL, US |
| 510(k) history | 49 submissions · 49 cleared · 1979-1989 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k810662/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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