

K812950 QAC CONTROLDec 17, 1981
57 days to decisionK812950 · Product code: **DIF** · Toxicology
Source: <https://www.510kdatabase.net/k812950/>**SUBMISSION DETAILS**

| | |
|-----------------------|--------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Drug Mixture Control Materials (DIF) |
| Date received | Oct 21, 1981 |
| Decision date | Dec 17, 1981 |
| Days to decision | 57 days |
| Third-party review | No |

APPLICANT

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|----------------|--|
| Company | The Quality Assurance Service Corp. |
| Location | Walker, MI, US |
| 510(k) history | 1 submissions · 1 cleared · 1981-1981 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k812950/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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