

K221417 CLAROX plus (Model: VX-100)Sep 13, 2022
120 days to decisionK221417 · Product code: **IZL** · Radiology
Source: <https://www.510kdatabase.net/k221417/>**SUBMISSION DETAILS**

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | System, X-ray, Mobile (IZL) |
| Date received | May 16, 2022 |
| Decision date | Sep 13, 2022 |
| Days to decision | 120 days |
| Third-party review | No |
| Combination product | No |
| PCCP authorized | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Vsi Co., Ltd. |
| Location | Cheongju-Si, KR |
| Contact | Jake Yu |
| 510(k) history | 2 submissions · 2 cleared · 2022-2022 |

REGULATORY CONSULTANT

| | |
|-----------------|----------------------------|
| Consulting firm | Lighten Bridge, LLC |
| Contact | Edward Park |

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k221417/> 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 26, 2026