# California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Ethicon Endo-Surgery Inc Endopath Echelon Vascular White Reload

| Recall Date | Product Description | Recalling Firm | Recall Reason |
|---|---|---|---|
| 05/20/2025 | Endopath Echelon Vascular White Reload for Advanced Placement Tip (35mm, 4 row)<br><br>Product code: VASECR35 | Ethicon Endo-Surgery Inc | Firm has received an increase in reports regarding inadvertent instrument lockout during surgical procedures involving the device. If lockout occurs, the device will momentarily activate but will not cut or staple tissue, and additional steps will be required to open it and remove it from tissue. If user is unable to remove the device from the tissue, potential harms to patient include surgical delay, bleeding/hemorrhage, life threatening hemorrhage/hemorrhagic shock, and conversion to open therapy. Two adverse events received, including one patient death. |

| Recall Class | Product Identification | Distribution | Affected Dates |
|---|---|---|---|
| I | UDI-DI: 10705036014591<br><br>Packaging Batches:<br>917C65 - 214D17 | 678,526 units nationwide | April and prior |

For additional information, please visit the FDA Website.