



California Medical Device Recall Information



Recall Name

**Edwards Lifesciences Recalls the EMBOL-X Glide Protection System Cannulae
Due to a Deformed Cannula Tip**

| Recall Date | Product Description | Recalling Firm | Recall Reason |
|--------------|---|---|--|
| 9/16/13 | EMBOL-X Glide Protection System | Edwards Lifesciences, LLC. Draper, UT | <i>Potential for deformed cannula tips.</i> |
| Recall Class | Product Identification | Distribution | Affected Dates |
| I | EMBOL-X Glide Protection System Cannulae Model Numbers recalled: <ul style="list-style-type: none"> • EXGF24D • EXGF24LLD • EXGF24MMD • EXGF24SSD • EXGF24XLD • EXGF24XSD • EXGFXS2D • EXGF24SS2D • EXGF24MM2D • EXGF24LL2D • EXGF24XL2D <p>All Lots recalled.</p> | CA , nationwide | Distributed from: Sept. 21, 2010 through Aug 22, 2013 |

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm373525.htm>