## CALIFORNIA DEVICE RECALL INFORMATION SHEET

**LumiraDx SteriPack Sterile Polyester Spun Swab for Failed Post-Sterilization Sterile Swabs Cantilever Testing**

<table>
<thead>
<tr>
<th>Recall Date</th>
<th>Product Description</th>
<th>Recalling Firm</th>
<th>Recall Reason</th>
</tr>
</thead>
</table>
| 6/21/2023   | **SteriPack Sterile Polyester Spun Swabs**  
(25 units/Pack)- Intended for sample collection in the intended nasal area. Catalog Number: 60566RevB | LumiraDx  
Waltham, MA | Fails Post-Sterilization Sterile Swabs Cantilever (Bend) Testing and may be more susceptible to breakage. If the swab breaks in the nasal cavity of a patient may cause injury or medical intervention to remove part of the swab |

<table>
<thead>
<tr>
<th>Recall Class</th>
<th>Product Identification</th>
<th>Distribution</th>
<th>Affected Dates</th>
</tr>
</thead>
</table>
| II           | **SteriPack Sterile Polyester Spun Swabs**  
GTIN-DI: 00850027193205  
Pack Lot Number : 85438;  
Individual Swab Lot Number: 86445  
Exp Date: 28-Oct-2023 | 284 Units in California | April 2023 and Prior |

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE [FDA WEBSITE](https://www.fda.gov)