

**U.S. Food and Drug Administration**  
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## Establishment Registration & Device Listing


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| <b>Proprietary Name:</b>                | Cenefom                                |
| <b>Classification Name:</b>             | BALLOON, EPISTAXIS                     |
| <b>Product Code:</b>                    | <a href="#">EMX</a>                    |
| <b>Device Class:</b>                    | 1                                      |
| <b>Regulation Number:</b>               | <a href="#">874.4100</a>               |
| <b>Medical Specialty:</b>               | Ear Nose & Throat                      |
| <b>Registered Establishment Name:</b>   | <a href="#">CENEFOM CORP</a>           |
| <b>Registered Establishment Number:</b> | 3010610457                             |
| <b>Owner/Operator:</b>                  | <a href="#">Cenefom Corp</a>           |
| <b>Owner/Operator Number:</b>           | 10045592                               |
| <b>Establishment Operations:</b>        | Manufacturer                           |

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### U.S. Food and Drug Administration

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