



May 24, 2023

RE: **SDS Exemption for Mölnlycke Exufiber® and Exufiber® Ag**

To Whom It May Concern,

Per Occupational Safety and Health (OSHA) regulation 29 CFR 1910.1200(b)(5)(iii), the following are **exempt from the hazard communication system requirements** (including hazard labels and safety data sheets):

Any food, food additive, color additive, drug, cosmetic, or **medical or veterinary device or product**, including materials intended for use as ingredients in such products (e.g. flavors and fragrances), as such terms are defined in the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 301 et seq.) or the Virus-Serum-Toxin Act of 1913 (21 U.S.C. 151 et seq.), and regulations issued under those Acts, when they are subject to the labeling requirements under those Acts by either the Food and Drug Administration or the Department of Agriculture

Exufiber and Exufiber Ag are devices that are subject to FDA labeling requirements and, as such, are exempt from SDS requirements. FDA classification information is provided below for reference.

Exufiber:

- Device: Dressing, Wound, Hydrophilic
- Regulation: 21 CFR 878.4018
- Classification: Class 1
- Product Code: NAC

Exufiber Ag:

- Device: Dressing, Wound, Drug
- Regulation: Pre-Amendment
- Classification: Unclassified
- Product Code: FRO

If you have any questions or need additional information, I can be contacted by email at megan.bevill@molnlycke.com.

Sincerely,

A handwritten signature in black ink that reads "Megan Bevill".

Megan Bevill
Regulatory Affairs Director, Americas and Antiseptics