We, Mölnlycke Health Care AB, Gamlestadsvägen 3C, Box 13080, SE-402 52 Göteborg, Sweden being exclusively responsible manufacturer for conformity of the device/s; declare that the devices listed in the attached schedule are in conformity with the provisions of the Council Directive 93/42/EEC of 14 June 1993, as amended by 2007/47/EEC, concerning medical devices, the Medical Devices Act SFS 1993:584 and the Swedish Medical Product Agency regulations and guidelines: Medical Devices, LVFS 2003:11, as amended by LVFS 2009:18.

Trade name/ Product name:	**Avance Solo Foam**

Product classification:	**IIb**
Sterility Status:	**Sterile**
Measuring function:	**No**

This declaration is supported by a conformity assessment procedure in accordance with	
Annex/es:	**II**

Certificate number:	**CE 01965**
Issued by:	**BSI (2797)**
For non sterile, non-measuring Class I products, no certificate is issued by a Notified Body.	

Mölnlycke Health Care issues this declaration in recognition of all applicable harmonised standards.

Signed for and on behalf of Mölnlycke Health Care

Date: **2020-05-13** Function: **Regulatory Affairs Manager Compliance**

Name: **Karin Darle Olsson** Signature:

Product(s) covered by this declaration:

Product Reference:	Product Descriptor:	GMDN Code:
882000	Foam wound filler	47406 Negative - pressure wound therapy system foam dressing