Valid for the following product/s as delivered by Henkel: LOCTITE AA 3921 LIGHT CURE FLOURESCENT known as 3921 Light Cure Medical Device

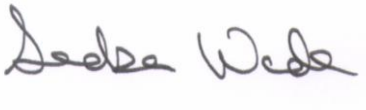
Referenced Document/s in terms of enquiry/ies, specs, standard/s or substance list/s:

EU Directive 2011/65/EU on the restriction of the use of certain hazardous substances in electrical and electronic equipment (**RoHS**), OJ 13.02.2003; esp. its Article 4, as last amended by Commission Delegated Directive (EU) 2015/863.

Confirmation:
We herewith confirm, that the above mentioned product does not contain any of the restricted substances as listed in the above referenced documents in concentrations above the limits as specified therein.

Disclaimer:
Please note that this confirmation is given to the best of our present knowledge and belief.
Nothing herein represents and/or may be interpreted as warranty within the meaning of the applicable warranty law.

Date: October 9, 2026

 _____ Senthil Pichandi Vice President Global Product Safety & Regulatory Affairs Henkel Corporation One Henkel Way Rocky Hill, CT 06067 USA	 _____ Sandra Wade Regional Head Product Safety & Regulatory Affairs – Europe Henkel Ireland Operations and Research Ltd Tallaght Business park Whitestown Tallaght, Dublin 24 Ireland