Valid for the following product/s as delivered by Henkel: LOCTITE AA 3971 LIGHT CURE MED. DEV. ADH. known as 3971 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: April 24, 2024

Senthil Pichandi
Vice President
Global Product Safety & Regulatory Affairs
Henkel Corporation
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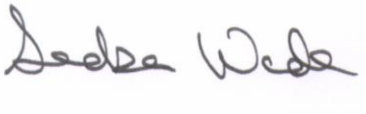

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