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Switchcraft[®]

CERTIFICATE OF COMPLIANCE – RoHS General Compliance Statement

This statement confirms that Switchcraft is aware of the requirements of EU Directive 2011/65/EU (RoHS2) as amended (EU) 2015/863 (RoHS3) and intends to comply with the requirement in regards to material content for new product releases and legacy products identified as RoHS compliant. Many legacy products and customer specific products continue to be offered in the original configuration which will not be RoHS compliant. In most cases RoHS compliant versions of the legacy products are available and are identified by adding an "X" suffix to the part number. Customer specific products are converted and re-identified as requested.

The plastic resins, base metals, metal plating or coatings used to manufacture RoHS compliant Switchcraft products do not contain any intentionally added substances listed below at or above the thresholds noted unless applied exemptions are identified:

Lead (0,1 %) * Mercury (0,1 %) Cadmium (0,01 %) ** Hexavalent chromium (0,1 %) Polybrominated biphenyls (PBB) (0,1 %) Polybrominated diphenyl ethers (PBDE) (0,1 %) Bis(2-ethylhexyl) phthalate (DEHP) (0,1 %) *** Butyl benzyl phthalate (BBP) (0,1 %) *** Dibutyl phthalate (DBP) (0,1 %) *** Diisobutyl phthalate (DIBP) (0,1 %) ***

* Per Exemption 6c, copper alloys may contain up to 4% Lead. Per exemption 6b aluminum alloys may contain up to 0.4% Lead.

** Per Exemption 8b, electrical contacts may contain Cadmium or its compounds.

*** The restriction of DEHP, BBP, DBP and DIBP shall apply to medical devices, including in vitro medical devices, and monitoring and control instruments, including industrial monitoring and control instruments, from 22 July 2021. Switchcraft does not dictate the end use of products.

*** The restriction of DEHP, BBP, DBP and DIBP shall not apply to cables or spare parts for the repair, the reuse, the updating of functionalities or upgrading of capacity of EEE placed on the market before 22 July 2019, and of medical devices, including in vitro medical devices, and monitoring and control instruments, including industrial monitoring and control instruments, placed on the market before 22 July 2021. Switchcraft does not dictate the end use of products.

*** The restriction of DEHP, BBP and DBP shall not apply to toys which are already subject to the restriction of DEHP, BBP and DBP through entry 51 of Annex XVII to Regulation (EC) No 1907/2006. Switchcraft does not dictate the end use of products.

Specific product RoHS2 Certificates of Compliance are available on request.

We reserve the right to make material changes and substitutions in our product that will not impact this certification.

None of our materials were analyzed for content, relying on the material supplier for Certifications and available MSDS.

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Signature:

Name:

Brian Smeenge

Title: Director of Engineering

Date: _27 August 2018__ Phone: _773-792-2700 X375_ email: bsmeenge@switchcraft.com