



Environmental Compliance Declaration for All Products

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Reell Precision Manufacturing (Reell) is dedicated to ensuring that our products and packaging fully comply with global environmental requirements. Reell declares that all Reell products, including our catalog and custom designed products, comply with the following:

Global RoHS Compliance

All Reell products are designed to comply with the applicable restricted substance requirements of the European Union's Restriction of the use of certain Hazardous Substances in Electrical and Electronic Equipment (RoHS) Directive (2011/65/EU) as amended by the EU RoHS Recast Directive (2015/863/EU). Reell also achieves compliance with other countries' laws that duplicate the RoHS Directive's substance restrictions, including, but not limited to, China Management Methods for Controlling Pollution by Electronic Information Products (China RoHS), Japan's Law for Promotion of Effective Utilization of Resources, and California's Electronic Waste Recycling Act.

EU REACH Compliance (**includes Reach List dated 1-19-2021**)

All Reell products comply with European Union's Regulation on the Registration, Evaluation, Authorization and Restriction of Chemicals (REACH) (2006/1907/EC) entered into force June 1, 2007. Reell maintains compliance to every update as they become relevant and in accordance with information provided by its suppliers.

California Proposition 65 (Prop 65):

The Safe Drinking Water and Toxic Enforcement Act of 1986 entered into force November 1986. None of Reell's products contain chemicals in amounts that would trigger a notification or warning under California Proposition 65.

Volatile Organic Compound (VOCs):

All Reell products comply with the United States Environmental Protection Agency (EPA) Standard 40 CFR 59.

Persistent Organic Pollutants (POPs):

All Reell products comply with the European Regulation EU 2019/1021.

Waste Framework Directive (WFD):

None of Reell's products exceed the any substances of very high concern (SVHC), therefore, there is nothing to report in the Substances of Concern in Articles (SCIP) database.

EU Medical Device Regulation (MDR):

NONE of Reell's products fall under the jurisdiction / classification of the EU 2017/745 EU entered into force 25 May 2017 and effective May 2020. The definition does not apply to Reell products.



David Titchenal

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