

Declaration of RoHS Compliance

The EU Directive 2011/65/EU & 2015/863/EU & (EU)2020/659, which refers to “Restriction of Hazardous Substances” or “RoHS”, is made by the European Union (EU) and took effect on 1 July 2011. This restricts the use of the following those hazardous substances in the manufactures of various types of the electrical and electronic devices:

- 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 %)

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.

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.

TEGUAR®

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We, Teguar Corp., hereby declare that the following component is fully compliant with the RoHS Directive 2011/65/EU & 2015/863/EU & (EU)2020/659.

15.6" TEGUAR MEDICAL DISPLAY: TMD-25-15

Tom Poplawski
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Teguar Corporation

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Tom Poplawski