## CONFORMIT



Reference No. : WTH22H05104977R1E
Applicant (Holder) : G.S.D. GROUP INC.

Address : 180 Bellerose Ouest Suite 100, Laval, H7L 6A2

Manufacturer : G.S.D. GROUP INC.

Address : 180 Bellerose Ouest Suite 100, Laval, H7L 6A2

Product Name : DIGITAL VIDEO RECORDER

Trade Mark : GSD GROUP

X=Blank or 0 - 9, A - Z, a - z

Test Model : GSD-DVRFNC4K-08-N-AI

Rating : Input: === 12V,5A

**Test Standards:** 

EN 55032:2015+A1:2020, EN 55035:2017+A11:2020, EN 50130-4:2011+A1:2014,

EN IEC 61000-3-2:2019+A1:2021, EN 61000-3-3:2013+A2:2021.

The above product has been tested by us with the listed standards and found in compliance with the 2014/30/EU European Electromagnetic Compatibility.

It is possible to use CE marking to demonstrate the compliance with this EMC Directive.

EN 55032: Electromagnetic compatibility of multimedia equipment - Emission Requirements

EN 55035: Electromagnetic compatibility of multimedia equipment — Immunity requirements

EN 50130-4: Alarm systems - Part 4: Electromagnetic compatibility - Product family standard:

Immunity requirements for components of fire, intruder, hold up, CCTV, access control and social alarm systems

EN IEC 61000-3-2: Electromagnetic compatibility (EMC) - Part 3-2: Limits - Limits for harmonic current emissions (equipment input current ≤16 A per phase)

EN 61000-3-3: Electromagnetic compatibility (EMC) - Part 3-3: Limits - Limitation of voltage changes, voltage fluctuations and flicker in public low-voltage supply systems, for equipment with rated current ≤ 16 A per phase and not subject to conditional connection

The referred test report(s) show that the product complies with standard(s) recognized as giving presumption of compliance with the essential requirements in the above mentioned EU Directive. Other relevant Directives have to be observed.

After preparation of the necessary technical documentation as well as the conformity declaration, the CE marking as shown below can be affixed on the equipment.

CE

Galthean / Project Manager

The statement is based on a single evaluation of the sample of above medical product(s). It does not imply an assessment of the whole production.

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