

# ATTESTATION OF CONFORMITY

**Name and Address of Attestation Holder:**

Taizhou Grandfar International Trading Co.,Ltd.  
23/F, B, No.190, Donghuan Street, Taizhou, Zhejiang,  
China

**Document Number**

CE-1390-01-070623

**Date of Issue:**

07.06.2023

**Name and Address of Manufacturer:**

Zhejiang Grandfar Pump Industry Co., Ltd.  
Building 61, Huigu S&T Park, Taizhou, Zhejiang, China

**Expiration Date:**

07.06.2028

**Brand:**

GRANDFAR

**Test Report Number:**

WBOViS202305-063M  
WBOViS202305-063L  
WBOViS202305-063E

**Product Name:**

Circulation Pump

**Product Model:**

See Annex I

**Test Required:**

EN ISO 12100:2010  
EN 809:1998+A1:2009+AC:2010  
EN 60204-1:2018  
EN 60335-1:2012+A11:2014+A13:2017+A1  
:2019+A14:2019+ A2:2019+A15:2021  
EN 60335-2-51:2003+A2:2012  
EN 62233:2008+AC:2008  
EN 60034-1:2010+AC:2010  
EN IEC 55014-1:2021  
EN IEC 55014-2:2021  
EN IEC 61000-3-2:2019+A1:2021  
EN 61000-3-3:2013+A1:2019+A2:2021

The product meets the technical requirement of the above standards as mentioned in the reference test reports and hence fulfils the technical requirements of the following directives

**2014/35/EU Low Voltage Directive**  
**2014/30/EU Electromagnetic Compatibility Directive**  
**2006/42/EC Machinery Directive**

CGS Test confirms type which is mentioned above according to the [Annex I] Essential Health and Safety Regulations of 2006/42/EC Machinery Directive with inspection report. Manufacturer must ensure that assessment of conformity with internal checks on the manufacture of above product according to the Annex VIII of 2006/42/EC.

This document is only valid for the equipment and configuration described, in conjunction with the test data detailed above reference test reports. Document was issued on voluntary basis and does not imply meeting Notified Body conformity assessment procedure for the product.

The CE Mark, under the responsibility of the manufacturer or the importer, after completion of an EC Declaration of Conformity and compliance with all relevant EC Directives.

**SIGNATURE**

# ATTESTATION OF CONFORMITY

## Annex I

**Product Model:**

GEA20-4-130, GEA20-6-130, GEA25-4-130,  
GEA25-5-130, GEA25-6-130, GEA25-4-180,  
GEA25-5-180, GEA25-6-180, GEA32-4-180,  
GEA32-6-180, GEA25-8-180, GEA25-10-180,  
GEA25-12-180, GEA32-8-180, GEA32-10-180,  
GEA32-12-180, GEA20-4-130U, GEA20-6-130U,  
GEA25-4-130U, GEA25-5-130U, GEA25-6-130U,  
GEA25-4-180U, GEA25-5-180U, GEA25-6-180U,  
GEA25-10-180U, GEA25-12-180U, GEA32-8-180U,  
GEA32-10-180U, GEA32-12-180U, GEB25-4 -180,  
GEB25-6 -180, GEB25-8 -180, GEB25-10 -180,  
GEB25-12 -180, GEB32-4 -180, GEB32-6 -180,  
GEB32-8 -180, GEB32-10 -180, GEB32-12 -180,  
GEB32-4 -220F, GEB32-6 -220F, GEB32-8 -220F,  
GEB32-10-220F, GEB25-4 -180N, GEB25-6 -180N,  
GEB25-8 -180N, GEB25-10 -180N, GEB25-12 -180N,  
GEB32-4 -180N, GEB32-6 -180N, GEB32-8 -180N,  
GEB32-10 -180N, GEB32-12 -180N, GEB32-4-220FN,  
GEB32-6-220FN, GEB32-8-220FN, GEB32-10-220FN,  
GEB12/1.2,

**Document Number**

CE-1390-01-070623

**Date of Issue:**

07.06.2023

**Expiration Date:**

07.06.2028

**Test Report Number:**

WBOViS202305-063M  
WBOViS202305-063L  
WBOViS202305-063E

**Test Required:**

EN ISO 12100:2010  
EN 809:1998+A1:2009+AC:2010  
EN 60204-1:2018  
EN 60335-1:2012+A11:2014+A13:2017+A1  
:2019+A14:2019+ A2:2019+A15:2021  
EN 60335-2-51:2003+A2:2012  
EN 62233:2008+AC:2008  
EN 60034-1:2010+AC:2010  
EN IEC 55014-1:2021  
EN IEC 55014-2:2021  
EN IEC 61000-3-2:2019+A1:2021  
EN 61000-3-3:2013+A1:2019+A2:2021

The product meets the technical requirement of the above standards as mentioned in the reference test reports and hence fulfils the technical requirements of the following directives

**2014/35/EU Low Voltage Directive****2014/30/EU Electromagnetic Compatibility Directive****2006/42/EC Machinery Directive**

CGS Test confirms type which is mentioned above according to the [Annex I] Essential Health and Safety Regulations of 2006/42/EC Machinery Directive with inspection report. Manufacturer must ensure that assessment of conformity with internal checks on the manufacture of above product according to the Annex VIII of 2006/42/EC.

This document is only valid for the equipment and configuration described, in conjunction with the test data detailed above reference test reports. Document was issued on voluntary basis and does not imply meeting Notified Body conformity assessment procedure for the product.

The CE Mark, under the responsibility of the manufacturer or the importer, after completion of an EC Declaration of Conformity and compliance with all relevant EC Directives.

**SIGNATURE**