

Către Agenția Medicamentului  
și Dispozitivelor Medicale

**Notificare**  
pentru înregistrarea dispozitivelor medicale în Registrul de stat  
al dispozitivelor medicale  
nr. 58 din 03.07.2023

Solicitantul „Neotec” SRL, cu sediul mun.Chisinau, str.Zaikin, 37, tel./fax: 022 852250/ 022 852252, e-mail [office@neotec.md](mailto:office@neotec.md), [agb@neotec.md](mailto:agb@neotec.md), solicit înregistrarea în Registrul de stat al dispozitivelor medicale a următoarelor categorii și tipuri de dispozitive medicale pentru introducerea și punerea la dispoziție pe piață a:

**1. Centrifuga, de laborator (24 tuburi) viteza redusa CFG-5B (WS) cu Swing Rotor Cross, No.3 Tube Rack(15ml\*24)**

**Se anexează următoarele acte:**

1. Actul prin care producătorul își desemnează reprezentantul.
2. Declarație pe proprie răspundere
3. EC Declaration Of Conformity
4. Ce Certificate

Data 03.07.2023

Digitally signed by Botnaru Andrei  
Date: 2023.06.23 10:56:10 EEST  
Reason: My Signature  
Location: Moldova



**Tabelul de recepționare a notificării**

(se completează de către Agenție în momentul depunerii notificării de către solicitant)

Comentarii cu privire la acceptul/refuzul recepționării notificării, inclusiv motivul refuzului	
Data/nr. de ordine atribuit notificării de către Agenție (în cazul acceptării recepționării)	
Numele, prenumele, funcția persoanei responsabile de recepționarea dosarului	
Semnătura persoanei responsabile	

*Anexa nr. 2*  
*La Procedurile administrative pentru notificarea*  
*dispozitivelor medicale care dețin marcajul CE*

Către Agenția Medicamentului și Dispozitive Medicale

**DECLARAȚIE PE PROPRIE RĂSPUNDERE**

Solicitant: „Neotec” SRL, cu sediul în mun.Chisinau, str.Zaikin, 37,

declar pe proprie răspundere, cunoscând prevederile art. 352<sup>1</sup>, Codul Penal al Republicii Moldova cu privire la falsul în declarații, că documentele și datele furnizate pentru notificarea dispozitivului medical:

- 1. Centrifuga, de laborator (24 tuburi) viteza redusa CFG-5B (WS) cu Swing Rotor Cross, No.3 Tube Rack(15ml\*24)**

**Sunt autentice și corespund realității.**

**Botnaru Andrei- Director**

*Numele, prenumele și funcția*

*Semnătura* \_\_\_\_\_

*Data* 03.07.2023



We, BIOEVOPEAK CO., LTD.,

based in BLDG. 3, LIGAOGUOJIHUAYUAN, NO. 1222, WEST AOTI ROAD, LIXIA DISTRICT, JINAN, SHANDONG P.C.: 250100, assign Neotec SRL, based in Str I. Zaikin 37, Chisinau MD -2005, Moldova, as **authorized representative** in correspondence with the conditions of directive 93/42/EEC, 98/79/EEC and 90/385/EEC.

We declare that the company mentioned above is authorized to register, notify, renew or modify the registration of medical devices on the territory of the Republic of Moldova.

Place: Sales representative

Date: March 30, 2023

Signed: 2023.3.30



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# Bioevoke Co., Ltd.

NO.4 BLDG MINGSHI HAOTING, JINGSHI ROAD, LIXIA DISTRICT,  
JINAN CITY, SHANDONG PROVINCE, CHINA. P.C.: 250100  
TEL: +86-531-88982330 FAX: +86-531-88983691  
EMAIL: info@bioevoke.com WEBSITE: www.bioevoke.com

## DECLARATION OF CONFORMITY

Technical file of the company mentioned below has been inspected and audit has been completed successfully 2014/35/EU Low Voltage Directive and 2014/30/EU Electromagnetic Compatibility Directive have been taken as reference for these processes

Company Name: Bioevoke Co., Ltd.

Brand: BIOEVOKE

Related Directives and Annex: 2014/35/EU Low Voltage Directive (LVD)  
2014/30/EU Electromagnetic Compatibility (EMC)

Related Standards: EN 61326-1:2013; EN 61010-1:2010

Product(s): Centrifuge

Type(s)/Model(s): CFG-5BL; CFGR-5B; CFGR-5BL; CFGR-B8B; CFGR-B5B;  
CFG-550; CFG-4.5D ; CFG-4B; CFG-4BA; CFG-4D; CFG-4D-E ;  
CFG-4Z; CFG-4ZA; CFG-4ZB; CFG-4ZBP; CFG-4ZC;  
CFG-4ZCP; CFG-4ZL; CFG-4ZN; CFG-4ZS ; CFG-5B; CFG-5B;  
CFG-5B-A; CFG-5D; CFG-6B; CFG-5B; CFG-6D; CFG-6J;  
CFG-550J; CFG-B5BL; CFG-12D; CFG-14D; CFG-15D;  
CFG-18.5J; CFG-21J; CFG-16B; CFG-18.5B; CFG-16.5B;  
CFG-20.5B; CFG-580; CFGR-23J; CFGR-18.5J; CFGR-24J;  
CFGR-24JL; CFGR-550J; CFGR-B550J; CFGR-B16.5B;  
CFGR-17B; CFGR-12BP; CFGR-20.5B ; CFGR-25BP;  
CFGR-22BP; CFGR-21BP; CFGR-B20B; CFGR-B18B;  
CFGR-B16B; CFGR-B21B; CFGR-B580; CFG-Mini4B;  
CFG-MINI5D; CFG-MINI7D; CFG-Mini6B; CFG-Mini6R;  
CFG-Mini10B; CFG-Mini15D; CFG-Mini15R; CFG-MP2R;  
CFG-4.5D(PRP); CFG-5B(PRP); CFG-4B(PRP); CFG-4B(oil);  
CFG-5B(oil); CFG-12B(HCT); CFG-3B (Cyto);  
CFG-4B (BT); CFG-4B (BW); CFG-5B(FT)

Parameters: AC110-240 V, 50Hz/60Hz

Classification: Centrifuge

Examination Period: March 29, 2021

Date of Expiry: March 29, 2026

Review Result: We, Bioevoke Co., Ltd. declare that during the self-testing and performance evaluation, no Non-compliance according to the requirements of the Low Voltage Directive 2014/35/EC and Electromagnetic Compatibility Directive 2014/30/EU was detected.

Year of DOC marking: 2021

Signed for and on behalf of

Company: Bioevoke Co., Ltd.

General Manager

Document No: BEPSD-2119002



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Location: Moldova



# C E R T I F I C A T E

## ATTESTATION CERTIFICATE OF ELECTROMAGNETIC COMPATIBILITY AND LOW VOLTAGE DIRECTIVES

Technical file of the company mentioned below has been observed and audit has been completed successfully.

2014/30/EU Electromagnetic Compatibility Directive and  
2014/35/EU Low Voltage Directives have been taken as references for these processes

Company Name	: Bioevopeak Co., Ltd.
Company Address	: 1-1509, No.4 Bldg Mingshi Haoting, Jingshi Rd, Jinan City, Shandong Province, China
Related Directives and Annex	: 2014/35/EU Low Voltage Directive 2014/30/EU Electromagnetic Compatibility Directive
Related Standards	: EN 61010-1:2010; EN 61010-2-020:2006; EN 61326-1:2013
Product Name	: Centrifuge
Report No and Date	: EC.BIO.20210507001-R-2
Product Brand/Model/Type	: CFGR-25BP, CFGR-22BP, CFGR-21BP, CFGR-12BP, CFGR-25B, CFGR-22B, CFGR-21B, CFGR-20.5B, CFGR-20B, CFGR-12B, CFGR-B20B, CFGR-B18B, CFGR-B16.5B, CFGR-B16B, CFGR-6B, CFGR-8BL, CFGR-5B, CFGR-B21B, CFGR-8B, CFGR-5BL, CFGR-B8B, CFGR-B5B, CFGR-B580, CFGR-24JL, CFGR-24J, CFGR-23J, CFGR-18.5J, CFGR-550J, CFGR-B550J, CFG-18.5B, CFG-16.5B, CFG-16BA, CFG-16B, CFG-16.8BA, CFG-5BL, CFG-6B, CFG-B5BL(WS), CFG-B5BL, CFG-5B-A, CFG-5B(WS), CFG-5B, CFG-4B, CFG-4B(WS), CFG-4BA, CFG-4BA(WS), CFG-20.5B, CFG-580, CFG-550, CFG-5B(PRP), CFG-4B(PRP), CFG-5B(FT), CFG-12B(HCT), CFG-4B(BW), CFG-4B (BT), CFG-3B (Cyto), CFG-5B(GL), CFG-5B(FC), CFG-5B(BD), CFG-4B (Oil), CFG-5B (Oil), CFG-21J, CFG-18.5J, CFG-6J, CFG-550J, CFG-5J, CFG-5J(PRP), CFG-4ZN, CFG-4ZA, CFG-4Z, CFG-4ZS, CFG-4ZB, CFG-4ZBP, CFG-4ZCP, CFG-4ZC, CFG-4ZL, CFG-Mini4B, CFG-Mini6B, CFG-Mini10B, CFG-Mini6R, CFG-MP2R, CFG-Mini15R, CFG-Mini4M, CFG-Mini6M, CFG-Mini7M, CFG-Mini10M

Certificate Number	: M.2021.206.C64662
Initial Assessment Date	: 18.05.2021
Registration Date	: 19.05.2021
Reissue Date/No	: -
Expiry Date	: 18.05.2026

  
UDEM International Certification  
Auditing Training Centre Industry  
and Trade Inc. Co.

The validity of the certificate can be checked through [www.udem.com.tr](http://www.udem.com.tr). The CE mark shown on the right can only be used under the responsibility of the manufacturer with the completion of EC Declaration of Conformity for all the relevant Directives. This certificate remains the property of UDEM International Certification Auditing Training Centre Industry and Trade Inc. Co. to whom it must be returned upon request. The above named firm must keep a copy of this certificate for 15 years from the registration of certificate. This certificate only covers the product(s) stated above and UDEM must be noticed in case of any changes on the product(s)

Address: Mutlukent Mahallesi 2073 Sokak (Eski 93 Sokak) No:10 Çankaya - Ankara - TURKEY  
Phone: +90 0312 443 03 90 Fax: +90 0312 443 03 76  
E-mail: [info@udemltd.com.tr](mailto:info@udemltd.com.tr) [www.udem.com.tr](http://www.udem.com.tr)

