

## EC DECLARATION OF CONFORMITY

**We certify that the following designated product**

Product Name: Doctor Centrifuge

Model No.: i fuge D06

**Is in fully conformity with the harmonized standard(s)**

EN 61326-1:2013 , EN 61326-2-6:2013, EN 61010-1, EN 61010-2-020, EN 61010-2-101

**Under the EU Council Directive**

2014/30/EU on Electromagnetic Compatibility(EMC),  
Low Voltage Directive (LVD) 2014/35/EU,  
In vitro diagnostic medical device Directive(IVD) 98/79/EC  
RoHS Directive 2015/863/EU,  
WEEE Directive 2012/19/EU

**The designated product was subject to sample testing for which  
issued test report number:**

YMPL/296204/81081 , YMPL/296209/81082, YMPL/296206/81088, YMPL/297004/83192,  
YMPL/296218/81090, YMPL/292500/81091, YMPL/296201/81092, YMPL/296202/81093  
RP-1718-000025, RP-1718-000007, RP-1718-000016  
ITC/TEST/NN/1611/05-E1 , ITC/TEST/NN/1611/05-E2, ITC/TEST/NN/1611/ 05-E3

**The declaration is the sole responsibility of the manufacturer / importer**

Neuation Technologies Pvt Ltd  
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17/Sep/2019

Date



Signature & Company stamp