Our Case Ref.:
#AR-NuzoneMedical_USA_I_150923#
M: +1-509-481-1919 F: 888-812-7474

Your Ref./Contact Persons:
Mr Dan Ogilvie, CEO
E: dan@nuzonemedical.com

To:

Nuzone Medical LLC.
(c/o Mr Dan Ogilvie)
E. 1418 14th Ave,
Spokane, WA 99202,
USA

CE
European Authorised Representative For Medical Devices

Auth Rep Certificate

No. **ARMDD151356A151012LN**
Valid until: **22 Sept 2020**

This is to certify that **Wellkang Ltd** has formally accepted the appointment as the European Authorised Representative (EC Rep) for manufacturer- **Nuzone Medical LLC.,** located at: **E. 1418 14th Ave, Spokane, WA 99202, USA,** as required by the EU MDD Directive 93/42/EEC of 14 June 1993 amended by Directive 2007/47/EC. The manufacturer may use Wellkang's name & address as the EC Rep for the CE-marked products represented by Wellkang. The examples of use of Wellkang's name/address can be found at: **http://www.ce-marking.com/ec-rep.html**

This representation including the information on the products represented is subject to the terms and conditions stated in the Authorised Representative Agreement signed between our two companies, and has been published online for verification by third parties at -
http://www.cemark.info/mdd/NuzoneMedical.html

When you start to place the above-mentioned product(s) on the EEA Market, please make sure to properly affix the CE Marking on the product(s), the labelling, packaging, and/or other accompanying materials according to the related EU directives and guidelines which have already been provided to you by us.

Please be advised that Wellkang is NOT involved in the Design, Manufacture and/or Marketing, Distribution, Sales, Supply, Installation of your products, it is therefore your responsibility to provide the Instruction For Use (IFU), if applicable and required by the legislation of the EEA member state(s), in the official language(s) of the EEA member state(s) in which the products are placed on the market. You need to provide us with the true, accurate and most updated Technical Documentation (Technical Files) including IFU, per EU Decision No. 768/2008/EC, prior to any shipment of your products, which carry the name of Wellkang Ltd as the Authorised Representative, to the EEA market.

Please inform us immediately whenever there is a change about either the above-mentioned product(s) or your company details. So we can update your CE Marking documentation promptly and properly.

London/UK, 12th Oct 2015.

For and on behalf of
WELLKANG LIMITED

Authorized Signature(s)

Signature of Responsible
Wellkang Ltd