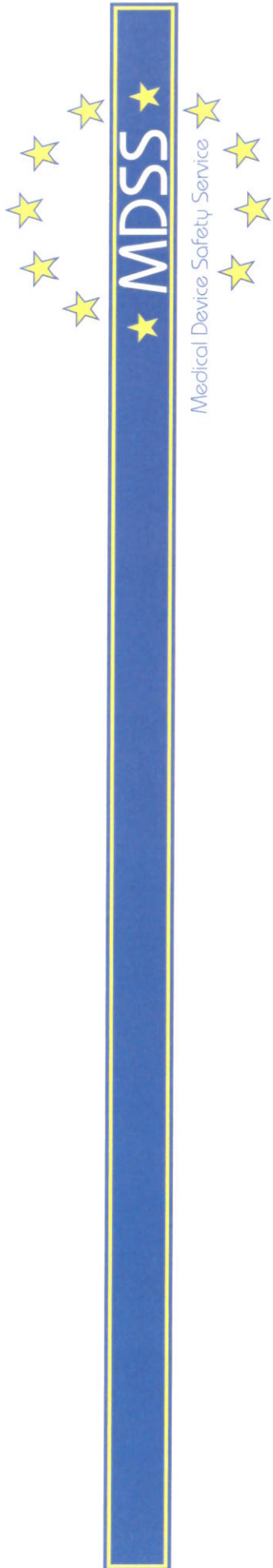


Certificate of CE-Registration



This is to certify that, in accordance with the *In Vitro* Diagnostic Medical Device Directive 98/79/EC, Medical Device Safety Service GmbH (MDSS) agrees to perform all duties and responsibilities as the Authorized Representative for:

**MacBrud Corporation
14021 SW 143 COURT #6
MIAMI, FL 33186
UNITED STATES OF AMERICA**

as stipulated and demanded by the aforementioned Directive. The European Databank on Medical Devices (EUDAMED) is established as of May 1, 2011. The German Competent Authority is notified of the manufacturer's *in vitro* diagnostic medical devices and has allocated registration numbers shown in:

Annex A dated January 11, 2016

The Manufacturer has provided MDSS with the appropriate Declaration(s) of Conformity confirming that the *in vitro* diagnostic medical devices fulfill the applicable requirements of Directive 98/79/EC. In compliance with German law, a safety officer has been appointed for Germany.

January 11, 2016


Joy Grimm
Senior Consultant - IVD
MDSS GmbH

Annex A: January 11, 2016
Manufacturer: MacBrud Corporation



| THE FOLLOWING NOTIFICATIONS WERE SUBMITTED AFTER EUDAMED IMPLEMENTATION | |
|---|--|
| Registration No.: | DE/CA09/0170/IVD/5651 |
| EDMA Code: | 51 09 10 01 |
| EDMA Description: | Other containers for samples of human origin |
| Risk Class: | "Other" |
| EC Certificate: | N/A |
| Certificate Expiry: | N/A |
| Device Name(s): | SpecBoard CE |
| Manufacturer's Device Identification | |
| SpecBoard CE | 777 |