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 NOTFICATIONS 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