SIEMENS

Test Certificate

on the Effect of Third Party Products
on MRI Systems

1. Test Certificate, device designation and manufacturer

This is to certify that the following device

BrainAmp MR plus 64-channel EEG system
tested with AMP0611591MR plus and AMP0801787MR plus

BrainCap MR electrode cap 64-channel
tested with 3/578

Power Pack Box 5,6V
tested with PP0712306

of the manufacturer:

Brain Products GmbH
Zeppelinstrasse 7

D-82205 Gilching

does not impair the functioning and safety of the Siemens AG, Healthcare Sector, Imaging & IT Division, Magnetic Resonance Systems described in section 2 as long as the "Instructions for Use" of both the MRI system and the BrainAmp EEG system are strictly followed.

Possible functional restrictions of the MRI Systems are indicated in section 4.

Possible adverse effects of the MRI Systems on the device described above are explicitly not subject to the test on which this certificate is based. Therefore consequently, this certificate does not imply non-interference of above-mentioned product through the MRI system.

H30-S1002-N41 500.000 10.03 87783

2. MAGNETOM Systems affected

Seq. no.	MRI System or option (type designation)
1	Trio 3T
2	Trio, A Tim System 3T
3	Verio 3T
4	Verio DOT 3T
5	Skyra 3T
6	Skyra^{fit} 3T
7	Spectra 3T
8	Prisma 3T
9	Prisma^{fit} 3T

3. Scope

In connection with the MRI application the EEG de*vice* is intended for the following use to which the Test Certificate refers:

- EEG recording

4. Restrictions

During use of the device there are the following restrictions of functions and/or application possibilities of the MRI System, the systems, options, accessories:

- n.a.

5. Warnings

When using this device in connection with the MRI Systems described in section 2 the following precautions must be observed:

- n.a.

6. Validity:

This Test Certificate is valid until revoked by Siemens AG, Healthcare Sector.

Erlangen, 2014-09-23

Siemens Aktiengesellschaft

i. V. Bank, H MR SPR

i. V. ~~Dr. Beckmann~~ Reiner Lochner