PICO-TESLA

the *MAGNECEUTICAL*® company

Certificate of Compliance

Date: 3/14/2012

Product Name: Magnesphere™

Catalog/Model Number: 7001

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The Magnesphere™ is classified under the U.S. Food and Drug Administration (FDA) regulation 21 CFR 890.5660. Under this product regulation the Magnesphere™ device is classified as a Class I medical device, exempt from 510(k) Premarket Submission. The FDA Product Code for the Magnesphere™ is ISA by the Physical Medicine Review Panel. The Magnesphere™ FDA product listing number is D134631.

Pico Tesla Magnetic Therapies, LLC, the manufacturer of the Magnesphere™, carries an FDA Establishment Registration number of 3009428929.

The Magnesphere™ has been designed and manufactured to meet FDA Quality System Regulation requirements. In addition, Pico Tesla Magnetic Therapies Quality Management System has been certified to ISO13485:2003, by the British Standards Institute, under certificate number FM 568199.

Allen Braswell Jr.,
President / Managing Member
Pico-Tesla Magnetic Therapies, LLC.