

QUALITY ASSURANCE

Tiller MIND BODY, INC. San Antonio, Texas (USA) is the manufacturer and distributor of "The LIBBE". LIBBE Devices are found world wide in conformance as follows:

- USA FDA (Food & Drug Administration 21 CFR § part 820)
- USA FDA (Registered Device Establishment License)
- Texas Licensed Medical Device Manufacturer
- Quality System with ISO 13485:2003
- CE 0459 Certification through Europe Quality & Production Assurance
- Health Canada Medical Device License No.78207
- Australia ((ARTG) Licensed
- I-ACT Recognized and Colon Hydrotherapy School
- Texas Licensed Education Provider

The LIBBE Colonic Irrigation Device is an FDA-registered Class II device for colon cleansing when medically indicated, such as before radiological or endoscopic examination. Colon irrigation takes place in an integrated clinical setting, affording the patient both modesty and dignity and the practitioner ease-of-use. The LIBBE Device is comfortable and allows the patient the simple release of the contents of the large Intestine (colon) during a 45-minute session. Each session uses 10 gallons or less of filtered, ultraviolet purified Water, which is controlled by a temperature control sensor. Should the water temperature go above 104 F (40 C) the water flow will automatically shut off.

The LIBBE Device was developed to assist the healthcare practitioner and patient for a simpler comfortable preparatory method prior to medical procedures, such as before a colonoscopy or when medically indicated.

LIBBE Device Training is required at a LIBBE Recognized School. The LIBBE Device has grown in acceptance worldwide as the colon hydrotherapy device that provides its patients and therapists with ease of use, safety, proper sanitization, easy maintenance & cleaning, Quick Reference Guides, low-cost disposables and supplies, and an optional Worldwide Internet Listing with over 400,000 monthly inquiries searching for a LIBBE Center/Therapist!

For Information E-mail: LIBBE@colonic.net

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FDA U.S. Food and Drug Administration
Class II Medical Device - Since 1995
Quality System FDA 21 CFR Part 820

ISO 13485: 2003

EC	REP
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CE 0459

United States
Australia
Health Canada
European Union
Others

EMERGO EUROPE
Malentsraat 15
2513 BH, The Hague
The Netherlands

EMERGO CANADA
1275 West 6th Ave,
Vancouver, BC Canada

EMERGO AUSTRALIA
Level 20, Tower II, Darling Park
201 Sussex Street
Sydney, NSW 2000 Australia

Rx
Required for
Device/Nozzle
Purchase

www.colonic.net