EMORD & ASSOCIATES, P.C.

THE DO'S AND DON'TS OF HOMEOPATHIC DRUGS
(from Ingredients to Labeling)

AN EMORD & ASSOCIATES TELECONFERENCE

Teleconference Date:

April 17, 2008

What Rules Govern Homeopathic drugs and their labels?

The Food and Drug Administration does not require pre-market approval for homeopathic drugs, but many do not understand the limits FDA places on the ingredients that may be used in homeopathy and the labeling content permitted for homeopathic drugs. In this informative teleconference, the attorneys at Emord & Associates will explain:

1. The legal limits on addition off non-homeopathic ingredients to homeopathic drugs
2. The FDA's definition of homeopathic drugs and what ingredients qualify for exemption from the drug pre-market approval process
3. The claims permitted on the labels and in the labeling of homeopathic drugs and the consequences of deviating from permitted claims
4. The risks posed by the FTC to companies that advertise homeopathic drugs

To participate, call Emord & Associates, P.C., (202) 466-6937 or email us at jemord.com The fee for participation (payable by credit card) is $225. A pass code and phone number for the April 17th program will be supplied to you upon paying the fee.

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