

UNITED STATES OF AMERICA
BEFORE FEDERAL TRADE COMMISSION

COMMISSIONERS: Robert Pitofsky, Chairman
Mary L. Azcuenaga
Janet D. Steiger
Roscoe B. Starek, III

In the Matter of)	DOCKET NO. C-3770
)	
KAVE ELAHIE d/b/a M.E.K. INTERNATIONAL,)	
)	DECISION AND ORDER
a sole proprietorship.)	
)	

The Federal Trade Commission having initiated an investigation of certain acts and practices of the respondent named in the caption hereof, and the respondent having been furnished thereafter with a copy of a draft of complaint which the San Francisco Regional Office proposed to present to the Commission for its consideration and which, if issued by the Commission, would charge respondent with violation of the Federal Trade Commission Act; and

The respondent and counsel for the Commission having thereafter executed an agreement containing a consent order, an admission by the respondent of all the jurisdictional facts set forth in the aforesaid draft of complaint, a statement that the signing of said agreement is for settlement purposes only and does not constitute an admission by respondent that the law has been violated as alleged in such complaint, or that the facts as alleged in such complaint, other than jurisdictional facts, are true and waivers and other provisions as required by the Commission's Rules; and

The Commission having thereafter considered the matter and having determined that it had reason to believe that the respondent has violated the said Act, and that a complaint should issue stating its charges in that respect, and having thereupon accepted the executed consent agreement and placed such agreement on the public record for a period of sixty (60) days, now in further conformity with the procedure prescribed in § 2.34 of its

Rules, the Commission hereby issues its complaint, makes the following jurisdictional findings and enters the following order:

1. Respondent Kave Elahie is the sole proprietor of M.E.K. International, a California company with its principal office or place of business at 1669 Emeric Street, Simi Valley, California 93065.

2. The Federal Trade Commission has jurisdiction of the subject matter of this proceeding and of the respondent, and the proceeding is in the public interest.

ORDER

DEFINITIONS

For the purposes of this order, the following definitions shall apply:

1. "Competent and reliable scientific evidence" shall mean tests, analyses, research, studies, or other evidence based on the expertise of professionals in the relevant area, that has been conducted and evaluated in an objective manner by persons qualified to do so, using procedures generally accepted in the profession to yield accurate and reliable results; and

2. Unless otherwise specified, "respondent" shall mean Kave Elahie, individually and doing business as M.E.K. International, his successors and assigns and each of his officers agents, representatives, and employees.

3. "Commerce" shall mean as defined in Section 4 of the Federal Trade Commission Act, 15 U.S.C. § 44.

I.

IT IS ORDERED that respondent, directly or through any corporation, subsidiary, division, or other device, in connection with the labeling, advertising, promotion, offering for sale, sale, or distribution of the "NutraTrim Bio-Active Cellulite Control Cream," or "NutraTrim Weight Loss" tablets, or any other food, drug, or dietary supplement, as "food" and "drug" are defined in Section 15 of the Federal Trade Commission Act, in or affecting commerce, shall not make any representation, in any manner, expressly or by implication:

- A. That such product causes, aids, facilitates or contributes to reducing body fat;
- B. That such product causes, aids, facilitates or contributes to causing rapid weight or body fat loss;
- C. That such product causes or assists in causing weight or fat loss without dieting or strenuous exercise;
- D. That such product reduces serum cholesterol levels;
- E. That such product increases human metabolism;
- F. That such product controls appetite;
- G. That such product increases energy or stamina; or
- H. That such product eliminates cellulite or fat;

unless at the time it is made, respondent possesses and relies upon competent and reliable scientific evidence that substantiates the representation.

II.

IT IS FURTHER ORDERED that respondent, directly or through any corporation, subsidiary, division, or other device, in connection with the labeling, advertising, promotion, offering for sale, sale, or distribution of the "NutraTrim Bio-Active Cellulite Control Cream," or "NutraTrim Weight Loss" tablets, or any other food, drug, or dietary supplement, as "food" and "drug" are defined in Section 15 of the Federal Trade Commission Act, in or affecting commerce, shall not make any representation, in any manner, expressly or by implication, regarding the performance, benefits, efficacy, or safety of such product, unless at the time it is made, respondent possesses and relies upon competent and reliable scientific evidence that substantiates the representation.

III.

IT IS FURTHER ORDERED that respondent, directly or through any corporation, subsidiary, division, or other device, in connection with the labeling, advertising, promotion, offering for sale, sale, or distribution of the "NutraTrim Bio-Active Cellulite Control Cream," or "NutraTrim Weight Loss" tablets, or any other food, drug, or dietary supplement, as "food" and "drug" are defined in Section 15 of the Federal Trade Commission Act, in or affecting commerce, shall not make any misrepresentation, in any manner, expressly or by implication, regarding the existence,

contents, validity, results, conclusions, or interpretations of any test or study.

IV.

IT IS FURTHER ORDERED that respondent, directly or through any corporation, subsidiary, division, or other device, in connection with the labeling, advertising, promotion, offering for sale, sale, or distribution of any product in or affecting commerce, shall not represent, in any manner, expressly or by implication, that the experience represented by any user testimonial or endorsement of the product represents the typical or ordinary experience of members of the public who use the product, unless:

- A. At the time it is made, respondent possesses and relies upon competent and reliable scientific evidence that substantiates the representation; or
- B. Respondent discloses, clearly and prominently, and in close proximity to the endorsement or testimonial, either:
 - 1. what the generally expected results would be for users of the product, or
 - 2. the limited applicability of the endorser's experience to what consumers may generally expect to achieve, that is, that consumers should not expect to experience similar results.

For purposes of this Part, "endorsement" shall mean as defined in 16 C.F.R. § 255.0(b).

V.

Nothing in this order shall prohibit respondent from making any representation for any product that is specifically permitted in labeling for such product by regulations promulgated by the Food and Drug Administration pursuant to the Nutrition Labeling and Education Act of 1990.

VI.

Nothing in this order shall prohibit respondent from making any representation for any drug that is permitted in labeling for such drug under any tentative final or final standard promulgated by the Food and Drug Administration, or under any new drug application approved by the Food and Drug Administration.

VII.

IT IS FURTHER ORDERED that respondent shall, for five (5) years after the last date of dissemination of any representation covered by this order, maintain and upon request make available to the Federal Trade Commission for inspection and copying:

A.

IT IS FURTHER ORDERED that respondent shall, within sixty (60) days after the date of service of this order, and at such other times as the Federal Trade Commission may require, file with the Commission a report, in writing, setting forth in detail the manner and form in which he has complied with this order.

XI.

This order will terminate on September 19, 2017, or twenty (20) years from the most recent date that the United States or the Federal Trade Commission files a complaint (with or without an accompanying consent decree) in federal court alleging any violation of the order, whichever comes later; provided, however, that the filing of such a complaint will not affect the duration of: