The Federal Trade Commission ("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 a complaint which the Western Region-San Francisco proposed to present to the Commission for its consideration and which, if issued, would charge the respondent with violations of the Federal Trade Commission Act; and

The respondent, its attorney, and counsel for the Commission having thereafter executed an Agreement Containing Consent Order ("consent agreement"), which includes: a statement by respondent that it neither admits nor denies any of the allegations in the draft complaint except as specifically stated in the consent agreement, and, only for purposes of this action, admits the facts necessary to establish jurisdiction; 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 Federal Trade Commission 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 thirty (30) days for the receipt and consideration of public comments, and having duly considered the comments received from interested persons pursuant to Commission Rule 2.34, 16 C.F.R. § 2.34, now in further conformity with the procedure prescribed in Commission Rule 2.34, the Commission hereby issues its complaint, makes the following jurisdictional findings and enters the following order:

1. Respondent Lornamead, Inc., is a Delaware corporation with its principal office or place of business at 175 Cooper Avenue, Tonawanda, New York 14150.

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 purposes of this order, the following definitions shall apply:

1. Unless otherwise specified, “respondent” shall mean Lornamead, Inc., a corporation, its successors and assigns, and its officers, agents, representatives, and employees.

2. “Adequate and well-controlled human clinical study” means a human clinical study that is randomized, double-blind, placebo controlled, and conducted by persons qualified by training and experience to conduct such study.


4. “Covered Product” means any drug, cosmetic, or pesticide, including but not limited to Lice Shield Products.


6. “Essentially Equivalent Product” means a product that contains the identical ingredients, except for inactive ingredients (e.g., binders, colors, fillers, excipients), in the same form and dosage, and with the same route of administration (e.g., orally, sublingually), as the Covered Product; provided that the Covered Product may contain additional ingredients if reliable scientific evidence generally accepted by experts in the field demonstrates that the amount and combination of additional ingredients is unlikely to impede or inhibit the effectiveness of the ingredients in the Essentially Equivalent Product.

7. “Lice Shield Products” means any lice repellent product containing essential oils such as citronella, including, but not limited to Lice Shield Shampoo & Conditioner in 1, Lice Shield Leave In Spray, Lice Shield Gear Guard, and Lice Shield Long Lasting Spot Stick.

8. “Pediculosis” means infestation of the scalp by head lice.

9. The term “including” in this order means “without limitation.”

10. The terms “and” and “or” in this order shall be construed conjunctively or disjunctively as necessary, to make the applicable phrase or sentence inclusive rather than exclusive.
I.

**IT IS ORDERED** that respondent, directly or through any corporation, partnership, subsidiary, division, trade name, or other device, in connection with the manufacturing, labeling, advertising, promotion, offering for sale, sale, or distribution of any Covered Product, in or affecting commerce, shall not make any representation, in any manner, expressly or by implication, including through the use of a product name, endorsement, depiction, or illustration, that the Covered Product is effective in: a) preventing pediculosis, b) eliminating or reducing the risk of pediculosis by a specific percentage or amount, or c) repelling all lice, or a specific percentage or amount of lice, from a person’s head, unless the representation is non-misleading, and, at the time it is made, respondent possesses and relies upon competent and reliable scientific evidence that substantiates that the representation is true. For purposes of this Part I, competent and reliable scientific evidence shall consist of at least one adequate and well-controlled human clinical study of the Covered Product, or of an Essentially Equivalent Product, that conforms to an acceptable design and protocol and whose results, when considered in light of the entire body of relevant and reliable scientific evidence, are sufficient to substantiate that the representation is true. Respondent shall have the burden of proving that a product satisfies the definition of Essentially Equivalent Product.

II.

**IT IS FURTHER ORDERED** that respondent, directly or through any corporation, partnership, subsidiary, division, trade name, or other device, in connection with the manufacturing, labeling, advertising, promotion, offering for sale, sale, or distribution of any Covered Product, in or affecting commerce, shall not make any representation, other than representations covered by Part I of this order, in any manner, expressly or by implication, including through the use of a product name, endorsement, depiction, or illustration, that the Covered Product will reduce the risk of a head lice infestation or repel head lice, unless the representation is non-misleading, and, at the time of making such representation, the respondent possesses and relies upon competent and reliable scientific evidence that is sufficient in quality and quantity based on standards generally accepted in the relevant scientific fields, when considered in light of the entire body of relevant and reliable scientific evidence, to substantiate that the representation is true. For purposes of this Part II, competent and reliable scientific evidence means tests, analyses, research, or studies that have been conducted and evaluated in an objective manner by qualified persons, and that are generally accepted in the profession to yield accurate and reliable results.

III.

**IT IS FURTHER ORDERED** that respondent, directly or through any corporation, partnership, subsidiary, division, trade name, or other device, in connection with the manufacturing, labeling, advertising, promotion, offering for sale, sale, or distribution of any Covered Product, in or affecting commerce, shall not make any representation, other than representations covered under Part I of this order, in any manner, expressly or by implication, including through the use of a product name, endorsement, depiction, or illustration, about the
health benefits of any Covered Product unless the representation is non-misleading, and, at the
time of making such representation, the respondent possesses and relies upon competent and
reliable scientific evidence that is sufficient in quality and quantity based on standards generally
accepted in the relevant scientific fields, when considered in light of the entire body of relevant
and reliable scientific evidence, to substantiate that the representation is true. For purposes of
this Part III, competent and reliable scientific evidence means tests, analyses, research, or studies
that have been conducted and evaluated in an objective manner by qualified persons, and that are
generally accepted in the profession to yield accurate and reliable results.

IV.

IT IS FURTHER ORDERED that respondent, directly or through any corporation,
partnership, subsidiary, division, trade name, or other device, in connection with the
manufacturing, labeling, advertising, promotion, offering for sale, sale, or distribution of any
Covered Product in or affecting commerce, shall not misrepresent, in any manner, expressly or
by implication, including through the use of any product name or endorsement, depiction, or
illustration, the existence, contents, validity, results, conclusions, or interpretations of any test,
study, or research.

V.

IT IS FURTHER ORDERED that nothing in this order shall prohibit respondent from
making any representation for any drug that is permitted in the 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.

VI.

IT IS FURTHER ORDERED that respondent shall pay to the Federal Trade
Commission the sum of five hundred thousand dollars ($500,000). This payment shall be made
in the following manner:

A. The payment shall be made by electronic funds transfer within ten (10) days after
the date that this order becomes final and in accordance with instructions
provided by a representative of the Federal Trade Commission.

B. In the event of default on any obligation to make payment under this order,
interest, computed pursuant to 28 U.S.C. § 1961(a), shall accrue from the date of
default to the date of payment. In the event such default continues for ten (10)
calendar days beyond the date that payment is due, the entire amount shall
immediately become due and payable.

C. All funds paid to the Commission pursuant to this order shall be deposited into
an account administered by the Commission or its agents to be used for equitable
relief, including restitution, and any attendant expenses for the administration of
such equitable relief. In the event that direct redress to consumers is wholly or partially impracticable or funds remain after the redress to consumers (which shall be the first priority for dispensing the funds set forth above) is completed, the Commission may apply any remaining funds for such other equitable relief (including consumer information remedies) as it determines to be reasonably related to respondent’s practices alleged in the complaint. Any funds not used for such equitable relief shall be deposited in the United States Treasury as disgorgement. Respondent shall be notified as to how the funds are distributed, but shall have no right to challenge the Commission’s choice of remedies under this Part. Respondent shall have no right to contest the manner of distribution chosen by the Commission. No portion of any payment under this Part shall be deemed a payment of any fine, penalty, or punitive assessment.

D. Respondent relinquishes all dominion, control, and title to the funds paid to the fullest extent permitted by law. Respondent shall make no claim to or demand for return of the funds, directly or indirectly, through counsel or otherwise.

E. Respondent agrees that the facts as alleged in the complaint filed in this action shall be taken as true without further proof in any bankruptcy case or subsequent civil litigation pursued by the Commission to enforce its rights to any payment or money judgment pursuant to this order, including but not limited to a nondischargeability complaint in any bankruptcy case. Respondent further agrees that the facts alleged in the complaint establish all elements necessary to sustain an action by the Commission pursuant to Section 523(a)(2)(A) of the Bankruptcy Code, 11 U.S.C. § 523(a)(2)(A), and that this order shall have collateral estoppel effect for such purposes.

F. In accordance with 31 U.S.C. § 7701, respondent is hereby required, unless it has done so already, to furnish to the Commission its taxpayer identifying number, which shall be used for the purposes of collecting and reporting on any delinquent amount arising out of respondent’s relationship with the government.

G. Proceedings instituted under this Part are in addition to, and not in lieu of, any other civil or criminal remedies that may be provided by law, including any other proceedings the Commission may initiate to enforce this order.

VII.

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

A. All advertisements and promotional materials containing the representation;

B. All materials that were relied upon in disseminating the representation; and
C. All tests, reports, studies, surveys, demonstrations, or other evidence in its possession or control that contradict, qualify, or call into question the representation, or the basis relied upon for the representation, including complaints and other communications with consumers or with governmental or consumer protection organizations.

VIII.

IT IS FURTHER ORDERED that respondent Lornamead, Inc., and its successors and assigns shall deliver a copy of this order to all current and, for the next three (3) years, all future principals, officers, directors, and other employees having primary responsibilities with respect to the subject matter of this order, and shall secure from each such person a signed and dated statement acknowledging receipt of the order. Respondent Lornamead, Inc., and its successors and assigns shall deliver this order to current personnel within thirty (30) days after the date of service of this order, and to future personnel within thirty (30) days after the person assumes such position or responsibilities.

IX.

IT IS FURTHER ORDERED that respondent Lornamead, Inc., and its successors and assigns shall notify the Commission at least thirty (30) days prior to any change in the corporation that may affect compliance obligations arising under this order, including but not limited to a dissolution, assignment, sale, merger, or other action that would result in the emergence of a successor corporation; the creation or dissolution of a subsidiary, parent, or affiliate that engages in any acts or practices subject to this order; the proposed filing of a bankruptcy petition; or a change in the corporate name or address. Provided, however, that, with respect to any proposed change in the corporation about which respondent learns less than thirty (30) days prior to the date such action is to take place, respondent shall notify the Commission as soon as is practicable after obtaining such knowledge. Unless otherwise directed by a representative of the Commission, all notices required by this Part shall be sent by overnight courier (not the U.S. Postal Service) to the Associate Director of Enforcement, Bureau of Consumer Protection, Federal Trade Commission, 600 Pennsylvania Avenue NW, Washington, DC 20580, with the subject line: In the Matter of Lornamead, Inc., FTC File Number 122-3255. Provided, however, that, in lieu of overnight courier, notices may be sent by first-class mail, but only if an electronic version of such notices is contemporaneously sent to the Commission at Debrief@ftc.gov.

X.

IT IS FURTHER ORDERED that respondent Lornamead, Inc., and its successors and assigns shall, within sixty (60) days after the date of service of this order, file with the Commission a true and accurate report, in writing, setting forth in detail the manner and form of its own compliance with this order. Within ten (10) days of receipt of written notice from a representative of the Commission, they shall submit additional true and accurate written reports.
XI.

This order will terminate on September 16, 2034, 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:

A. Any Part in this order that terminates in less than twenty (20) years;

B. This order’s application to any respondent that is not named as a defendant in such complaint; and

C. This order if such complaint is filed after the order has terminated pursuant to this Part.

Provided, further, that if such complaint is dismissed or a federal court rules that the respondent did not violate any provision of the order, and the dismissal or ruling is either not appealed or upheld on appeal, then the order will terminate according to this Part as though the complaint had never been filed, except that the order will not terminate between the date such complaint is filed and the later of the deadline for appealing such dismissal or ruling and the date such dismissal or ruling is upheld on appeal.

By the Commission, Commissioner McSweeny not participating.

Donald S. Clark  
Secretary

SEAL:  
ISSUED: September 16, 2014