2023080

**UNITED STATES OF AMERICA**
**BEFORE THE FEDERAL TRADE COMMISSION**

**COMMISSIONERS:** **Rebecca Kelly Slaughter, Acting Chair**
**Noah Joshua Phillips**
**Rohit Chopra**
**Christine S. Wilson**

| | |
|---|---|
| **In the Matter of**<br><br>**CBD MEDS, INC., a corporation,**<br><br>**G2 HEMP, INC., a corporation, and**<br><br>**LAWRENCE MOSES, a/k/a LAWRENCE D. MOSES, JR.,**<br>**individually and as an officer of**<br>**CBD MEDS, INC. and G2 HEMP, INC.** | **DECISION AND ORDER**<br><br>**DOCKET NO. C-4735** |

## DECISION

The Federal Trade Commission ("Commission") initiated an investigation of certain acts and practices of the Respondents named in the caption. The Commission's Bureau of Consumer Protection ("BCP") prepared and furnished to Respondents a draft Complaint. BCP proposed to present the draft Complaint to the Commission for its consideration. If issued by the Commission, the draft Complaint would charge the Respondents with violations of the Federal Trade Commission Act.

Respondents and BCP thereafter executed an Agreement Containing Consent Order ("Consent Agreement"). The Consent Agreement includes: 1) statements by Respondents that they neither admit nor deny any of the allegations in the Complaint, except as specifically stated in this Decision and Order, and that only for purposes of this action, they admit the facts necessary to establish jurisdiction; and 2) waivers and other provisions as required by the Commission's Rules.

The Commission considered the matter and determined that it had reason to believe that Respondents have violated the Federal Trade Commission Act, and that a Complaint should issue stating its charges in that respect. The Commission accepted the executed Consent Agreement and placed it on the public record for a period of 30 days for the receipt and

consideration of public comments. The Commission duly considered any comments received from interested persons pursuant to Section 2.34 of its Rules, 16 C.F.R. § 2.34. Now, in further conformity with the procedure prescribed in Rule 2.34, the Commission issues its Complaint, makes the following Findings, and issues the following Order:

## Findings

1. The Respondents are:

   a. Respondent CBD Meds, Inc., ("CBD Meds") is a California nonprofit mutual benefit corporation. Pursuant to California law, a nonprofit mutual benefit corporation is set up for the benefit of its members and may conduct business at a profit. Cal. Corp. Code §§ 7110 cmt., 7140(1). Thus, CBD Meds is organized to carry on business for its own profit or the profit of its members within the meaning of Section 4 of the FTC Act. 15 U.S.C. § 44. Its principal office or place of business is in Winchester, California 92596.

   b. Respondent G2 Hemp, Inc. ("G2 Hemp") is a California corporation. At times relevant to this Complaint, G2 Hemp operated a website that advertised and sold cannabidiol products. Its principal office or place of business is in Winchester, California 92596.

   c. Respondent Lawrence Moses, also known as Lawrence D. Moses, Jr., is the owner and CEO of CBD Meds and G2 Hemp. Individually or in concert with others, he controlled or had the authority to control, or participated in the acts and practices of CBD Meds and G2 Hemp, including the acts and practices alleged in the Complaint. His principal office or place of business is the same as that of CBD Meds and G2 Hemp.

2. The Commission has jurisdiction over the subject matter of this proceeding and over the Respondents, and the proceeding is in the public interest.

## ORDER

### Definitions

For purposes of this Order, the following definitions apply:

A. "**CBD Product**" means any Dietary Supplement, Food, or Drug containing cannabidiol.

B. "**Covered Product**" means any Dietary Supplement, Food, or Drug, including but not limited to CBD Products.

C. "**Dietary Supplement**" means: (1) any product labeled as a dietary supplement or otherwise represented as a dietary supplement; or (2) any pill, tablet, capsule, powder, softgel, gelcap, liquid, or other similar form containing one or more ingredients that are a vitamin, mineral, herb or other botanical, amino acid, probiotic, or other dietary substance for use by humans to supplement the diet by increasing the total dietary intake, or a concentrate, metabolite, constituent, extract, or combination of any ingredient described above, that is intended to be ingested, and is not represented to be used as a conventional food or as a sole item of a meal or the diet.

D. "**Drug**" means: (1) articles recognized in the official United States Pharmacopoeia, official Homoeopathic Pharmacopoeia of the United States, or official National Formulary, or any supplement to any of them; (2) articles intended for use in the diagnosis, cure, mitigation, treatment, or prevention of disease in humans or other animals; (3) articles (other than food) intended to affect the structure or any function of the body of humans or other animals; and (4) articles intended for use as a component of any article specified in (1), (2), or (3); but does not include devices or their components, parts, or accessories.

E. "**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 indicates that the amount and combination of additional ingredients is unlikely to impede or inhibit the effectiveness of the ingredients in the Essentially Equivalent Product.

F. "**Food**" means: (1) any article used for food or drink for humans or other animals; (2) chewing gum; and (3) any article used for components of any such article.

G. "**Respondents**" means all of the Corporate Respondents and the Individual Respondent, individually, collectively, or in any combination.

1. "**Corporate Respondents**" means CBD Meds, Inc., a corporation, and G2 Hemp, Inc., a corporation, and their successors and assigns.

2. "**Individual Respondent**" means Lawrence Moses, a/k/a Lawrence D. Moses, Jr.

## Provisions

## I. Prohibited Representations: Regarding Health-Related Claims Requiring Human Clinical Testing For Substantiation

**IT IS ORDERED** that Respondents, Respondents' officers, agents, employees, and attorneys, and all other persons in active concert or participation with any of them, who receive actual notice of this Order, whether acting directly or indirectly, in connection with the manufacturing, labeling, advertising, promotion, offering for sale, sale, or distribution of any Covered Product, must not make, or assist others in making, expressly or by implication, any representation that such product:

A. treats blood pressure conditions or gastrointestinal disorders; reduces seizures and convulsions; or reduces blood sugar levels; or

B. cures, mitigates, or treats any disease, including but not limited to cancer, age-related bone disease, arthritis, diabetes, glaucoma, strokes, Alzheimer's disease, multiple sclerosis, Parkinson's disease, epilepsy, autism, post traumatic stress disorder, bipolar disorders, schizophrenia, psoriasis, or HIV dementia,

unless the representation is non-misleading, and, at the time of making such representation, they possess and rely upon competent and reliable scientific evidence substantiating that the representation is true. For purposes of this Section, competent and reliable scientific evidence must consist of human clinical testing of the Covered Product, or of an Essentially Equivalent Product, that is sufficient in quality and quantity based on standards generally accepted by experts in the relevant disease, condition, or function to which the representation relates, when considered in light of the entire body of relevant and reliable scientific evidence, to substantiate that the representation is true. Such testing must be: (1) randomized, double-blind, and placebo-controlled; and (2) conducted by researchers qualified by training and experience to conduct such testing. In addition, all underlying or supporting data and documents generally accepted by experts in the field as relevant to an assessment of such testing as described in the Section entitled Preservation of Records Relating to Competent and Reliable Human Clinical Tests or Studies must be available for inspection and production to the Commission. Persons covered by this Section have the burden of proving that a product satisfies the definition of Essentially Equivalent Product.

## II. Prohibited Representations: Other Health-Related Claims

**IT IS FURTHER ORDERED** that Respondents, Respondents' officers, agents, employees, and attorneys, and all other persons in active concert or participation with any of them, who receive actual notice of this Order, whether acting directly or indirectly, in connection with the manufacturing, labeling, advertising, promotion, offering for sale, sale, or distribution of any Covered Product, must not make, or assist others in making, expressly or by implication, any representation, other than representations covered under the Section of this Order entitled Prohibited Representations: Regarding Health-Related Claims Requiring Human Clinical Testing For Substantiation, about the health benefits, performance, efficacy, safety, or side effects of any Covered Product, including that such product prevents artery blockage, dementia, seizures and

convulsions, cancer, age-related bone disease, arthritis, blood pressure conditions, diabetes, gastrointestinal disorders, glaucoma, Alzheimer's disease, multiple sclerosis, Parkinson's disease, epilepsy, autism, post traumatic stress disorder, bipolar disorders, or schizophrenia, unless the representation is non-misleading, and, at the time of making such representation, they possess and rely upon competent and reliable scientific evidence that is sufficient in quality and quantity based on standards generally accepted by experts in the relevant disease, condition, or function to which the representation relates, 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 Section, competent and reliable scientific evidence means tests, analyses, research, or studies (1) that have been conducted and evaluated in an objective manner by experts in the relevant disease, condition, or function to which the representation relates; (2) that are generally accepted by such experts to yield accurate and reliable results; and (3) that are randomized, double-blind, and placebo-controlled human clinical testing of the Covered Product, or of an Essentially Equivalent Product, when such experts would generally require such human clinical testing to substantiate that the representation is true. In addition, when such tests or studies are human clinical tests or studies, all underlying or supporting data and documents generally accepted by experts in the field as relevant to an assessment of such testing as set forth in the Section entitled Preservation of Records Relating to Competent and Reliable Human Clinical Tests or Studies must be available for inspection and production to the Commission. Persons covered by this Section have the burden of proving that a product satisfies the definition of Essentially Equivalent Product.

## III. Preservation of Records Relating to Competent and Reliable Human Clinical Tests or Studies

**IT IS FURTHER ORDERED** that, with regard to any human clinical test or study ("test") upon which Respondents rely to substantiate any claim covered by this Order, Respondents must secure and preserve all underlying or supporting data and documents generally accepted by experts in the field as relevant to an assessment of the test, including:

A. All protocols and protocol amendments, reports, articles, write-ups, or other accounts of the results of the test, and drafts of such documents reviewed by the test sponsor or any other person not employed by the research entity;

B. All documents referring or relating to recruitment; randomization; instructions, including oral instructions, to participants; and participant compliance;

C. Documents sufficient to identify all test participants, including any participants who did not complete the test, and all communications with any participants relating to the test; all raw data collected from participants enrolled in the test, including any participants who did not complete the test; source documents for such data; any data dictionaries; and any case report forms;

D. All documents referring or relating to any statistical analysis of any test data, including any pretest analysis, intent-to-treat analysis, or between-group analysis performed on any test data; and

E. All documents referring or relating to the sponsorship of the test, including all communications and contracts between any sponsor and the test's researchers.

*Provided*, *however*, the preceding preservation requirement does not apply to a reliably reported test, unless the test was conducted, controlled, or sponsored, in whole or in part by (1) any Respondent; (2) any Respondent's officers, agents, representatives, or employees; (3) any other person or entity in active concert or participation with any Respondent; (4) any person or entity affiliated with or acting on behalf of any Respondent; (5) any supplier of any ingredient contained in the product at issue to any of the foregoing or to the product's manufacturer; or (6) the supplier or manufacturer of such product.

For purposes of this Provision, "reliably reported test" means a report of the test has been published in a peer-reviewed journal, and such published report provides sufficient information about the test for experts in the relevant field to assess the reliability of the results.

For any test conducted, controlled, or sponsored, in whole or in part, by Respondents, Respondents must establish and maintain reasonable procedures to protect the confidentiality, security, and integrity of any personal information collected from or about participants. These procedures must be documented in writing and must contain administrative, technical, and physical safeguards appropriate to Respondents' size and complexity, the nature and scope of Respondents' activities, and the sensitivity of the personal information collected from or about the participants.

## IV. Prohibited Misrepresentations Regarding Tests, Studies, or Other Research

**IT IS FURTHER ORDERED** that Respondents, Respondents' officers, agents, employees, and attorneys, and all other persons in active concert or participation with any of them, who receive actual notice of this Order, whether acting directly or indirectly, in connection with the manufacturing, labeling, advertising, promotion, offering for sale, sale, or distribution of any product must not misrepresent, in any manner, expressly or by implication:

A. that any Covered Product is scientifically proven to prevent seizures; treat cancer; treat or prevent strokes, Alzheimer's disease, Parkinson's disease, or HIV dementia; or make chemotherapy more effective and increase cancer cell death without harming normal cells;

B. that the performance or benefits of any product are scientifically or clinically proven or otherwise established;

C. the existence, contents, validity, results, conclusions, or interpretations of any test, study, or other research;

D. that a U.S. government laboratory study showed that any Covered Product may make chemotherapy more effective and increase cancer cell death without harming normal cells; or

E. that the U.S. government has stated that any Covered Product is scientifically proven to have antioxidant and neuroprotectant properties, limit neurological damage following ischemic insults, such as stroke and trauma, and treat neurogenerative diseases, such as Alzheimer's disease, Parkinson's disease and HIV dementia.

## V. FDA Approved Claims

**IT IS FURTHER ORDERED** that nothing in this Order prohibits Respondents, Respondents' officers, agents, employees, and attorneys, or all other persons in active concert or participation with any of them, from:

A. for any Drug, making a representation that is approved in labeling for such Drug under any tentative final or final monograph promulgated by the Food and Drug Administration ("FDA"), or under any new drug application approved by the FDA; and

B. for any product, making a representation that is specifically authorized for use in labeling for such product by regulations promulgated by the FDA pursuant to the Nutrition Labeling and Education Act of 1990 or permitted under Sections 303-304 of the Food and Drug Administration Modernization Act of 1997.

## VI. Notices to Customers

**IT IS FURTHER ORDERED** that Respondents must notify customers as follows:

A. Respondents must identify all consumers who purchased the CBD Products on or after January 9, 2017 and through the Order's effective date ("eligible customers").

   1. Such eligible customers, and their contact information, must be identified to the extent such information is in Respondents' possession, custody or control;

   2. Eligible customers include those identified at any time including after Respondents' execution of the Agreement through the eligibility period, which runs for 1 year after the issuance date of the Order.

B. Respondents must notify all identified eligible customers by mailing each a notice:

   1. The letter must be in the form shown in Attachment A.

   2. The envelope containing the letter must be in the form shown in Attachment B.

   3. The mailing of the notification letter must not include any other enclosures.

4. The mailing must be sent by first-class mail, postage prepaid, address correction service requested with forwarding and return postage guaranteed. For any mailings returned as undeliverable, Respondents must use standard address search methodologies such as re-checking Respondents' records and the Postal Service's National Change of Address database and re-mailing to the corrected address within 8 days.

C. Respondents must notify all eligible customers within 180 days after the issuance date of this Order and any eligible customers identified thereafter within 30 days of their identification.

D. Respondents must provide a notice on their websites' landing pages. Such notice must link to a copy of the Order. The notice must be posted not later than 3 days after the effective date of the Order and for at least 1 year after the Order's effective date.

E. Respondents must report on their notification program under penalty of perjury:

1. Respondents must submit a report at the conclusion of the notification program summarizing their compliance, including the total number of notices sent or re-sent, the dates sent, and eligible customers identified.

2. If a representative of the Commission requests any information regarding the program, including any of the underlying customer data, Respondents must submit it within 10 days of the request.

3. Failure to provide required notices or any requested information will be treated as a continuing failure to obey this Order.

## VII. Acknowledgments of the Order

**IT IS FURTHER ORDERED** that Respondents obtain acknowledgments of receipt of this Order:

A. Each Respondent, within 10 days after the effective date of this Order, must submit to the Commission an acknowledgment of receipt of this Order sworn under penalty of perjury.

B. For 20 years after the issuance date of this Order, the Individual Respondent for any business that such Respondent, individually or collectively with any other Respondents, is the majority owner or controls directly or indirectly, unless such business cannot violate the Order, and each Corporate Respondent, must deliver a copy of this Order to: (1) all principals, officers, directors, and LLC managers and members; (2) all employees having managerial responsibilities for conduct related to the subject matter of the Order and all agents and representatives who participate in conduct related to the subject matter of the Order; and (3) any business entity resulting from any change in structure as set forth in the Provision titled Compliance Report and Notices. Delivery must occur within 10 days after the effective date of this Order for current personnel. For all others,

delivery must occur before they assume their responsibilities.

C. From each individual or entity to which a Respondent delivered a copy of this Order, that Respondent must obtain, within 30 days, a signed and dated acknowledgment of receipt of this Order.

## VIII. Compliance Report and Notices

**IT IS FURTHER ORDERED** that Respondents make timely submissions to the Commission:

A. Sixty days after the issuance date of this Order, each Respondent must submit a compliance report, sworn under penalty of perjury, in which:

1. Each Respondent must: (a) identify the primary physical, postal, and email address and telephone number, as designated points of contact, which representatives of the Commission, may use to communicate with Respondent; (b) identify all of that Respondent's businesses by all of their names, telephone numbers, and physical, postal, email, and Internet addresses; (c) describe the activities of each business, including the goods and services offered, the means of advertising, marketing, and sales, and the involvement of any other Respondent (which Individual Respondents must describe if they know or should know due to their own involvement); (d) describe in detail whether and how that Respondent is in compliance with each Provision of this Order, including a discussion of all of the changes the Respondent made to comply with the Order; and (e) provide a copy of each Acknowledgment of the Order obtained pursuant to this Order, unless previously submitted to the Commission.

2. Additionally, the Individual Respondent must: (a) identify all his telephone numbers and all his physical, postal, email and Internet addresses, including all residences; (b) identify all his business activities, including any business for which such Respondent performs services whether as an employee or otherwise and any entity in which such Respondent has any ownership interest; and (c) describe in detail such Respondent's involvement in each such business activity, including title, role, responsibilities, participation, authority, control, and any ownership.

B. Each Respondent must submit a compliance notice, sworn under penalty of perjury, within 14 days of any change in the following:

1. Each Respondent must submit notice of any change in: (a) any designated point of contact; or (b) the structure of any Corporate Respondent or any entity that Respondent has any ownership interest in or controls directly or indirectly that may affect compliance obligations arising under this Order, including: creation, merger, sale, or dissolution of the entity or any subsidiary, parent, or affiliate that engages in any acts or practices subject to this Order.

2. Additionally, the Individual Respondent must submit notice of any change in: (a) name, including alias or fictitious name, or residence address; or (b) title or role in any business activity, including (i) any business for which such Respondent performs services whether as an employee or otherwise and (ii) any entity in which such Respondent has any ownership interest and over which Respondents have direct or indirect control. For each such business activity, also identify its name, physical address, and any Internet address.

C. Each Respondent must submit notice of the filing of any bankruptcy petition, insolvency proceeding, or similar proceeding by or against such Respondent within 14 days of its filing.

D. Any submission to the Commission required by this Order to be sworn under penalty of perjury must be true and accurate and comply with 28 U.S.C. § 1746, such as by concluding: "I declare under penalty of perjury under the laws of the United States of America that the foregoing is true and correct. Executed on: _____" and supplying the date, signatory's full name, title (if applicable), and signature.

E. Unless otherwise directed by a Commission representative in writing, all submissions to the Commission pursuant to this Order must be emailed to DEbrief@ftc.gov or sent by overnight courier (not the U.S. Postal Service) to: Associate Director for Enforcement, Bureau of Consumer Protection, Federal Trade Commission, 600 Pennsylvania Avenue NW, Washington, DC 20580. The subject line must begin: In re CBD Meds, Inc.

## IX. Recordkeeping

**IT IS FURTHER ORDERED** that Respondents must create certain records for 20 years after the issuance date of the Order, and retain each such record for 5 years, unless otherwise specified below. Specifically, Corporate Respondents and the Individual Respondent for any business that such Respondent, individually or collectively with any other Respondents, is a majority owner or controls directly or indirectly, must create and retain the following records:

A. accounting records showing the revenues from all goods or services sold, the costs incurred in generating those revenues, and resulting net profit or loss;

B. personnel records showing, for each person providing services in relation to any aspect of the Order, whether as an employee or otherwise, that person's: name; addresses; telephone numbers; job title or position; dates of service; and (if applicable) the reason for termination;

C. copies or records of all consumer complaints and refund requests concerning the subject matter of the Order, whether received directly or indirectly, such as through a third party, and any response;

D. a copy of each unique advertisement or other marketing material making a representation

subject to this Order.

E. For 5 years from the date of the last dissemination of any representation covered by this Order:

1. all materials that were relied upon in making the representation; and

2. all tests, studies, analysis, other research or other such evidence in Respondent's possession, custody, or control that contradicts, qualifies, or otherwise calls 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.

F. for 5 years from the date received, copies of all subpoenas and other communications with law enforcement, if such communication relate to Respondents' compliance with this Order.

G. all records necessary to demonstrate full compliance with each provision of this Order, including all submissions to the Commission.

## X. Compliance Monitoring

**IT IS FURTHER ORDERED** that, for the purpose of monitoring Respondents' compliance with this Order:

A. Within 10 days of receipt of a written request from a representative of the Commission, each Respondent must: submit additional compliance reports or other requested information, which must be sworn under penalty of perjury, and produce records for inspection and copying.

B. For matters concerning this Order, representatives of the Commission are authorized to communicate directly with each Respondent. Respondents must permit representatives of the Commission to interview anyone affiliated with any Respondent who has agreed to such an interview. The interviewee may have counsel present.

C. The Commission may use all other lawful means, including posing through its representatives as consumers, suppliers, or other individuals or entities, to Respondents or any individual or entity affiliated with Respondents, without the necessity of identification or prior notice. Nothing in this Order limits the Commission's lawful use of compulsory process, pursuant to Sections 9 and 20 of the FTC Act, 15 U.S.C. §§ 49, 57b-1.

D. Upon written request from a representative of the Commission, any consumer reporting agency must furnish consumer reports concerning Individual Respondents, pursuant to Section 604(2) of the Fair Credit Reporting Act, 15 U.S.C. § 1681b(a)(2).

## XI. Order Effective Dates

**IT IS FURTHER ORDERED** that this Order is final and effective upon the date of its publication on the Commission's website (ftc.gov) as a final order. This Order will terminate 20 years from the date of its issuance (which date may be stated at the end of this Order, near the Commission's seal), or 20 years from the most recent date that the United States or the Commission files a complaint (with or without an accompanying settlement) in federal court alleging any violation of this Order, whichever comes later; *provided, however,* that the filing of such a complaint will not affect the duration of:

A. Any Provision in this Order that terminates in less than 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 Provision.

*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 Provision 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.

April J. Tabor
Secretary

SEAL:
ISSUED: February 2, 2021

**ATTACHMENT A TO THE ORDER**

CLAIMS ABOUT PRODUCTS CONTAINING CBD
*In the Matter of CBD Meds, Inc., et al.*

<Date>

<Name of customer>
<mailing address of customer
including zip code>

Subject: CBD Products sold by CBD Meds and G2 Hemp

Dear <Name of customer>:

Our records show that you bought CBD products from CBD Meds and G2 Hemp. We are writing to tell you that the Federal Trade Commission, (FTC), the nation's consumer protection agency, has charged us with deceptive or false advertising.

The FTC brought a lawsuit against our companies for making misleading claims that our CBD products can effectively prevent, treat, or ease serious diseases or health conditions, including the following:

> Artery blockage; dementia; blood sugar levels; seizures and convulsions; psoriasis; HIV dementia; cancer; age-related bone disease; arthritis; blood pressure conditions; diabetes; gastrointestinal disorders; glaucoma; strokes; Alzheimer's disease; multiple sclerosis; Parkinson's disease; epilepsy; autism; post traumatic stress disorder; bipolar disorders; and schizophrenia.

To settle the FTC's lawsuit, we're contacting our customers to tell them that we don't have proof that our CBD products will effectively prevent, treat, or improve the serious diseases and health conditions listed above. In addition, the U.S. government has not validated those claims.

If you have other questions about this lawsuit, visit [add URL].

CBD oil and other alternative treatments might be harmful to your medical care, and could interfere with your prescriptions. CBD products could also be dangerous if you take them with other medicines or at a high dose. Talk to your doctor before you take any treatments or stop any prescriptions. For more information about protecting yourself from bogus health product claims visit ftc.gov/health.

Sincerely,

[signature]

Lawrence Moses
CEO, CBD Meds, Inc. and G2 Hemp, Inc.

**ATTACHMENT B to the Order – Envelope Template:**

The envelope for the notification letter must be in the following form, with the underlined text completed as directed:

*CBD MEDS, INC. AND G2 HEMP, INC.*
*Street Address*
*City, State and Zip Code*

FORWARDING AND RETURN POSTAGE GUARANTEED ADDRESS CORRECTION SERVICE REQUESTED

[name and
mailing address of customer,
including zip code]

**ABOUT YOUR PURCHASE FROM CBD MEDS, INC. AND G2 HEMP, INC.**