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                    UNITED STATES DISTRICT COURT
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                   CENTRAL DISTRICT OF CALIFORNIA
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   FEDERAL TRADE COMMISSION,
                                       ) Case No.: 2:20-cy-3775
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                         Plaintiff,
                                       ) [PROPOSED] STIPULATED
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                    v.
                                       ) PRELIMINARY INJUNCTION
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                                       ) PURSUANT TO SECTIONS 13(a)
                                       ) AND (b) OF THE FEDERAL
   MARC CHING, individually and also
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                                         TRADE COMMISSION ACT
   doing business as WHOLE LEAF
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   ORGANICS,
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                         Defendant.
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5. No security is required of any agency of the United States for issuance of a preliminary injunction. Fed. R. Civ. P. 65(c).

DEFINITIONS

For purposes of this Order, the following definitions apply:

- A. "Covered Product" means Thrive, CBEX, CBD-RX, or CBD-Max or any other Drug, Food, or Dietary Supplement.
- B. "Dietary Supplement" means:

- 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, cibuent, 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.
- C. "Drug" means: (a) 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; (b) articles
 intended for use in the diagnosis, cure, mitigation, treatment, or prevention of
 disease in humans or other animals; (c) articles (other than Food) intended to affect
 the structure or any function of the body of humans or other animals; and (d)
 articles intended for use as a component of any article specified in (a), (b), or (c);
 but does not in ded devices or their components, parts, or accessories.
- D. "Essentially equivalent product" means a product that contains the identical ingredients, except for inactive ingredients (e.g., inactive binders, colors, fillers,

relates; (2) that are generally accepted by such experts to yield accurate and reliable results; and (3)hat are randomized, doubthind, and placeboontrolled 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 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 described in the Provision of this Order titled Preservation of Records Relating to Competent and Reliable Human Clinical Tests or Studies must be available for inspection and production to the Commission. Defending have the burden of proving that a product satisfies the definition of an Essentially Equivalent Product.

III. PROHIBITED MISREPRESENTATIONS REGARDING TESTS,
STUDIES, OR OTHER RESEARCH
IT IS FURTHER ORDERED that Defendantid Defendant's agents,

IV. FDA APPROVED CLAIMS

IT IS FURTHER ORDERED that nothing in this Order prohibits Defendant, or Defendant's 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 (3 (inis)88.5 (on 20g A)o8c0d Drll ou(ve)3.6d la teuodla s ine (

obtained in connection with any activity that pertains to the subject matter of this Order.

Provided, however, that Defendantay disclose such identifying information to a law enforcement agencyhtsattorneys as required fbis defense this or the pending administrative action, as required by any law, regulation, or court order, or in any filings, pleadings or discovery in this action in the manner required by the Feedl Rules of Civil Procedure and by any protective order in the case.

VI. PRESERVATION OF RECORDS

IT IS FURTHER ORDERED that Defendant, Defendant ents, 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 o(e)3.6 (c)3.5 <68.5 (pa)12wJ 0 -1i1 0 Td ()Tj 0.0rctis Order, whe0 T3 (e)3.6 (tua)12.0

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person (including any financial institution) that may have possessescustody or control of anydocument of Defendant, or that may be subject to any provision of this Order pursuant to Rule 65(d)(2) of the Federal Rules of Civil Procedure. For purposes of this Section, service upon any branch, subsidiary, affiliate or office of any entity shall effect service upon the entire entity.

VIII. CORRESPONDENCE AND SERVICE ON PLAINTIFF
IT IS FURTHER ORDERED that, for the purpose of this Order, all
correspondence and service of pleadings on Plaintiff shall beineenthail to:

TAWANA E. DAVIS tdavis@ftc.gov; (202) 32@755 AMBER LEE alee5@ftc.go,v(202) 3262764 Federal Trade Commission 600 Pennsylvania Avenue, NW Washington, DC 20580 Fax: (202) 326-3259

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1	X. RETENTION OF JURISDICTION		
2	IT IS FURTHER ORDERED that this Court shall retain jurisdiction of this		
3	matter for all purposes.		
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5	SO ORDERED, this da	y of, 2020, atm	
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7	UNITED STATES DISTRICT JUDGE		
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