1 2 3 4 5	FARNESE P.C. Peter J. Farnese (SBN 251204) 700 S. Flower St., Suite 1000 Los Angeles, California 90017 Telephone: 310-356-4668 Facsimile: 310-388-1232 Email: pif@farneselaw.com	Clerk of the Superior Court OCT 12 2023 By D. HER. 2 DEPUTY CLERK
5, 6 7, 8	Attorneys for Plaintiff	ASSIGNED TO JUDGE ALESIA JONES 10 12 22 23 FOR ALL PURPOSES
9	SUPERIOR COURT FOR TH	IE STATE OF CALIFORNIA Judge Kafe 1-1-24 Following
	FOR THE COUN	TY OF SOLANO
10 11	DANIELLE SKARPNES, individually and on	CASE NO. C U 2 3 - 0 4 6 3 8
12	behalf of all others similarly situated,	CLASS ACTION
13	Plaintiffs,	COMPLAINT FOR:
14 15	v.	1. VIOLATION OF CAL. BUS. & PROF. § 17200, ET SEQ. (Unlawful, Unfair, and
16	ELIXIR COSMETICS OPCO, LLC.	Fraudulent Prongs)
17	Defendants.	2. VIOLATION OF CAL. BUS. & PROF. § 17500, ET SEQ. (False and Misleading Advertising)
18		3. VIOLATION OF CAL. CIV. CODE §
19 20		1750, ET SEQ. (Consumers Legal Remedies Act)
21		4. UNJUST ENRICHMENT
22		5. BREACH OF WARRANTY
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24		JURY TRIAL DEMANDED
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Farnese PC	CLASS ACTION	N COMPLAINT

Plaintiff Danielle Skarpnes (hereafter "Plaintiff"), individually and on behalf of all other similarly situated purchasers (hereafter the "Class") of the Babe Lash Essential, Lash Serum, Babe Lash Eyelash Serum, Babe Amplifying Brow Serum and Babe Brow Serum (collectively hereafter the "Product(s)"), brings this consumer class action against Elixir Cosmetics OPCO, LLC (hereafter "Defendant") and alleges as follows:

JURISDICTION AND VENUE

- 1. This Court has jurisdiction over all causes of action asserted herein pursuant to the California Constitution, Article VI, § 10, as this case is a cause not given by statute to other trial courts.
- 2. Defendant has sufficient minimum contacts with California, and/or otherwise has intentionally availed itself of the markets in California through the manufacture, promotion, marketing, and sale of their products in California. Moreover, Defendant, directly and through its agents, has substantial contacts with and receive benefits and income from and through the State of California such that the exercise of jurisdiction by this Court is permissible under traditional notions of fair play and substantial justice.
- 3. Defendant made numerous misrepresentations and material omissions which had a substantial effect in Solano County, including, but not limited to internet advertisements as well as sales in retail stores located in Solano County. Additionally, Defendant received considerable compensation from sales in Solano County.
- 4. Venue is proper in this Court as Plaintiff at all relevant times is and was a resident of Solano County, California and Plaintiff's purchase of the Products occurred in Solano County, California.
- 5. Out-of-state participants can be brought before this Court pursuant to the provisions of Code of Civil Procedure § 395.5.

THE PARTIES

6. Plaintiff is an individual residing in the State of California. Plaintiff purchased the Babe Lash Essential Lash Serum from a Costco retail store located in Vacaville, California in 2023 for \$43.99 plus tax, for personal use. In doing so, Plaintiff relied upon Defendant's advertising,

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packaging, labeling and other promotional materials, which were jointly prepared and approved by Defendant and its agents and disseminated through advertising media, social media, Defendant's website, and the internet, containing the misrepresentations, concealments alleged herein. Plaintiff would not have purchased the Product, nor would she have paid the price that she did, had she known that it contained a prostaglandin associated with a variety of side effects and health risks, and is sold in California in violation of California's Sherman Food Drug and Cosmetic Law.

- 7. Defendant Elixir is a limited liability company with its principal place of business in Texas. At times relevant to this Complaint, Defendant has advertised, marketed, and sold a variety of cosmetic products, including the Products at issue, to consumers throughout the United States and the State of California. Defendant has sold the Products directly to consumers via the Internet through its own website, as well as third party retail stores throughout the United States, including in this County.
- 8. Defendant transacts or has transacted business in this County and throughout the United States. Defendant, directly and through its agents, have substantial contacts with and receives substantial benefits and income from and through the State of California and this County.

FACTUAL ALLEGATIONS

Background

- 9. Women have long strived to possess long, thick, and dark eyelashes. Prominent eyes and eyelashes are often considered a sign of beauty and can be associated with increased levels of attractiveness, confidence, and well-being.
- 10. Numerous options may improve the appearance of eyelashes. In fact, women spend over a billion dollars a year on products that enhance the appearance of eyelashes. They coat them with mascara to make them look thicker, glue on fake eyelashes and extensions, tint them with dye to make them look darker, and even have hairs transplanted to their lids.
- 11. Since approximately 2007, the new craze in the eyelash industry has been eyelash enhancing serums that claim to cause a user's eyelashes to grow longer, thicker, and darker.
- 12. The active ingredient contained in various eyelash enhancing serums was originally discovered by mistake. Lumigan eye drops, used to treat glaucoma, contain the active ingredient

bimatoprost which is in a category of compounds known as prostamides. Prostamides are related to a category of compounds known as prostaglandins ("PGs"). Eyelash growth was found to be an unexpected, positive side effect of Lumigan, due to the PGs contained therein. However, adverse side effects were also found to be associated with PGs. These adverse side effects include but are not limited to: ocular irritation, conjunctival hyperemia, iris color change, eyelid pigmentation, redness, macular edema, ocular inflammation, unwanted hair growth, eye pruritus, blurred vision and blindness.

- 13. The manufacturer of Lumigan, Allergan Inc., took the information regarding PGs and its effectiveness to grow eyelashes and created Latisse, an eyelash enhancing serum containing bimatoprost.
- 14. The Federal Food, Drug and Cosmetic Act ("FDCA") defines drugs by their intended use, as "articles intended for use in the diagnosis, cure, mitigation, treatment, or prevention of disease" and "articles (other than food) intended to affect the structure or any function of the body of man or other animals." (FD&C Act, sec 201(g)(1)). Under the FDCA, drugs must either receive premarket approval by FDA through the New Drug Application (NDA) process, or conform to a monograph for a particular drug category (as established by the FDA's Over-the-Counter Drug Review), before they go on the market. California's Health and Safety Code mirrors the FDCA.
- 15. Latisse is intended to affect the structure of the human body by elongating and thickening eyelashes, and consequently, is considered a drug for purposes of the FDA. Accordingly, Allergan needed to receive FDA approval in order to put Latisse on the market. Because there was no monograph for eyelash serums, Allergan was required to receive premarket approval by the FDA through the NDA process.
- 16. Following Allergan's NDA process and based on Latisse's documented safety record and proof that bimatoprost .03% grows eyelashes when applied to eyelid skin, in December of 2008, Latisse became the first (and to date remains the only) federally approved eyelash enhancing serum.
- 17. Latisse is not available over-the-counter and must be prescribed to customers through a physician.
- 18. In addition to requiring a prescription after consultation with a physician, Allergan warns of the risks of prostaglandins. The Latisse warnings and precautions advise of effects on:

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intraocular pressure, iris pigmentation changes, lid pigmentation, hair growth outside the treatment area, intraocular inflammation, and macular edema.

- 19. Since 2007, a variety of companies sought to capitalize on the market created by Latisse by selling eyelash enhancing serums, also containing prostaglandin analogs. These companies attempt to take advantage of the fact that Latisse is expensive, requires a doctor's prescription, and advises consumers of potential significant side effects of using the product.
- 20. Thus, they market their products as purported "cosmetics" and not as "drugs", in an attempt to circumvent the rigorous process required to seek regulatory approval. Further, these companies sell the serums directly to customers and at retail without a prescription. The ease of purchase for customers leads to greater sales for vendors and is a critical part to their success.
- 21. But, FDA has previously warned manufacturers marketing eyelash serums formulated with ICP that these products violated the FDCA because they are unapproved new drugs and misbranded drugs. FDA also noted the harmful side effects associated with prostaglandin analogs: "[o]ther potential adverse events associated with prostaglandin analogs for ophthalmic use include ocular irritation, hyperemia, iris color change, macular edema, ocular inflammation, and interference with glaucoma therapy. In addition, prostaglandin analogs for ophthalmic use are currently classified as Pregnancy Class C." See U.S. Food and Drug Administration Warning Letter to Lifetech Resources, LLC, dated April 18, 2011.

Defendant's Advertising and Marketing of the Products is False and Deceptive

22. With the expansion of social media platforms, digital marketing channels have lowered barriers to entry into the beauty and personal care market, enabling the creation of digitally native, "direct-to-consumer" cosmetics brands, like Defendant's "Babe Original" brand¹. These brands have disrupted the high-margin cosmetics industry that has long been dominated by large, billion-dollar

Defendant originally marketed the Products under the brand "Babe Lash", however, in or about 2020, transitioned to "Babe Original" as its blanket brand for the Products. The formulations of the products (and intended uses as described below) remained.

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solution), (b)(6) (travoprost ophthalmic solution) and to treat hypotrichosis of the eyelashes (e.g., (b)(6) (bimatoprost ophthalmic solution)). Prostaglandin analogs are well known to have an effect on the structure or function of the body. The presence of the prostaglandin analog, isopropyl cloprostenate, along with appearance claims such as 'enhance the appearance of your lashes and brows,' 'fuller healthier-looking lashes,' and 'fuller healthier-looking brows' indicate that your products are intended to affect the structure or function of the body. Accordingly, [these products] are drugs as defined by section 201 (g)(1)(C) of the Act (21 U.S.C. § 321(g)(I)(C))." See U.S. Food and Drug Administration Warning Letter to Lifetech Resources, LLC, dated April 18, 2011.

The Products Are Sold In Violation of the California Sherman Law

- 40. Defendant has marketed, labeled, and sold unapproved drugs in violation of California's Sherman Food, Drug, and Cosmetic Law, Cal. Health & Safety Code § 109875 et seq. ("Sherman Law") and federal law.
- 41. Under California's Sherman law, a product is a "drug" if it is intended to "affect" "any function" of the "human body". Cal. Health & Safety Code § 109925(c). Federal law is identical. See, e.g., 21 U.S.C. § 321(g)(1)(C). "The drug definition is to be given a liberal interpretation in light of the remedial purposes of the legislation." National Nutritional Foods Ass'n v. Mathews, 557 F.2d 325, 336 (2d Cir.1977) (citing United States v. An Article of Drug... Bacto-Unidisk, 394 U.S. 784, 792, 798 (1968).
- 42. A product's "intended use" is not determined by the seller's subjective explanations. Rather, it is determined by the seller's objective intent as "derived or inferred from labeling, promotional material, advertising, or any other relevant source." Allergan, Inc. v. Athena Cosmetics, Inc., supra at 1357 citing United States v. Storage Spaces Designated Nos. "8" & "49", 777 F.2d 1363, 1366 (9th Cir. 1985).
- 43. The "intended use" of eyelash serums, like the Products, has been extensively litigated in California. Namely, the district court in the Allergan/Athena Cosmetics line of cases identified several factors that are to be considered in making the determination of the "intended use" of various eyelash serums (and in turn, were "drugs" under the California Health & Safety Code and federal law), including: the product name, formulation, marketing, marketing structure, advertising claims,

formulation, method of application, time needed to work, pricing, and reports of consumers' use of the Product, among others. *See, e.g, Allergan, Inc. v. Athena Cosmetics, Inc.*, 8:07-cv-01316-JVS-RNB, Dkt. No. 711 (C.D. Cal. July 19, 2012). These facts are "all considered in light of prevalent claims about making one's lashes 'look' or 'appear' longer, fuller, and more beautiful; statements championing the replacement of extensions and false eyelashes; and statements about combining use with mascara." *Id.* at pp. 14-15 (emphasis added).

44. Here, as described above, the Products' advertising history, formulation, clinical testing, pricing, before and after pictures, time to provide results, use of ICP, method of application, and pricing (ranging between \$966.67 and \$684.62 per fl oz, depending on size), among other things, demonstrate that Defendants' objective intent is to affect the structure or function of the human body with respect to eyelash and eyebrow growth and, accordingly, the Products are unapproved and misbranded drugs under the California Sherman law (and identical federal law).

CLASS ACTION ALLEGATIONS

45. <u>Class Definition</u>: Plaintiff brings this class action on behalf of herself, and as a class action on behalf of the following putative class (the "Class"):

All persons in the United States or its territories who purchased any Elixir Product for personal, family, household, or professional purposes between June 1, 2019 and the date of the entry of an order granting preliminary approval to the Settlement Agreement excluding (a) any individuals who have pending litigation against Elixir; (b) any Settlement Class Members who file a timely request for exclusion; (c) any officers, directors, or employees, or immediate family members of the officers, directors, or employees, of Elixir or any entity in which Elixir has a controlling interest; (d) any person who has acted as a consultant of Elixir; (e) any legal counsel or employee of legal counsel for Elixir; (f) any federal, state, or local government entities; and (g) any judicial officers presiding over the Action and the members of their immediate family and judicial staff.

Plaintiff reserves the right to amend the Class definition if further investigation and discovery indicates that the Class definitions should be narrowed, expanded, or otherwise modified.

46. <u>Numerosity and Ascertainability</u>: Plaintiff does not know the exact number of members of the putative classes. Plaintiff is informed and believes that the total number of Class members is at least in the many tens of thousands, and that members of the Class are numerous and geographically dispersed throughout the United States and California. While the exact number and identities of the

Class members are unknown at this time, such information can be ascertained through appropriate investigation and discovery, including Defendants' records, either manually or through computerized searches.

- 47. <u>Typicality and Adequacy</u>: Plaintiff's claims are typical of those of the proposed Class, and Plaintiff will fairly and adequately represent and protect the interests of the proposed Class. Plaintiff does not have any interests that are antagonistic to those of the proposed Class. Plaintiff has retained counsel competent and experienced in the prosecution of this type of litigation.
- 48. <u>Commonality</u>: The questions of law and fact common to the Class members, some of which are set out below, predominate over any questions affecting only individual Class members:
 - a. whether Defendants claimed that the Products may cause or assist in eyelash growth;
 - b. whether the surrounding facts and circumstances demonstrate that the intended use of the Products is to cause or assist in eyelash and eyebrow growth;
 - c. whether the Products contain ICP;
 - d. whether Defendant's advertising and labeling, along with the concealment of certain information regarding Defendant's Products, is material to consumers;
 - e. whether Defendant's advertising and marketing claims set forth above are unlawful, untrue, or are misleading, or reasonably likely to deceive;
 - f. whether Defendant's conduct is fraudulent and/or violates public policy;
 - g. whether Defendant engaged in unfair, unlawful and/or fraudulent business practices in marketing and distributing the Products;
 - h. whether Defendant made express and/or implied warranties for the Products;
 - i. whether the Products are adulterated and/or misbranded under the California Health & Safety Code and identical federal law;
 - j. whether Defendant knew or should have known that the representations were false;
 - k. whether Defendant engaged in false advertising with respect to the Products;
 - 1. whether Defendant knowingly concealed or misrepresented material facts for the purpose of inducing consumers into spending money on the Products;

- m. whether Defendant's representations, concealments and non-disclosures concerning the Products are likely to deceive the consumer;
- n. whether Defendant's representations, concealments and non-disclosures concerning the Products violate the CLRA, FAL, UCL, and/or the common law;
- o. whether Defendants should be permanently enjoined from making the claims at issue; and
- p. whether Plaintiff and the Class are entitled to restitution and damages.
- 49. Predominance and Superiority: Common questions, some of which are set out above, predominate over any questions affecting only individual Class members. A class action is the superior method for the fair and just adjudication of this controversy. The expense and burden of individual suits makes it impossible and impracticable for members of the proposed Class to prosecute their claims individually and multiplies the burden on the judicial system presented by the complex legal and factual issues of this case. Individualized litigation also presents a potential for inconsistent or contradictory judgments. In contrast, the class action device presents far fewer management difficulties and provides the benefits of single adjudication, economy of scale, and comprehensive supervision by a single court on the issue of Defendants' liability. Class treatment of the liability issues will ensure that all claims and claimants are before this Court for consistent adjudication of the liability issues.
- 50. <u>Manageability</u>: The trial and litigation of Plaintiff's and the proposed Class's claims are manageable. Defendants have acted and refused to act on grounds generally applicable to the Class, making appropriate final injunctive relief and declaratory relief with respect to the Class as a whole.
- Notice: If necessary, notice of this action may be affected to the proposed Class through publication in a manner authorized in the California Rules of Court, Civil Code, and/or the Federal Rules of Civil Procedure. Also, Class members may be notified of the pendency of this action by mail and/or email, through the distribution records of Defendant, third party retailers, and vendors.

FIRST CAUSE OF ACTION

VIOLATION OF UNFAIR COMPETITION LAW (CAL. BUS. & PROF. CODE § 17200, et seq.)

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(Unlawful, Unfair, and Fraudulent Prongs of the Act)

- 52. Plaintiff incorporates by this reference the allegations contained in the preceding paragraphs as if fully set forth herein.
- 53. Plaintiff brings this claim individually and on behalf of the proposed Class against Defendants.
- California Business and Professions Code § 17200 prohibits any "any unlawful, unfair 54. or fraudulent business act or practice and unfair, deceptive, untrue or misleading advertising". For the reasons discussed above, Defendants have engaged in unlawful, unfair, and deceptive acts, and untrue and misleading advertising in violation of California Business & Professions Code §17200.
- As alleged herein, Plaintiff has standing to pursue this claim as Plaintiff has suffered 55. injury in fact and has lost money or property as a result of Defendant's actions. Specifically, Plaintiff purchased the Products for her own personal use. In so doing, Plaintiff relied upon the representations referenced above. Plaintiff would not have purchased the Product, nor would she have paid the price that she did, had she known that it contained a prostaglandin associated with a variety of side effects and health risks, and is sold in California in violation of California's Sherman Food Drug and Cosmetic Law as an unlawful drug.
- 56. Unlawful Business Practices: The actions of Defendants, as alleged herein, constitute illegal and unlawful practices committed in violation of the Business & Professions Code §17200.
- 57. In addition, Defendants' have committed unlawful business practices by, inter alia, making the misrepresentations and omissions of material facts, as set forth more fully herein, by violating: (1) sections 1770(a)(5), 1770(a)(7), 1770(a)(8) and 1770(a)(9) of the CLRA, Civil Code § 1750, et seq.; (2) Cal. Business and Professions Code § 17500, et seq.; and (3) sections 111550(a)(1), 111550(b), 111330, 111375, 110389, and 111440 of the California Health & Safety Code.
- 58. Plaintiff and the Class reserve the right to allege other violations of law which constitute other unlawful business acts or practices. Such conduct is ongoing and continues to this date.
- 59. Unfair Business Practices: California Business & Professions Code § 17200 also prohibits any "unfair ... business act or practice."
 - 60. Defendants' acts, omissions, misrepresentations, practices and non-disclosures as

alleged herein also constitute "unfair" business acts and practices within the meaning of Business & Professions Code § 17200 et seq. in that its conduct is substantially injurious to consumers, offends public policy, and is immoral, unethical, oppressive, and unscrupulous as the gravity of the conduct outweighs any alleged benefits attributable to such conduct.

- 61. There were reasonably available alternatives to further Defendants' legitimate business interests, other than the conduct described herein.
- 62. <u>Fraudulent Business Practices</u>: California Business & Professions Code § 17200 also prohibits any "fraudulent business act or practice."
- 63. Defendant's claims, nondisclosures and misleading statements with respect to the Products, as more fully set forth above, were false, misleading and/or likely to deceive the consuming public within the meaning of Business & Professions Code § 17200.
- 64. Defendant's conduct caused substantial injury to Plaintiff and the other Class members. Plaintiff has suffered injury in fact and has lost money as a result of Defendants' unfair conduct.
- 65. Pursuant to section 17203 of the California Business & Professions Code, Plaintiff and the Class seek an order of this court enjoining Defendant from continuing to engage in unlawful, unfair, or deceptive business practices and any other act prohibited by law, including, but not limited to: (a) selling, marketing, or advertising the Products with false representations set forth above; (b) engaging in any of the illegal, fraudulent, misleading, unlawful, unfair and/or deceptive conduct described herein; and (c) engaging in any other conduct found by the Court to be illegal, fraudulent, misleading, unlawful, unfair and/or deceptive conduct.
- 66. In addition, Plaintiff requests that this Court enter such orders or judgments as may be necessary to restore to any person in interest any money which may have been acquired by means of such illegal practices as provided in Business & Professions Code § 17203, and for such other relief as set forth below.
- 67. Plaintiff engaged counsel to prosecute this action and is entitled to recover costs and reasonable attorney's fees according to proof at trial.

SECOND CAUSE OF ACTION

FALSE AND MISLEADING ADVERTISING (CAL. BUS. & PROF. CODE § 17500, et seq.)

- 68. Plaintiff incorporates by this reference the allegations contained in the preceding paragraphs as if fully set forth herein.
- 69. Plaintiff brings this claim individually and on behalf of the proposed Class against Defendants.
- 70. As alleged herein, Plaintiff has standing to pursue this claim as Plaintiff has suffered injury in fact and has lost money or property as a result of Defendant's actions. Specifically, Plaintiff purchased the Products for her own personal use. In so doing, Plaintiff relied upon the representations referenced above. Plaintiff would not have purchased the Product, nor would she have paid the price that she did, had she known that it contained a prostaglandin associated with a variety of side effects and health risks, and is sold in California in violation of California's Sherman Food Drug and Cosmetic Law as an unlawful drug.
- 71. Defendant violated Business & Professions Code § 17500 by publicly disseminating false and misleading advertisements regarding the Products.
- 72. Defendant's false and misleading advertisements were disseminated to increase the sales of the Products.
- 73. Defendant knew or should have known that their advertisements for the Products were false and misleading and that those advertisements would induce consumers to purchase the Products. Such advertisements have deceived and are likely to deceive the consuming public, in violation of Business & Professions Code § 17500.
- 74. Furthermore, Defendant publicly disseminated the false and misleading advertisements as part of a plan or scheme and with the intent to sell unproven and ineffective products.
- 75. Plaintiff and the members of the Class have suffered harm as a result of these violations of the FAL because they have incurred charges and/or paid monies for the Products that they otherwise would not have incurred or paid.
- 76. Defendant are aware, or by the exercise of reasonable care should have been aware, that the representations were untrue or misleading and that such conduct is in violation of the current injunction.

- 77. Plaintiff and the members of the Class have suffered injury in fact and have lost money as a result of Defendant's false representations and false advertising.
- 78. Pursuant to Business & Professions Code § 17535, Plaintiff and the members of the putative Class seek an order of this Court enjoining Defendant from continuing to engage, use, or employ their practice of advertising the Products.
- 79. Likewise, Plaintiff and the members of the putative Class seek an order requiring Defendant to disclose such misrepresentations, and additionally request an order awarding Plaintiff and other members of the putative class restitution of the money wrongfully acquired by Defendants by means of responsibility attached to Defendants' failure to disclose the existence and significance of said misrepresentations.

THIRD CAUSE OF ACTION

VIOLATION OF CALIFORNIA LEGAL REMEDIES ACT (CAL. CIV. CODE § 1750 et seq.)

- 79. Plaintiff incorporates by this reference the allegations contained in the preceding paragraphs as if fully set forth herein.
- 80. Plaintiff brings this claim individually and on behalf of the proposed Class against Defendants.
- 80. Plaintiff is a consumer as defined by California Civil Code § 1761(d). The Products are a goods within the meaning of the Act. Specifically, prior to the filing of this action, as alleged herein, Plaintiff purchased the Products for her own personal use. In so doing, Plaintiff relied upon the representations referenced above. Plaintiff would not have purchased the Product, nor would she have paid the price that she did, had she known that it contained a prostaglandin associated with a variety of side effects and health risks, and is sold in California in violation of California's Sherman Food Drug and Cosmetic Law as an unlawful drug.
- 81. Plaintiff has standing to pursue this claim as Plaintiff has suffered injury in fact and has lost money or property as a result of Defendants' actions as set forth herein.
- 82. Plaintiff has concurrently filed the declaration of venue required by Civil Code §1780(d) with this complaint.

1	based on Defendant's warranties; and (c) the Products do not have the characteristics, uses, or benefits		
2	as promised.		
3	101.	Plaintiffs mailed a pre-suit notice letter to Defendant consistent with Cal. Com. Code §	
4	2607(3)(a) and U.C.C. 2-607(3)(A). The letter was sent on behalf of Plaintiff and all other persons		
5	similarly situated.		
6	PRAYER FOR RELIEF		
7	WHEREFORE, Plaintiff, on behalf of herself and as representative of all other persons		
8	similarly situated, prays for judgment against Defendant, as follows:		
9	1.	An order certifying that the action may be maintained as a Class Action;	
10	2.	An order permanently enjoining Defendant from pursuing the policies, acts, and	
11	practices complained of herein;		
12	3.	An order requiring Defendant to pay restitution and all other forms of equitable	
13	monetary relief;		
14	4.	An order requiring Defendant to pay compensatory, statutory, and punitive damages in	
15	amounts to be determined by the Court and/or jury;		
16	5.	For pre-judgment interest from the date of filing this suit;	
17	6.	For reasonable attorneys' fees;	
18	7.	Costs of this suit; and,	
19	8.	Such other and further relief as the Court may deem necessary and appropriate.	
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21	DATED: October 10, 2023 FARNESE P.C.		
22		By: A	
23		Peter J. Farnese	
24		Attorneys for Plainitff	
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26		DEMAND FOR JURY TRIAL	
27	Plaintiff hereby demands a jury trial on all issues so triable.		
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2	DATED: October 10, 2023 FARNESE P.C.
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4	By: Poter I Formace
5	Peter J. Farnese
6	Attorneys for Plainitff
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FARNESE PC	-19- CLASS ACTION COMPLAINT