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UNITED STATES DISTRICT COURT
NORTHERN DISTRICT OF CALIFORNIA

ALEXIS SLATEN,
Plaintiff,
v.
CHRISTIAN DIOR, INC.,
Defendant.

Case No. 23-cv-00409-JSC

ORDER RE: MOTION TO DISMISS

Re: Dkt. No. 23

Plaintiff alleges Defendant misleadingly advertises the sun protection factor (“SPF”) benefits of its cosmetic products. (Dkt. No. 1.)¹ Before the Court is Defendant’s motion to dismiss. (Dkt. No. 23.) After carefully considering the briefing, and with the benefit of oral argument on May 11, 2023, the Court GRANTS the motion in part and DENIES it in part.

COMPLAINT ALLEGATIONS

Plaintiff, a California resident, has purchased Defendant’s Dior Forever Foundation from a Macy’s retail store in Daly City, California for several years. (Dkt. No. 1 ¶ 51.) She read and relied on the front of the Dior Forever Foundation box, which states:

TRANSFER-PROOF – 24H FOUNDATION
HIGH PERFECTION
CONCENTRATED FLORAL SKINCARE
WITH SUNSCREEN

BROAD SPECTRUM SPF 15

(*Id.* ¶ 21.) She believed the product would provide both cosmetic coverage and SPF protection for 24 hours. (*Id.* ¶¶ 22, 52.) The back of the product bottle (inside the box) also states:

¹ Record citations are to material in the Electronic Case File (“ECF”); pinpoint citations are to the ECF-generated page numbers at the top of the documents.

24H WEAR HIGH PERFECTION
SKIN-CARING FOUNDATION
WITH SUNSCREEN

LASTING COMFORT AND CARE

BROAD SPECTRUM SPF 35

(*Id.* ¶ 37.) It is not clear whether the product is offered in multiple SPF options, given that the complaint quotes both SPF 15 and 35. (*See id.* ¶¶ 22, 52.)

However, the product’s SPF protection lasts for two hours at most. (*Id.* ¶ 23.) The drug facts panel on the back of the product box directs consumers to “reapply at least every 2 hours.” (*Id.* ¶ 37.) Plaintiff would not have purchased, or would have paid less for, the product had she known Defendant’s labeling was deceptive and misleading. (*Id.* ¶¶ 54–55.) Plaintiff asserts if “the Products were reformulated and/or relabeled without the misleading 24 hour SPF claims, [she] would likely purchase the Products again in the future.” (*Id.* ¶ 56.) Plaintiff also challenges the labels on the Dior Forever Skin Glow Foundation product and any other on which Defendant “make[s] an SPF claim and a claim that the Products will last longer than two hours.” (*Id.* ¶ 20.)

Plaintiff asserts the product labels are misleading because a reasonable consumer will think they mean the products provide all benefits—including cosmetic coverage and SPF protection—for 24 hours without the need to reapply. (*Id.* ¶ 22.) On behalf of a putative nationwide class and California subclass who bought the products, Plaintiff brings claims for: (1) violation of California’s Consumer Legal Remedies Act (“CLRA”); (2) violation of California’s False Advertising Law (“FAL”); (3) fraud, deceit, and/or misrepresentation; (4) violation of all three prongs (unlawful, unfair, and fraudulent) of California’s Unfair Competition Law (“UCL”); and (5) unjust enrichment. (*Id.* ¶¶ 57, 66–111.) Plaintiff disclaims any causes of action under the Federal Food, Drug, and Cosmetic Act (“FDCA”) and regulations promulgated by the Food and Drug Administration (“FDA”). (*Id.* ¶ 65.) Accordingly, she relies on the FDCA and FDA regulations only to the extent they are also enacted under California state law or regulation, or provide a predicate for liability under state law. (*See id.*)

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1 **DISCUSSION**

2 **I. EXPRESS PREEMPTION**

3 Defendant argues the FDCA expressly preempts Plaintiff’s state claims. “FDCA
4 preemption, like all federal preemption, is an affirmative defense. Only when the plaintiff pleads
5 itself out of court—that is, admits all the ingredients of an impenetrable defense—may a
6 complaint that otherwise states a claim be dismissed under Rule 12(b)(6).” *Durnford v.*
7 *MusclePharm Corp.*, 907 F.3d 595, 603 n.8 (9th Cir. 2018) (cleaned up). In analyzing express
8 preemption,

9 our focus is on the plain meaning of [the statute]. That’s because the
10 plain wording of the clause necessarily contains the best evidence of
11 Congress’ preemptive intent. In discerning its meaning, we look to
12 [the statute’s] text, structure, and context. And we apply this textual
13 analysis without any presumptive thumb on the scale for or against
14 preemption.

15 *Cal. Rest. Ass’n v. City of Berkeley*, 65 F.4th 1045, 1050 (9th Cir. 2023) (cleaned up).

16 The FDCA’s preemption clause says,

17 [N]o State or political subdivision of a State may establish or continue
18 in effect any requirement—

19 (1) that relates to the regulation of a drug that is not subject to the
20 requirements of section 353(b)(1) [related to prescription drugs] or
21 353(f)(1)(A) [related to veterinary prescription drugs] of this title; and

22 (2) that is different from or in addition to, or that is otherwise not
23 identical with, a requirement under this chapter

24 21 U.S.C. § 379r(a). The statute regulates sunscreen as a drug. Among other requirements,
25 sunscreen products must include a drug facts label stating, “reapply at least every 2 hours.” 21
26 C.F.R. § 201.327(e)(4). Sunscreen products and other drugs “shall be deemed to be misbranded”
27 if their “labeling is false or misleading in any particular.” 21 U.S.C. § 352(a)(1). In particular,

28 There are claims that would be false and/or misleading on sunscreen
products. These claims include but are not limited to the following:
“Sunblock,” “sweatproof,” and “waterproof.” These or similar claims
will cause the product to be misbranded under [21 U.S.C. § 352].

21 C.F.R. § 201.327(g).

There are two alternative reasons the FDCA does not preempt Plaintiff’s claims. First,
Plaintiff’s claims do not implicate any FDCA “requirement” for a sunscreen product. 21 U.S.C. §

1 379r(a)(2). The FDCA does not require the front of Defendant’s sunscreen package, also known
2 as the “principal display panel,” 21 C.F.R. § 201.60, to say “24H” or anything else about how long
3 it lasts. There is no FDCA requirement “on point,” so no preemption. *Hollins v. Walmart Inc.*,
4 No. 21-56031, 2023 WL 3364616, at *8 (9th Cir. May 11, 2023). For example, in *Durnford*, 907
5 F.3d at 603, the label misleadingly suggested the product’s protein came from certain ingredients.
6 That claim was not preempted because FDA regulations set out a protocol to test protein content,
7 but not protein composition. *Id.* at 603–05; *see Hollins*, 2023 WL 3364616, at *8. And in *Astiana*
8 *v. Hain Celestial Grp., Inc.*, the products were labeled “All Natural” or “Pure Natural,” which was
9 not required by FDA regulations. 783 F.3d 753, 758 (9th Cir. 2015). A claim requiring the
10 defendant “to remove these allegedly misleading advertising statements” did not “run afoul of the
11 FDCA” because it would not “modify or enhance any aspect of [the] cosmetics labels that are
12 required by federal law.” *Id.*; *see Meza*, 2023 WL 3082346, at *5 (“Put another way, a claim will
13 not be expressly preempted under the FDCA if defendant’s labeling duties would remain the same
14 if plaintiffs prevailed.”). Similarly here, FDA regulations require sunscreen labels to include
15 several pieces of information, but none on point. *See Meza*, 2023 WL 3082346, at *6 (“Plaintiff
16 alleges . . . the ‘24/25 HR’ claims misleadingly suggest the Products will provide sun protection
17 for that amount of time when in fact they only provide two hours of protection. As in *Astiana*,
18 FDA regulations do not mandate that sunscreen manufacturers include claims pertaining to the
19 product’s durational capability. Instead, the regulations state that non-water-resistant products
20 include ‘reapply at least every 2 hours.’ As such, Plaintiff’s claims that the durational statements
21 violate state law would not lead to any additional or different requirements to the existing
22 regulations.” (cleaned up)).

23 Alternatively, to the extent Plaintiff’s claims impose a requirement on a sunscreen product,
24 it is “identical to” the FDCA’s requirement that sunscreen labels not be false or misleading. 21
25 U.S.C. § 379r(a)(2); *see id.* § 352(a)(1). As the Ninth Circuit explained in *Ebner v. Fresh, Inc.*:

26 [B]oth the federal FDCA and California’s Sherman Law prohibit the
27 false or misleading labeling of a cosmetic. Viewed in this light,
28 Plaintiff is not asking Fresh to modify or enhance any aspect of its
cosmetics labels that are required by federal law. Rather, the state-
law duty that Plaintiff seeks to enforce under the Sherman Law is

1 identical to Fresh’s federal duty under the FDCA: the duty to avoid
false or misleading labeling.
2 838 F.3d 958, 965 (9th Cir. 2016) (cleaned up). Because the FDCA prohibits false or misleading
3 labeling, it does not preempt California’s laws allowing consumers to sue manufacturers who
4 violate the federal prohibition on false or misleading labeling. “Simply put, the availability of
5 state law damages for violations of federal law does not amount to an additional or different
6 requirement.” *Astiana*, 783 F.3d at 757 (cleaned up). Defendant has not identified any part of the
7 FDCA suggesting “24H” is not false or misleading as a matter of law. *Cf. Roffman v. Perfect Bar,*
8 *LLC*, No. 22-CV-02479-JSC, 2022 WL 4021714, at *8 (N.D. Cal. Sept. 2, 2022) (“FDA
9 regulations allow the nitrogen method for front-label protein claims That the regulations
10 simultaneously prohibit ‘misleading’ labeling, shows that nitrogen-method protein claims are not
11 inherently misleading under the FDA regulations.” (cleaned up)).

12 Defendant insists the FDA considered regulating cosmetic duration claims in products with
13 sunscreen but decided not to, and therefore Plaintiff’s claims are preempted. (Dkt. No. 33 at 15–
14 16.) Not so. Defendant’s characterization of what the FDA considered is wrong. Several
15 commenters on FDA-proposed sunscreen regulations argued that different drug facts labeling
16 (what is on the back of the product) should apply to cosmetics with sunscreen. The FDA
17 concluded cosmetics with sunscreen should have the same drug facts as non-cosmetic sunscreen.
18 It then explained:

19 To help consumers understand that the sunscreen directions apply to
20 the use of the product as a drug, for sun protection, we are allowing
21 the optional statement “for sunscreen use:” to appear as the first line
22 under “Directions.” Consumers who are using these products
23 primarily for cosmetic use will be more likely to understand that they
24 might not receive the intended sun protection if they do not follow the
25 directions in the Drug Facts label.
26 76 Fed. Reg. 35,639 (June 17, 2011). This conclusion says nothing about duration claims on the
27 front label, and it also does not suggest that consumers will not be confused by a duration claim on
28 the front label.

29 Accordingly, Plaintiff’s claims are not preempted. Defendant’s motion to dismiss on this
basis is DENIED.

1 **II. FALSE OR MISLEADING UNDER CLRA, FAL, & UCL**

2 Plaintiff asserts Defendant violated the CLRA by misrepresenting the products’ “source,
3 sponsorship, approval, or certification”; representing they have “sponsorship, approval,
4 characteristics, ingredients, uses, benefits, or quantities that they do not have”; representing they
5 “are of a particular standard, quality, or grade” but are of another; and “[d]isparaging the goods . .
6 . of another by false or misleading representation of fact.” Cal. Civ. Code §§ 1770(a)(2), (5), (7),
7 (8). She also asserts Defendant’s labels are “untrue or misleading” under the FAL and “unfair”
8 and “fraudulent” under the UCL. Cal. Bus. & Prof. Code §§ 17500, 17200. All of these claims
9 are governed by the “reasonable consumer” test, under which Plaintiff must adequately allege
10 “members of the public are likely to be deceived.” *Ebner*, 838 F.3d at 965 (“This requires . . . a
11 probability that a significant portion of the general consuming public or of targeted consumers,
12 acting reasonably in the circumstances, could be misled.” (cleaned up)); *see also Becerra v. Dr*
13 *Pepper/Seven Up, Inc.*, 945 F.3d 1225, 1228 (9th Cir. 2019) (“This requires more than a mere
14 possibility that [the] label might conceivably be misunderstood by some few consumers viewing it
15 in an unreasonable manner.” (cleaned up)).

16 Plaintiff must also meet Rule 9(b)’s heightened pleading requirement for claims “grounded
17 in” fraud. *Kearns v. Ford Motor Co.*, 567 F.3d 1120, 1126 (9th Cir. 2009) (explaining that if a
18 claim relies on an alleged “unified course of fraudulent conduct,” Rule 9(b) applies even if fraud is
19 not a necessary element of the claim); *see, e.g., Loh v. Future Motion, Inc.*, No. 5:21-CV-06088-
20 EJD, 2022 WL 2668380, at *5 (N.D. Cal. July 11, 2022) (“Plaintiffs’ theory that Defendant
21 harmed its customers by fraudulently representing the safety and craftsmanship of the Onewheel
22 underlies each claim.”). Plaintiff’s claims are grounded in fraud because they assert Defendant’s
23 “24H” marketing statement is false. Thus, for each claim she must allege “the who, what, when,
24 where, and how of the misconduct charged.” *Vess v. Ciba-Geigy Corp. USA*, 317 F.3d 1097, 1106
25 (9th Cir. 2003) (cleaned up).

26 Plaintiff plausibly alleges the product labels are false or misleading and reasonable
27 consumers are likely to be misled. She alleges facts supporting a reasonable inference that
28 consumers increasingly care about sun protection, which supports an inference they may choose

1 Defendant’s products believing that its SPF benefits will last for “24H.” *See Bell Atl. Corp. v.*
2 *Twombly*, 550 U.S. 544, 570 (2007). Plaintiff also meets Rule 9(b)’s requirement to plead “*why*
3 the [statements are] false,” *Davidson v. Kimberly-Clark Corp.*, 889 F.3d 956, 964 n.2 (9th Cir.
4 2018): because the SPF protection lasts only two hours.

5 Defendant argues the “24H” promise applies only to the product’s cosmetic benefits, but,
6 drawing all inferences in Plaintiff’s favor, reasonable consumers could think otherwise. *See Bell*
7 *v. Publix Super Markets, Inc.*, 982 F.3d 468, 476–77 (7th Cir. 2020) (“With the time afforded by
8 litigation, we can see how ‘100% Grated Parmesan Cheese’ might be interpreted as claiming only
9 that whatever it contains is ‘100% grated’ Another reading, though, and certainly a plausible
10 reading, is that ‘100%’ applies to all three words: it’s all cheese; all the cheese is Parmesan, and
11 it’s all grated.”). On the product box, “24H” is not so far away from or stylistically different from
12 “WITH SUNSCREEN,” (Dkt. No. 1 ¶ 21), that only unreasonable consumers would think the
13 former applies to the latter. *See Meza v. Coty, Inc.*, No. 22-CV-05291-NC, 2023 WL 3082346, at
14 *10 (N.D. Cal. Apr. 24, 2023) (“The CoverGirl product . . . features a durational claim and SPF
15 claim grouped in close proximity. . . . As Plaintiff alleges, when read in vertical fashion, it is
16 reasonable to interpret that the product provides 24 hours of full coverage at SPF 18.”). “24H”
17 modifies “FOUNDATION,” which in turn is “WITH SUNSCREEN.” (Dkt. No. 1 ¶ 21.) That
18 makes this label different from the one at issue in *Klammer v. Mondelez Int’l, Inc.*, on which “high
19 protein” modified “lentils” and “lentil flour,” ingredients within the product (chips). No. 22-CV-
20 02046-JSW, 2023 WL 105095, at *3 (N.D. Cal. Jan. 4, 2023) (“[T]he phrase ‘high protein’ would
21 not lead the reasonable consumer to believe that the chips themselves are high in protein.”).

22 Defendant suggests it is impossible for sunscreen to last 24 hours. Even if that is true, the
23 Court cannot conclude reasonable consumers would know that as a matter of law. In *Moore v.*
24 *Trader Joe’s Co.*, for example, the Ninth Circuit held reasonable sophisticated consumers of
25 Manuka honey, an “effete” product, know bees are foraging insects and “it is impossible to
26 exercise complete control over where bees forage down to each specific flower or plant.” 4 F.4th
27 874, 883, 884 n.11 (9th Cir. 2021). Therefore, “as a matter of law, other available information
28 about Trader Joe’s Manuka Honey would quickly dissuade a reasonable consumer from the belief

1 that Trader Joe’s Manuka Honey was derived from 100% Manuka flower nectar.” *Id.* at 883. The
2 scientific consensus about how long a drug lasts is not as obvious to the reasonable consumer. It
3 is nowhere near as obvious as that Froot Loops and Cap’n Crunch cereal, though colorful, are not
4 made of real fruit. *See id.*; *see also, e.g., Gitson v. Trader Joe’s Co.*, No. 13-CV-01333-VC, 2015
5 WL 9121232, at *1 (N.D. Cal. Dec. 1, 2015) (“The reasonable consumer (indeed, even the least
6 sophisticated consumer) does not think soymilk comes from a cow.”); *Red v. Kraft Foods, Inc.*,
7 No. CV 10–1028–GW(AGRx), 2012 WL 5504011, at *3 (C.D. Cal. Oct. 25, 2012) (“a reasonable
8 consumer will be familiar with the fact of life that a cracker is not composed of primarily fresh
9 vegetables”). Indeed, that the FDA requires sunscreen labels to tell consumers to reapply suggests
10 many do not know they should do so. *See POM Wonderful LLC v. Coca-Cola Co.*, 573 U.S. 102,
11 108 (2014) (“The FDCA statutory regime is designed primarily to protect the health and safety of
12 the public at large.”); *Wyeth v. Levine*, 555 U.S. 555, 574 (2009) (“Congress enacted the FDCA to
13 bolster consumer protection against harmful products.”).

14 Other than the drug facts panel’s instruction to reapply after two hours, nothing else about
15 the product, such as price, creates a “contextual inference[.]” that would lead the reasonable
16 consumer to know the SPF benefits last less than 24 hours. *Moore*, 4 F.4th at 882 (“information
17 available to a consumer is not limited to the physical label and may involve contextual inferences
18 regarding the product itself and its packaging”). And the drug facts panel does not cure, as a
19 matter of law, what is plausibly false or misleading about “24H.” *See Williams v. Gerber Prods.*
20 *Co.*, 552 F.3d 934, 939 (9th Cir. 2008) (“We disagree with the district court that reasonable
21 consumers should be expected to look beyond misleading representations on the front of the box
22 to discover the truth from the ingredient list in small print on the side of the box.”); *Ebner*, 838
23 F.3d at 966 (“*Williams* stands for the proposition that *if* the defendant commits an act of deception,
24 the presence of fine print revealing the truth is insufficient to dispel that deception.”).

25 * * *

26 Plaintiff adequately alleges Defendant’s statements are false or misleading under the
27 CLRA and FAL and unfair and fraudulent under the UCL. Accordingly, she also adequately
28 alleges the statements are “unlawful” under the UCL’s third prong. *Cel-Tech Commc’ns, Inc. v.*

1 *L.A. Cellular Tel. Co.*, 973 P.2d 527, 539–40 (1999) (“By proscribing ‘any unlawful’ business
2 practice, section 17200 borrows violations of other laws and treats them as unlawful practices that
3 the [UCL] makes independently actionable.” (cleaned up)). Defendant’s motion to dismiss on this
4 basis is DENIED.

5 **III. OTHER PRODUCTS**

6 “[A] plaintiff may have standing to assert claims for unnamed class members based on
7 products he or she did not purchase so long as the products and alleged misrepresentations are
8 substantially similar.” *Miller v. Ghirardelli Chocolate Co.*, 912 F. Supp. 2d 861, 869 (N.D.
9 Cal. 2012). “Although the Ninth Circuit has not directly addressed ‘substantial similarity’ for
10 purposes of consumer fraud-based class actions, district courts, driven by *Armstrong*, have taken a
11 broad approach.” *Cimoli v. Alacer Corp.*, 546 F. Supp. 3d 897, 907 (N.D. Cal. 2021); *see*
12 *Armstrong v. Davis*, 275 F.3d 849, 867 (9th Cir. 2001).

13 Plaintiff alleges the Dior Forever Skin Glow Foundation, which she did not buy,
14 “predominately, uniformly, and consistently include[s], on the principal display panel of the
15 product boxes and bottles, an SPF claim alongside a claim that the product[] last[s] longer than
16 two hours.” (Dkt. No. 1 ¶ 20.) But she does not quote or provide a photo of the label text. *See*
17 *Anderson v. The Hain Celestial Grp., Inc.*, 87 F. Supp. 3d 1226, 1233 (N.D. Cal. 2015). As such,
18 the Court cannot evaluate “the similarity of the claims and injuries flowing from the
19 misrepresentations on each product.” *Cimoli*, 546 F. Supp. 3d at 908; *cf. Miller*, 912 F. Supp. 2d
20 at 870–72 (“[T]he products are different: baking chips, three drink powders, and wafers. . . . They
21 look different, they are labeled differently (white mocha for a drink, Frappe Classico Classic
22 White for another drink) [Plaintiff’s] best argument is that the labels imply white chocolate
23 content, but the alleged misrepresentations vary widely.”). Similarly, the Court cannot evaluate
24 whether other unidentified products are similar to the product Plaintiff bought. *See Meza*, 2023
25 WL 3082346, at *4.

26 Accordingly, Plaintiff has not established her standing to assert class claims based on
27 products she did not buy. Defendant’s motion to dismiss on this basis is GRANTED.

28

1 **IV. INJUNCTIVE RELIEF**

2 As a matter of Article III standing to seek injunctive relief, a plaintiff must establish “an
3 actual and imminent, not conjectural or hypothetical threat of future harm.” *Davidson*, 889 F.3d at
4 969 (cleaned up). In the false advertising context,

5 a previously deceived consumer may have standing to seek an
6 injunction . . . even though the consumer now knows or suspects that
7 the advertising was false at the time of the original purchase
8 Knowledge that the advertisement or label was false in the past does
9 not equate to knowledge that it will remain false in the future. In some
10 cases, the threat of future harm may be the consumer’s plausible
11 allegations that she will be unable to rely on the product’s advertising
or labeling in the future, and so will not purchase the product although
she would like to. In other cases, the threat of future harm may be the
consumer’s plausible allegations that she might purchase the product
in the future, despite the fact it was once marred by false advertising
or labeling, as she may reasonably, but incorrectly, assume the
product was improved.

12 *Id.* at 969–70 (cleaned up). Plaintiff alleges she would like to buy Defendant’s products and
13 would likely do so in the future if they “were reformulated and/or relabeled without the misleading
14 24 hour SPF claims.” (Dkt. No. 1 ¶ 56.) Drawing all inferences in Plaintiff’s favor, suppose
15 Defendant changed its product to provide SPF protection for a longer or shorter length of time
16 than two hours; Plaintiff might not be able to tell whether the labels were accurate. *See Davidson*,
17 889 F.3d at 971. Or suppose Defendant changed its labels to state the SPF protection lasts for two
18 hours, not 24; Defendant might have to lower the price, and Plaintiff might be willing to pay that
19 lower price.

20 Accordingly, drawing all inferences in her favor, Plaintiff has established standing to seek
21 injunctive relief. Defendant’s motion to dismiss on this basis is DENIED.

22 **CONCLUSION**

23 Defendant’s motion is GRANTED as to Plaintiff’s claims based on products she did not
24 buy; they are dismissed with leave to amend. *See Yagman v. Garcetti*, 852 F.3d 859, 863 (9th Cir.
25 2017). Defendant’s motion is otherwise DENIED as to Plaintiff’s CLRA, FAL, UCL, and fraud,
26 deceit, and/or misrepresentation claims.

27 Plaintiff may file an amended complaint on or before **June 2, 2023**.

28 This Order disposes of Docket No. 23.

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IT IS SO ORDERED.

Dated: May 12, 2023



JACQUELINE SCOTT CORLEY
United States District Judge