

United States District Court
Northern District of California

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UNITED STATES DISTRICT COURT
NORTHERN DISTRICT OF CALIFORNIA
SAN JOSE DIVISION

MELISSA SANCHEZ, et al.,
Plaintiffs,
v.
NURTURE, INC.,
Defendant.

Case No. [5:21-cv-08566-EJD](#)

**ORDER GRANTING IN PART AND
DENYING IN PART MOTION TO
DISMISS**

Re: ECF No. 35

Plaintiffs Melissa Sanchez and Beverly Cassel bring this food labeling action on behalf of themselves and those similarly situated to challenge the labeling practices of Defendant Nurture, Inc.’s “Happy Baby” and “Happy Tot” line of food products. The Court had previously dismissed Plaintiff Sanchez’s complaint with leave to amend (ECF No. 24), which has since been amended to add an additional plaintiff and new theories of fraud and deceptive labeling. First Am. Class Action Compl. (“FAC”), ECF No. 29. Defendant moved again to dismiss the claims in the FAC (ECF No. 35, “Mot.”), and the Court heard oral arguments on April 27, 2023.

Based on the following, the Court GRANTS IN PART and DENIES IN PART Defendant’s motion to dismiss.

I. BACKGROUND

A. Defendant’s Products

Defendant Nurture, Inc. is a Delaware corporation that manufactures, distributes, markets, advertises, and sells a line of baby and toddler food products under the brand names, “Happy Baby” and “Happy Tot.” FAC ¶ 13. At issue here are forty-three (43) products (the “Products”) that allegedly contain “nutrient content” claims and are intended for children under the age of two.

1 FAC ¶¶ 13–14; *see also id.*, Ex. A (“Product Chart”).

2 The challenged nutrient content claims are prominently displayed on Defendant’s Products
 3 and contain language such as “2g of Protein, 4g of Fiber and 350 mg Omega-3 from Chia ALA.”
 4 *Id.* ¶ 37. The nutrient amounts reflected in the nutrition content claims match the information
 5 contained on the nutritional facts panel on the back of the Products. *Id.* Some Products also
 6 advertise that additional nutrients were added, such as protein and choline, *e.g.*, “+ 1 1/3 tsp Pea
 7 Protein.” *Id.* ¶ 39. The FAC alleges that the nutrient content claims on Defendant’s Product
 8 “deceive and mislead reasonable consumer into believing that the Products provide *physical health*
 9 *benefits* for their children when in fact, the Products are *harmful for children under two both*
 10 *nutritionally and developmentally.*” FAC ¶¶ 21 (emphasis added), 44. The FAC also alleges these
 11 claims are deceptive because they “mislead[] consumers into believing that foods for children
 12 under two should be purchased based on the quantities of the listed nutrients, when other
 13 considerations are just as, or more, important.” *Id.* ¶ 53.

14 The FAC cites three harmful effects of the Products on young children. First, the
 15 “Products have high amounts of both added sugars and free sugars.” *Id.* ¶ 56. Second, because
 16 many of Defendant’s Products are pureed, the raw ingredients are “stripped of insoluble fiber and
 17 the liver is no longer protected from the sugar in the food.” *Id.* ¶ 57. And third, the FAC alleges
 18 that the long-term use of puree pouches may be detrimental to children because they prevent
 19 children from learning to chew and swallow soft foods, which in turn may lead to “delays in motor
 20 development” and “bad long-term snacking habits and routine overeating.” *Id.* ¶¶ 59–64.

21 **B. Plaintiffs**

22 Plaintiff Melissa Sanchez is a California consumer who purchased nine (9) different Happy
 23 Baby and Happy Tots Products from Buy Buy Baby and Target stores in San Jose, California. *See*
 24 FAC ¶ 75 (listing products). Ms. Sanchez purchased these Products after reading the nutrient
 25 content claims on their labels, believing them to be “superior in providing nutrition for her child.”
 26 *Id.* ¶ 76. The FAC alleges that, without the nutrient content on the labels, Ms. Sanchez either
 27 would not have purchased Defendant’s Products or would not have been willing to pay a premium

1 for them. *Id.* ¶¶ 78–79. Ms. Sanchez regularly shops at stores where the Products are sold and, if
 2 Defendant’s Products did not contain the misleading labels, she would likely purchase the
 3 Products again in the future. *Id.* ¶ 80.

4 Plaintiff Beverly Cassel purchased four (4) different Happy Baby and Happy Tots Products
 5 from Safeway and Whole Foods stores in and around Santa Rosa, California. FAC ¶ 81. Like Ms.
 6 Sanchez, Ms. Cassel purchased these products after reading the nutrient content claims, believing
 7 them to be “healthful for her child under two.” *Id.* ¶ 82. Without the nutrient content claims, Ms.
 8 Cassel would not have purchased the Products or paid a premium for them, but if the Products’
 9 labels are no longer misleading, she would likely purchase them again. *Id.* ¶¶ 84–86.

10 The FAC further alleges that Plaintiffs have been economically injured because the
 11 Products are worth less than the price they had paid. FAC ¶¶ 87–88.

12 **C. Procedural History**

13 On November 3, 2021, Plaintiff Sanchez filed the original complaint against Defendant,
 14 alleging that Defendant’s products have been improperly labeled and misbranded in violation of
 15 several California and federal laws, including the CLRA, FAL, UCL, common law fraud, and
 16 unjust enrichment. ECF No. 1.

17 Defendant moved to dismiss the Complaint on January 14, 2022 (ECF No. 13), which the
 18 Court granted and denied in part on September 7, 2022. ECF No. 24 (“Sept. 7 Order”). In its
 19 September 7 Order, the Court found that Plaintiff had sufficiently stated a claim under the
 20 unlawful prong of the UCL and unjust enrichment claims, because the Products violated Food and
 21 Drug Administration (“FDA”) regulations by containing “nutrient content” claims on products
 22 intended for children under two. Sept. 7 Order 9–11. However, the Court dismissed the
 23 Complaint’s claims alleging deception and fraud, finding that a reasonable consumer would not be
 24 misled by truthful information regarding the Product’s nutrient content. *Id.* at 11–13.

25 On October 7, 2022, Plaintiff filed the First Amended Complaint, adding Ms. Cassel as an
 26 additional plaintiff and further allegations supporting the deception of the Products’ labels. *See*
 27 ECF No. 35-5 (redlined version of FAC). Defendant filed the instant motion to dismiss the FAC

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1 on November 8, 2022, and briefing was completed by January 20, 2023. The Court heard oral
2 arguments on April 27, 2023.

3 **II. LEGAL STANDARD**

4 **A. Rule 12(b)(6)**

5 Federal Rule of Civil Procedure 8(a) requires a plaintiff to plead each claim with enough
6 specificity to “give the defendant fair notice of what the . . . claim is and the grounds upon which
7 it rests.” *Bell Atl. Corp. v. Twombly*, 550 U.S. 544, 555 (2007) (internal quotations omitted). A
8 complaint which falls short of the Rule 8(a) standard may therefore be dismissed if it fails to state
9 a claim upon which relief can be granted. Fed. R. Civ. P. 12(b)(6). When deciding whether to
10 grant a motion to dismiss, the Court must accept as true all “well pleaded factual allegations” and
11 determine whether the allegations “plausibly give rise to an entitlement to relief.” *Ashcroft v.*
12 *Iqbal*, 556 U.S. 662, 679 (2009). The Court must also construe the alleged facts in the light most
13 favorable to the plaintiff. *Love v. United States*, 915 F.2d 1242, 1245 (9th Cir. 1989). While a
14 complaint need not contain detailed factual allegations, it “must contain sufficient factual matter,
15 accepted as true, to ‘state a claim to relief that is plausible on its face.’” *Iqbal*, 556 U.S. at 678
16 (quoting *Bell Atl. Corp.*, 550 U.S. at 570).

17 **B. Rule 9(b)**

18 Consumer protection claims that sound in fraud are subject to the heightened pleading
19 requirements of Federal Rule of Civil Procedure 9(b). *See Vess v. Ciba-Geigy Corp. USA*, 317
20 F.3d 1097, 1102 (9th Cir. 2003); *San Miguel v. HP Inc.*, 317 F. Supp. 3d 1075, 1084 (N.D. Cal.
21 2018). Rule 9(b) requires that “a party must state with particularity the circumstances constituting
22 fraud.” Fed. R. Civ. P. 9(b). The circumstances constituting the fraud must be “specific enough to
23 give defendants notice of the particular misconduct which is alleged to constitute the fraud
24 charged so that they can defend against the charge and not just deny that they have done anything
25 wrong.” *Semegen v. Weidner*, 780 F.2d 727, 731 (9th Cir. 1985). Therefore, a party alleging
26 fraud must set forth “the who, what, when, where, and how” of the misconduct. *Vess*, 317 F.3d at
27 1106 (quoting *Cooper v. Pickett*, 137 F.3d 616, 627 (9th Cir. 1997)).

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1 **III. DISCUSSION**

2 The FAC asserts five claims for relief: (1) violations of the CLRA; (2) violations of the
3 FAL; (3) common law fraud, deceit, and/or misrepresentation; (4) violations of the UCL; and (5)
4 unjust enrichment. Plaintiffs seek various damages, restitution, and injunctive relief. Defendant
5 moves to dismiss all claims under Rule 12(b)(6).

6 **A. UCL “Unlawful” Prong**

7 Defendant argues that Plaintiffs’ UCL “unlawful” claim should be dismissed because it is
8 preempted and includes claims that the Court had found lawful in its prior order. Mot. 16–19.
9 Defendant also challenges the “unlawful” UCL claim to the extent it is premised on Plaintiffs’
10 newly alleged violations of the FDA’s fortification policy. The Court addresses each in order.

11 **1. Federal Preemption**

12 Although the Court previously held that the original Complaint had sufficiently stated a
13 claim under the UCL’s “unlawful” prong based on the Products’ violation of certain FDA
14 regulations, Defendant again challenges the sufficiency of this claim. This time, Defendant argues
15 that—because the “unlawful” UCL claim is “derivative” of FDA regulations and only the federal
16 government has exclusive authority to enforce Food, Drug, and Cosmetic Act (“FDCA”)
17 regulations—Plaintiffs’ state UCL claim is preempted. Mot. 16 (citing *Buckman Co. v. Plaintiffs’*
18 *Legal Committee*, 531 U.S. 341 (2001) (holding that “state-law fraud-on-the-FDA claims conflict
19 with, and are therefore impliedly pre-empted by, federal law”). Plaintiffs respond by noting that
20 California has expressly incorporated the FDA’s regulations as state law under the Sherman Food,
21 Drug, and Cosmetic Law (“Sherman Law”). Opp. 6–9; Cal. Health & Safety Code § 110100(a)
22 (“All food labeling regulations and any amendments to those regulations adopted pursuant to the
23 [FDCA], in effect on January 1, 1993, or adopted on or after that date shall be the food labeling
24 regulations of this state.”).

25 As in the first round of motions to dismiss, Plaintiffs have the stronger argument.
26 Defendant attempts to portray the California Sherman Law as a mirror image reflection of the
27 federal regulations, such that if “the FDA were to remove the regulations on which Plaintiffs rely,

1 their claims would fail as a matter of law.” Opp. 16–17. As a result, Defendant argues, Plaintiffs’
 2 “unlawful” UCL claim “exist solely by virtue of the FDCA [] requirements,” and therefore is
 3 impliedly preempted per *Buckman*. 531 U.S. at 353; *but see McClellan v. I-Flow Corp.*, 776 F.3d
 4 1035, 1041 (9th Cir. 2015) (finding no *Buckman* preemption and rejecting suggestion that “any
 5 use of federal law to establish a standard of care is an attempt to enforce the underlying federal
 6 provisions”). Defendant’s characterization, however, is inconsistent with the Sherman Law’s
 7 statutory language, which purports to create a separate set of state regulations alongside federal
 8 regulations. The Sherman Law states that the FDA’s food labeling regulations “shall be the food
 9 labeling regulations” of California, but it also permits the California department to “adopt
 10 additional food labeling regulations.” Cal. Health & Safety Code § 110100(b). Additionally,
 11 although new FDA regulations (including rescissions) are generally adopted as California
 12 regulations as a matter of course, that adoption only occurs thirty days after the federal
 13 regulation’s implementation and only if there are no substantial objections to the regulation during
 14 that interim period; a “timely filing of substantial objections . . . stays the adoption of the
 15 regulation” in California. *Id.* § 110115 (emphasis added). These features of the Sherman Law
 16 indicate that it is more than a federal doppelganger and operates as an independent source of state
 17 law from the FDCA, thereby avoiding federal preemption under *Buckman*.¹

18 In short, Plaintiffs’ “unlawful” UCL claim is properly predicated upon violations of
 19 California’s Sherman Law and, therefore, do not implicate *Buckman*’s preemption holding.

20 2. Claims Addressed in Prior Order

21 Defendant argues that the Product Chart appended to the FAC contains claims that the
 22 Court previously found were not nutrient content claims. Mot. 18–19; Sept. 7 Order 10–11. After
 23

24 ¹ The Court notes that this conclusion accords with several others from this district that have
 25 “routinely reject[ed] the argument that the [Supreme] Court’s reasoning in *Buckman* justifies
 26 preemption of food labeling claims under the Sherman Law.” *Brown v. Van’s Int’l Foods, Inc.*,
 27 2022 WL 1471454, at *8 (N.D. Cal. May 10, 2022) (collecting cases). However, the Court
 28 notably parts company with the preemption analysis in *Davidson v. Sprout Foods Inc.*, No. 22-
 CV-01050-RS, 2022 WL 13801090, at *4 (N.D. Cal. Oct. 21, 2022), an opinion that this Court
 otherwise cites with approval later in this Order.

1 reviewing all the claims in the Product Chart, the Court finds that the claims for the “Happy Baby
2 Cereal” products (“With iron to support brain development”) are not nutrient content claims and
3 conflict with the Court’s prior holdings. Product Chart 12; *see* Sept. 7 Order 11 (“Claims about iron
4 are not unlawful.”). All other products in the Product Chart include some claim that would qualify as
5 a nutrient content claim. Accordingly, the Court GRANTS Defendant’s motion to dismiss the two
6 products under the heading “Happy Baby Cereal” without leave to amend.

7 3. Fortification Claims

8 In addition to the FDA’s regulations against content claims for children under two,
9 Plaintiffs also allege that certain Defendant’s labels violate the FDA’s policy against “random
10 fortification of foods.” FAC ¶ 39 (citing 21 C.F.R. § 104.20). Here, the labels at issue advertise
11 that certain Products have “+1 1/3 tsp Pea Protein” and “+ super chia.” *Id.*; *see also* Opp. 18 n.5.
12 Per the FAC’s Product Chart, the only claims that contain a “+” symbol are the “+1 1/3 tsp Pea
13 Protein” claims on “Fiber & Protein” pouches.² *See* Product Chart at 6–7.

14 However, neither Plaintiff is alleged to have purchased the “Fiber & Protein” *pouches*; the
15 FAC only alleges that Ms. Cassel purchased the “Fiber & Protein” *bars*, which do not contain the
16 challenged fortification claim. FAC ¶ 81; *see also* Product Chart at 1–2 (showing claims for the
17 “Fiber & Protein” bars”). In the Ninth Circuit, class plaintiffs may bring claims for products they
18 did not purchase only if the products and misrepresentations are “substantially similar” to those
19 they did purchase. *See, e.g., Corbett v. Pharmicare U.S., Inc.*, 544 F. Supp. 3d 996, 1011 (S.D.
20 Cal. 2021). In its prior order, the Court held that Plaintiffs’ purchases of pouches and puff
21 products do not also afford them standing to pursue claims made on bar products. *See* Sept. 7
22 Order 9. Likewise, the Court now finds that Ms. Cassel’s purchase of “Fiber & Protein” bars that
23 do not display any fortification claims are not “substantially similar” to “Fiber & Protein” pouches
24

25 _____
26 ² Plaintiffs’ attempt to cast “+ super chia” as an unlawful nutrient fortification claim is
27 unpersuasive. The “+ super chia” claim more closely resembles the addition of an *ingredient*, as
28 opposed to the “random fortification” of a *nutrient*, *e.g.*, a vitamin, mineral, or protein. *See* 21
C.F.R. § 104.20(d)(3) (listing regulated fortification nutrients, which does not include any food
ingredients); § 104.20(h) (regulating labels about the “addition of a vitamin, mineral, or protein”).
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1 that *do* contain the challenged fortification claim. Therefore, Plaintiffs do not have standing to
 2 pursue their “unlawful” UCL claim based on the FDA’s fortification policy.

3 * * *

4 Accordingly, the Court finds that Plaintiffs have stated a UCL “unlawful” claim for
 5 Products that violate the FDA’s regulations on nutrient content claims, and Defendant’s motion is
 6 once again DENIED in those respects. However, Plaintiffs have not established standing to sue
 7 for the “+1 1/3 tsp Pea Protein” alleged fortification claim that neither of them saw or purchased.
 8 The motion is, therefore, GRANTED to the extent Plaintiffs seek to bring a UCL “unlawful” claim
 9 that those Products violate the FDA’s policy against fortification. Because Plaintiffs have not yet
 10 had an opportunity to amend their complaint as to fortification claims, the Court will grant
 11 Plaintiffs LEAVE TO AMEND on this claim only, with the understanding that any new theories
 12 of liability will not be countenanced in an amended complaint.

13 **B. Fraud-Based Claims (Claims 1–4)**

14 Both Plaintiffs and Defendant address the CLRA, FAL, UCL, and common law fraud
 15 claims collectively, because they all turn on the same theories and allegations of deception from
 16 Defendant’s labeling. Mot. 4; Opp. 10, 20. The Court will also proceed accordingly.

17 Claims of consumer deception under California’s UCL, FAL, and CLRA statutes are
 18 governed by the “reasonable consumer” standard. *McGinity v. Procter & Gamble Co.*, 69 F.4th
 19 1093, 1097 (9th Cir. 2023). Under this standard, plaintiffs must show that “members of the public
 20 are likely to be deceived” by advertising that is either false, actually misleading, or “has a
 21 capacity, likelihood, or tendency to deceive or confuse the public.” *Id.* (citing *Kasky v. Nike, Inc.*,
 22 27 Cal. 4th 939, 951 (2002)). The “reasonable consumer” standard also contemplates a
 23 “probability that a significant portion of the general consuming public or of targeted consumers,
 24 acting reasonably in the circumstances, could be misled.” *Ebner v. Fresh, Inc.*, 838 F.3d 958, 965
 25 (9th Cir. 2016) (quoting *Lavie v. Procter & Gamble Co.*, 105 Cal. App. 4th 496, 508 (2003)).

26 **1. Reasonable Consumer**

27 Plaintiffs’ core theory of deception appears to be that the Products are misleadingly

1 advertised as “providing physical health benefits” when they are actually harmful—both
 2 nutritionally and developmentally—for children under two. FAC ¶¶ 44, 69, 100. With respect to
 3 the “physical health benefits,” the FAC alleges that the Products claim that protein is a “key
 4 building block for little growing bodies,” fiber “helps support the digestive system,” and DHA and
 5 choline provides “important nutrients to help support a healthy brain.” *Id.* ¶¶ 16–18. As for the
 6 nutritional and developmental harms, Plaintiffs provide allegations of the Products’ varying levels
 7 of added sugars (*id.* ¶¶ 54–57) and the fact that long-term use of pouches could stunt children’s
 8 motor development and create poor snacking habits (FAC ¶¶ 59–65).

9 The Court first notes that, although there has been a cornucopia of food labeling decisions
 10 from this district, Chief Judge Seeborg recently issued a decision involving nearly identical facts
 11 and allegations as the instant case. *See Davidson v. Sprout Foods Inc.*, No. 22-CV-01050-RS,
 12 2022 WL 13801090 (N.D. Cal. Oct. 21, 2022), *appeal filed*, No. 22-16656. The plaintiffs in
 13 *Sprout Foods*—represented by the same plaintiffs’ counsel in this case—asserted the same
 14 theories of fraud that Plaintiffs do here, namely that the defendant’s labels “communicate[d] a
 15 message that the Products provide physical health benefits for children; and second, that the
 16 Products are ‘harmful both nutritionally and developmentally.’” *Id.* at *3. The *Sprout Foods*
 17 plaintiffs even alleged the same harm, that the baby food products contained “high amounts of free
 18 sugars” and that “pouch-based foods may be unhealthy for developing children.” *Id.* In
 19 concluding that the plaintiffs had failed to plausibly allege harmful effects, Chief Judge Seeborg
 20 (1) rejected the allegations of “free sugars” harm because the plaintiffs did not allege “at what
 21 point ‘high’ sugar content crosses into harmful levels”; (2) rejected the allegations of “pouch-
 22 based” harm because they relied on “speculative research conclusions and hypothetical scenarios”;
 23 and (3) further reasoned that plaintiffs failed to allege that the harms would “outweigh any
 24 potential benefits of the Products . . . such that the Products no longer provide any physical health
 25 benefits.” *Id.* The Court finds the facts in *Sprout Foods* to be strikingly apt and Chief Judge
 26 Seeborg’s analysis in the “CLRA, FAL, Common Law Fraud, and UCL Fraudulent Practice
 27 Claims” section to be persuasive and well-reasoned. 2022 WL 2668481, at *4–6.

1 For all the reasons outlined in *Sprout Foods*, the Court finds that the near identical facts
2 here also fail to allege that a reasonable consumer could be misled by Defendant’s claims.

3 **a. “Physical Health Benefits”**

4 The Court, however, writes further to emphasize a particular deficiency that Judge Seeborg
5 highlighted, namely the FAC’s failure to allege any kind of nexus between the Products’ “physical
6 health benefits” representations and the harms stemming from free sugars and pouch usage.

7 In *Andrade-Heymfield v. NextFoods*, Judge Moskowitz dismissed a complaint where there
8 was a “mismatch” between plaintiff’s alleged harm (excessive amounts of free sugar) and the
9 challenged healthful claims (probiotics and promotion of digestive health). 2022 WL 1772262, at
10 *4–5 (S.D. Cal. Apr. 27, 2022). In so holding, Judge Moskowitz found that “a reasonable
11 consumer would not be misled given that the sugar content is clear, explicit, and otherwise not
12 contrary to the promotion of digestive health promoted on the packaging label.” *Id.* at *5.

13 Here, as well, the Court cannot plausibly infer that a reasonable consumer who is exposed
14 to claims of “3g Protein” and “PROTEIN is a key building block for little growing bodies” (FAC
15 ¶ 16) will be deceived as to the amounts of sugar in the Products or harmful effects of pouch
16 usage. Put differently, even if the Products’ sugar content and pouch-related concerns could result
17 in deleterious effects, the FAC does not allege that those effects would vitiate, negate, or even
18 relate to the specific “physical health benefits” conferred by the Products’ protein, fiber, or choline
19 content, as claimed. Just because a defendant truthfully advertises one of its product’s health
20 benefits does not mean that it has committed fraud or deceived consumers as to all other
21 conceivable unhealthy consequences that were not disclosed.³ Such a broad proposition would be
22 antithetical to both the heightened pleading standards of Rule 9(b) and the “reasonable consumer”
23 standard in UCL, FAL, and CLRA cases.

24 **b. “Happy and Healthy Start”**

25 Nor can the Court infer that the more general phrase “Here’s to a happy and healthy start!”
26

27 _____
28 ³ Plaintiffs do not purport to defend these claims as an omissions-based fraud claim.

1 would deceive a reasonable consumer into believing the product did not contain free sugars. The
 2 Court first clarifies that Plaintiffs disclaim any notion that they are alleging Defendant’s Products
 3 to be making a “health claim,” which is defined as any claim that “characterizes the relationship of
 4 any substance to a disease or health-related condition” and are preempted by federal regulations at
 5 21 C.F.R. § 101.14. Opp. 15; *see Clark v. Perfect Bar, LLC*, 816 F. App’x 141, 143 (9th Cir.
 6 2020) (finding that “applicable regulations are silent on whether sugar levels preclude a product
 7 from making health claims. . . . Allowing a claim of misbranding under California law based on
 8 misleading sugar level content would ‘indirectly establish’ a sugar labeling requirement ‘that is
 9 not identical to the federal requirements.’”). Accordingly, the Court will construe Plaintiffs’
 10 theories of fraud as challenging the Products only to the extent that their nutrient content claims —
 11 not any broader “health claims”—are misleading.

12 On this point, many courts in this district have rejected theories of fraud where plaintiffs
 13 alleged the presence of added sugars rendered a general health-related claim fraudulent. *See, e.g.,*
 14 *Yoshida v. Campbell Soup Co.*, 2022 WL 1819528, at *1 (N.D. Cal. May 27, 2022) (“The
 15 complaint alleges that the sugars occurring naturally in the fruits and vegetables . . . make label
 16 phrases such as ‘boost your morning nutrition’ and ‘healthy greens’ deceptive to consumers. . . .
 17 **No reasonable consumer would be misled by the challenged phrases** because the actual sugar
 18 content is plainly stated on the labels.”) (emphasis added); *Truxel v. Gen. Mills Sales, Inc.*, 2019
 19 WL 3940956, at *4 (N.D. Cal. Aug. 13, 2019) (“Defendant is under **no obligation to warn its**
 20 **consumers that certain levels of sugar may be associated with poor health results.** In fact,
 21 federal express preemption bars that demand as federal law for the disclosure of sugar content in
 22 food imposes no such requirement.”) (emphasis added); *Clark v. Perfect Bar, LLC*, 2018 WL
 23 7048788, at *1 (N.D. Cal. Dec. 21, 2018), *aff’d*, 816 F. App’x 141 (9th Cir. 2020) (“Plaintiffs’
 24 grievance is that the packaging led them to believe that the bars would be ‘healthy’ when, in
 25 supposed point of fact, the added sugar rendered them unhealthy or, in the alternative, less healthy
 26 from what they otherwise had believed. This is untenable. . . **Reasonable purchasers could**
 27 **decide for themselves how healthy or not the sugar content would be.**”) (emphasis added).

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1 That said, the Court recognizes that several other courts in this district have reached the opposite
 2 conclusion, finding general claims of healthfulness to be misleading where the food product
 3 contained excessive sugar. *See LeGrand v. Abbott Lab 'ys*, 2023 WL 1819159, at *10 (N.D. Cal.
 4 Feb. 8, 2023); *Johnson-Jack v. Health-Ade LLC*, 587 F. Supp. 3d 957, 968 (N.D. Cal. 2022);
 5 *Milan v. Clif Bar & Co.*, 2019 WL 3934918, at *2 (N.D. Cal. Aug. 20, 2019); *Krommenhock v.*
 6 *Post Foods, LLC*, 255 F. Supp. 3d 938, 964 (N.D. Cal. 2017).

7 Having considered the conclusions and accompanying facts in above-cited cases, this
 8 Court finds the reasoning in *Clark*, *Yoshida*, and *Truxel*—as well as Chief Judge Seeborg’s
 9 opinion in *Sprout Foods*—to be more persuasive. Accordingly, the Court holds that the mere
 10 presence of added sugar in a food product and the potential developmental concerns arising from
 11 pouch usage do not render Defendant’s claims of a “happy & health start” deceptive to a
 12 “reasonable consumer.”

13 2. FDA Administrative History

14 Plaintiffs also appear to contend that nutrient content claims would be *per se* misleading,
 15 so long as they are on products intended for children under two. Opp. 15. Plaintiffs and their
 16 cited case authorities rely almost entirely on the FDA’s 1991 commentary for this proposition.
 17 FAC ¶ 45; Opp. 12–13. However, after examining the complete administrative record, the Court
 18 found that the FDA had since retreated from the excerpts that Plaintiffs rely on, and the regulatory
 19 history does not support Plaintiffs’ theory of deception.

20 To begin, the quote Plaintiffs rely on originated from the FDA’s commentary to a 1991
 21 *Proposed Rule*, which reads:

22 Similarly, since many consumers have a limited knowledge and understanding of
 23 the amounts of nutrients that are recommended for daily consumption, a statement
 24 declaring that the product contained a specified amount of a nutrient **could be**
 25 **misleading**. By its very presence, such a statement could give consumers who were
 unfamiliar with the dietary recommendations the false impression that the product
 would assist them in maintaining healthy dietary practices relative to the amount of
 the nutrient consumed when it, in fact, would not.

26 Food Labeling: Nutrient Content Claims, 56 Fed. Reg. 60,421, 60,426 (proposed Nov. 27, 1991)
 27 (emphasis added). However, after receiving several comments during the notice-and-comment

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1 period, the FDA was persuaded to change its position considerably, recognizing instead that
 2 nutrient content claims are “generally useful to consumers” and acknowledging that the Proposed
 3 Rule would have been “too restrictive”:

4 These comments have convinced the agency to reconsider the proposed provisions
 5 for statements concerning the amount and percentage of nutrients in foods. **The**
 6 **agency believes that statements relating the amount and percentage of**
 7 **nutrients in foods are generally useful to consumers for such purposes as**
 8 **pointing out the level of a nutrient in the food and facilitating comparisons**
 9 **between foods.** The proposed provisions for amount and percentage statements
 would have limited the use of these statements to only foods that are “low” or
 “high” in the particular nutrient. **FDA believes that the provisions in the**
proposal were too restrictive because they would deny consumers the use of
such statements to evaluate many foods.

10 Food Labeling: Nutrient Content Claims, 58 Fed. Reg. 2302, 2309 (Jan. 6, 1993) (emphasis
 11 added). Moreover, the FDA’s Final Rule expressly departed from the “complete prohibition of
 12 nutrient content claims on foods for infants and children less than 2 years of age,” further
 13 undermining the support for Plaintiffs’ broad legal proposition:

14 In response to the comments, FDA has reconsidered the propriety of nutrient
 15 content claims on foods specifically intended for infants and children less than 2
 16 years of age. **The agency now believes that the complete prohibition of nutrient**
 17 **content claims on foods for infants and children less than 2 years of age may**
 18 **have been overly broad.** Although current dietary recommendations for
 Americans do not include infants and children less than 2 years of age, **there is no**
basis in the 1990 amendments to limit nutrient content claims to only foods
intended for the population over the age of 2.

19 Food Labeling, 58 Fed. Reg. at 2304.

20 Plaintiffs’ proposition is further eroded by the FDA’s express recognition that absolute
 21 nutrient claims—*e.g.*, “100 calories,” “5 grams of fat,” and the claims Plaintiffs challenge here,
 22 FAC ¶¶ 16–19—convey “no implied characterization of the level of the nutrient.” Food Labeling,
 23 58 Fed. Reg. at 2310. This observation ultimately led the FDA to include a provision in the Final
 24 Rule that permitted any nutrient content statement that “does not in any way implicitly
 25 characterize the level of the nutrient in the food and [] is not false or misleading in any respect.”
 26 21 C.F.R. § 101.13(i)(3). If, as Plaintiffs suggest, the FDA had believed *all* quantitative nutrient
 27 claims to be *per se* misleading, it would not have needed to clarify that this safe harbor did not

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1 extend to “false or misleading” statements. Indeed, if it had shared Plaintiffs’ understanding, the
 2 FDA would not have carved out a safe harbor for these quantitative nutrient content claims at all.⁴

3 As the administrative history indicates, Plaintiffs’ reliance on the commentary from the
 4 FDA’s proposed rule is misplaced, at best. Although the FAC and Plaintiffs’ opposition represent
 5 the FDA’s pre-comment positions as the agency’s definitive statements and warnings (FAC ¶¶ 45–
 6 49), the complete administrative record reflects the agency’s retreat from the statements that
 7 Plaintiffs rely on. *See* Food Labeling, 58 Fed. Reg. at 2309. To that end, Plaintiffs’ attempt to
 8 characterize the FDA’s pre-comment positions as the agency’s support for the *per se*
 9 deceptiveness of nutrient content claims is neither persuasive nor well-taken. Accordingly, the
 10 Court finds that Plaintiffs’ references to 30-year-old excerpts in the FDA’s administrative history
 11 does not give rise to any inference that nutrient claims are *per se* misleading or deceptive.⁵

12 * * *

13 In summary, the Court finds that Plaintiffs’ new theory of deception—that the Products
 14 claim they will yield “physical health benefit” but may result in harmful effects from added sugar
 15 and pouch usage—do not state a claim for fraud under the “reasonable consumer” standard. The
 16 Court does not find that a reasonable consumer exposed to the purported “physical health benefit”
 17 claims would be deceived as to other attributes of the Products, such as their sugar content or
 18 packaging. Accordingly, Plaintiffs’ FAL, CLRA, common law fraud, and UCL “fraud” claims are
 19 DISMISSED. Because Plaintiffs have already had an opportunity to amend these claims and have
 20 been unable to do so successfully, the Court finds that leave to amend would be futile. These
 21 dismissals, therefore, shall be WITHOUT LEAVE TO AMEND.

22 _____
 23 ⁴ It is worth noting that even Judge Chhabria—who had once suggested that any label that
 24 includes a prohibited nutrient content claim is *per se* misleading, *Howard v. Hain Celestial Grp.,*
 25 *Inc.*, 2022 WL 11044721, at *3 (N.D. Cal. Oct. 19, 2022)—has since receded from that position.
 26 *See Howard v. Gerber Prod. Co.*, 2023 WL 2716583, at *3 (N.D. Cal. Mar. 29, 2023) (“Even
 27 relying on the FDA’s guidance, it’s a step too far to say that a plaintiff can bring a claim sounding
 28 in fraud any time a product includes a prohibited nutrient content claim.”).

⁵ To clarify, the Court is not opining that Defendant’s nutrient content claims are compliant with
 FDA’s regulations; Plaintiffs’ “unlawful” UCL claim may still proceed based on the Sherman Law
 violations. The Court merely finds that the FDA’s 1991 observations do not support allegations
 that nutrient content claims on products for children under 2 are misleading, in and of themselves.

C. UCL “Unfair” Prong

Defendant moves to dismiss Plaintiffs’ UCL claim to the extent they assert a claim under the “unfair” prong. Mot. 19–20; FAC ¶¶ 125, 134. The Court previously dismissed the “unfair” UCL claim because Plaintiff did not respond to Defendant’s argument to dismiss. Sept. 7 Order 9 n.2. Plaintiffs again marshal no argument against Defendant’s motion to dismiss the UCL “unfair” prong, and the Court again GRANTS Defendant’s motion, this time WITHOUT LEAVE TO AMEND.

D. Unjust Enrichment

Defendant moves to dismiss the FAC’s unjust enrichment claim but does not assert any independent arguments, only that the claim should be dismissed “to the same extent as the remaining claims.” Mot. 24–25. In its earlier order, the Court had denied Defendant’s motion to dismiss based on the derivative nature of the argument and the fact that Plaintiff’s UCL “unlawful” claim survived. Sept. 7 Order 13–14. This reasoning continues to apply with equal force in the instant motion, where Defendant has not asserted a separate argument to dismiss the unjust enrichment claim and the Court again permitted the UCL “unlawful” prong to proceed.

Accordingly, the Court reaches the same conclusion in the Sept. 7 Order and DENIES Defendant’s motion to dismiss the unjust enrichment claim.

E. Equitable and Injunctive Relief

Finally, Defendant also moves to dismiss Plaintiffs’ request for equitable and injunctive relief on the basis that money damages would be an adequate remedy at law. Mot. 20–21. In its prior order, the Court rejected Defendant’s arguments regarding equitable relief and injunctive standing. Sept. 7 Order 14. Specifically, the Court found that, because Plaintiff’s claims for damages were inadequately pled and dismissed, “Defendant’s arguments concerning the availability of legal remedies under the CLRA and common law fraud claims [were] moot.” *Id.* To the extent Defendant’s prior arguments are raised here, the Court again finds them to be moot.

Defendant, however, also contends for the first time that Plaintiffs have failed to plausibly allege that they would likely purchase the Products again in the future because (1) health-

1 conscious consumers would not buy purportedly “harmful” products just because they were
2 relabeled; (2) Plaintiffs are now on notice of the Products’ composition and cannot claim that they
3 would be deceived again; and (3) Plaintiffs’ children have aged out of the age group for the
4 Products. Mot. 21–24.

5 Defendant’s first argument asks the Court to directly disbelieve the FAC’s allegations
6 regarding Plaintiffs’ intent to purchase the Products again in the future if they were properly
7 labeled. FAC ¶¶ 80, 86. This contravenes the standard for Rule 12(b)(6) motions for which the
8 Court must accept as true all well-pleaded factual allegations, *see, e.g., Ashcroft v. Iqbal*, 556 U.S.
9 at 679, and the Court declines Defendant’s invitation to do so otherwise. Defendant’s second
10 theory—that Plaintiffs cannot be deceived in the future because they are now on notice—has been
11 squarely rejected by the Ninth Circuit’s decision in *Davidson v. Kimberly-Clark Corp.*, 889 F.3d
12 956, 969 (9th Cir. 2018) (“We hold that a previously deceived consumer may have standing to
13 seek an injunction against false advertising or labeling, *even though the consumer now knows or*
14 *suspects that the advertising was false at the time of the original purchase.*”) (emphasis added).
15 And finally, with respect to Defendant’s third argument, Plaintiffs’ alleged intentions to purchase
16 the Products sufficiently allege a concrete and particularized risk of future informational injury.
17 *Cf. id.* at 971 (“[The Ninth Circuit] recognizes a history of lawsuits based on similar informational
18 injuries.”). To the extent Defendant asserts that Plaintiffs’ children have aged out of the Products’
19 “children under 2” age group during the course of this litigation, such an argument would appear
20 to run afoul of the “capable of repetition, yet evading review” exception to mootness. *See, e.g.,*
21 *Fed. Election Comm’n v. Wisconsin Right To Life, Inc.*, 551 U.S. 449, 462 (2007) (“The exception
22 applies where ‘(1) the challenged action is in its duration too short to be fully litigated prior to
23 cessation or expiration, and (2) there is a reasonable expectation that the same complaining party
24 will be subject to the same action again.’”).

25 In sum, Defendant’s arguments to dismiss the FAC’s requests for equitable and injunctive
26 relief are unavailing. The Court DENIES Defendant’s motion to the extent it seeks to dismiss the
27 FAC’s request for injunctive relief arising from the UCL “unlawful” claim.

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United States District Court
Northern District of California

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IV. CONCLUSION

Based on the foregoing, Defendant’s motion is GRANTED IN PART and DENIED IN PART, as follows:

1. The First, Second, and Third Claims are DISMISSED WITHOUT LEAVE TO AMEND;
2. The Fourth Claim is DISMISSED WITHOUT LEAVE TO AMEND as to the UCL “fraudulent” and “unfair” prongs;
3. Defendant’s motion is DENIED IN PART as to the Fourth Claim, to the extent it alleges a claim under the UCL “unlawful” prong based on Sherman Law violations;
4. The Fourth Claim is DISMISSED WITH LEAVE TO AMEND to the extent it alleges a UCL “unlawful” claim premised on the FDA’s fortification policy and DISMISSED WITHOUT LEAVE TO AMEND as to the Products listed in Section III.A.2 above;
5. Defendant’s motion is DENIED as to the Fifth Claim.

Any amendments to the FAC shall be filed within twenty-one (21) days of this Order.

IT IS SO ORDERED.

Dated: September 29, 2023



EDWARD J. DAVILA
United States District Judge