

CAED

(PS) Adams v. Abbott Laboratories

caed (March 24, 2025)

OPINION

(PS) Adams v. Abbott Laboratories

PER CURIAM.

1 2 3 4 5 6 7

8 UNITED STATES DISTRICT COURT

9 FOR THE EASTERN DISTRICT OF CALIFORNIA

10 11 ROLAND ADAMS, Case No. 2:24-cv-0555-DC-JDP (PS) 12 Plaintiff,

13 v. FINDINGS AND RECOMMENDATIONS

14 ABBOTT LABORATORIES,

15 Defendant. 16

17 18 Plaintiff Roland Adams brings this action for products liability against Defendant Abbott 19
Laboratories. Defendant moves for a second time to dismiss the complaint on the basis that it 20
fails to state a claim under Federal Rules of Civil Procedure 12(b)(6). For the reasons stated 21
below, I recommend that defendant's motion be granted and the complaint be dismissed with 22
leave to amend. 23 Procedural History 24 Plaintiff initiated this action in Sacramento Superior
Court, and defendant removed it on

25 February 22, 2024. ECF No. 1. Plaintiff alleged claims for strict products liability, negligence,
26 and breach of implied warranty. Id. Following removal, defendant moved to dismiss pursuant
to 27 Federal Rules of Civil Procedure 12(b)(6) & (8). ECF No. 4. The court granted defendant's
28 1 motion and gave plaintiff the opportunity to amend his complaint. ECF Nos. 17 & 21.
Plaintiff 2 has done so, ECF No. 18, and defendant moves again to dismiss, ECF No. 26. 3
Allegations 4 Plaintiff's amended complaint states that he suffers from diabetes and previously
used the

5 Libre 2 Sensor, manufactured by defendant, to monitor his diabetic symptoms. ECF No. 18 at 2.
6 In 2022, plaintiff's doctor changed his prescription to the upgraded Libre 3 sensor in an effort to
7 expanding the monitoring of plaintiff's diabetic symptoms. Id. Plaintiff first tried to use the new
8 sensor on December 11, 2022.¹ Id. In accordance with defendant's insurer's recommendation 9
and as advertised in defendant's promotional materials, plaintiff placed the sensor near the back 10
of his upper arm. Id. at 2-3. Shortly after he placed the sensor, plaintiff noticed blood on his shirt
11 and, once he removed the sensor, he saw that the sensor was completely covered in blood. Id.
at 12 3. Plaintiff sought medical treatment after using the sensor; a registered nurse prescribed
plaintiff 13 extra strength pain medication and antibiotics. Id. at 4. He was charged \$30 for the

treatment. 14 Id. 15 Following the incident, plaintiff sent the sensor back to defendant and filed an insurance 16 claim. Id. Before doing so, plaintiff examined the sensor and noticed that it was “atypical, 17 misshapen, and abrasive” and had “an irregular and jagged filament.” Id. at 3-4. Defendant’s 18 insurer denied plaintiff’s claim, stating that the sensor presented no issues after testing and that 19 plaintiff’s injury was likely due to incorrect placement of the device to high on plaintiff’s arm.

20 Id. at 4.

21 Plaintiff alleges three causes of action: strict product liability, negligence, and breach of 22 implied warranty. Id. at 5-8. He seeks \$10,000,000 in compensatory damages. Id. at 9. 23 Legal Standards 24 “Dismissal under Rule 12(b)(6) is proper when the complaint either (1) lacks a cognizable 25 legal theory or (2) fails to allege sufficient facts to support a cognizable legal theory.” *Somers v.* 26 27 1 The amended complaint also makes a passing reference that plaintiff had a similar experience with the sensor when he used it on August 9, 2024. Id. at 4. However, that incident 28 occurred well over two years after plaintiff filed this case and thus will not be considered. 1 *Apple, Inc.*, 729 F.3d 953, 959 (9th Cir. 2013). To survive a motion to dismiss for failure to state 2 a claim, a plaintiff must allege “enough facts to state a claim to relief that is plausible on its face.” 3 *Bell Atl. Corp. v. Twombly*, 550 U.S. 544, 570 (2007). A claim has facial plausibility when a 4 plaintiff “pleads factual content that allows the court to draw the reasonable inference that the 5 defendant is liable for the misconduct alleged.” *Ashcroft v. Iqbal*, 556 U.S. 662, 678 (2009). 6 In deciding motions under Rule 12(b)(6), the court generally considers only allegations 7 contained in the pleadings, exhibits attached to the complaint, and matters properly subject to 8 judicial notice, and construes all well-pleaded material factual allegations in the light most 9 favorable to the nonmoving party. *Chubb Custom Ins. Co. v. Space Sys./Loral, Inc.*, 710 F.3d 10 946, 956 (9th Cir. 2013); *Akhtar v. Mesa*, 698 F.3d 1202, 1212 (9th Cir. 2012). In certain 11 circumstances, the court may also consider documents referenced in but not included with the 12 complaint, or that form a basis of plaintiff’s claims. See *United States v. Ritchie*, 342 F.3d 903, 13 907 (9th Cir. 2003). 14 Discussion 15 The first amended complaint alleges three causes of action: strict products liability, 16 negligence, and breach of warranty. ECF No. 1. For the foregoing reasons, I find that the 17 amended complaint has failed to state a claim and recommend that this action be dismissed with 18 leave to amend. 19 Strict Products Liability: Design Defect 20 Plaintiff has advanced a theory of design defect. See ECF No. 18 at 5. Defendant asserts 21 three independent arguments for why plaintiff’s strict product liability claim fails. Defendant 22 argues: first that California does not recognize a design defect claim for prescription medical 23 devices. ECF No. 26 at 5. Second, that plaintiff has not alleged that there was an actual defect in 24 the design of the product. Id. at 6. And finally, that plaintiff has not plausibly alleged a design 25 defect under either the Risk Benefit Test or the Consumer Expectations Test. Id. at 7. In 26 response, plaintiff asserts that his pro se status should shield him from being held to the same 27 standard as an attorney and that this case cannot be properly adjudicated with discovery. ECF

28 No. 28.

1 A design defect claim requires that a product be built in accordance with its intended 2 specifications, and that the design itself is inherently defective. *Barker v. Lull Eng'g Co.*, 20 3 Cal.3d 413, 429 (1978). To be actionable under the design defect theory of product liability, a 4 product design must be defective under either the Consumer Expectations Test or the Risk- 5 Benefit Test. *Soule v. Gen. Motors Corp.*, 8 Cal. 4th 548, 566-67 (1994). The Consumer 6 Expectations Test considers whether the product “fail[s] to perform as safely as its ordinary 7 consumers would expect when used in an intended or reasonably foreseeable manner.” *Lucas v. 8 City of Visalia*, 726 F. Supp. 2d 1149, 1154 (E.D. Cal. 2010). Alternatively, the Risk-Benefit 9 Test considers whether “the design embodies ‘excessive preventable danger,’ that is, the risk of 10 danger inherent in the design outweighs the benefits of such design.” *Id.* Of particular relevance 11 here, California law precludes strict liability predicated on a theory of design defect against 12 manufacturers of prescription medical devices. See *Garrett v. Howmedica Osteonics Corp.*, 214 13 Cal. App. 4th 173, 182 (2013) (recognizing “an exemption from design defect strict products 14 liability for all implanted medical devices that are available only through the services of a 15 physician”). 16 Plaintiff alleges that the sensor was prescribed by his physician and that it is a 17 medical device. ECF No. 18 at 2. As noted above, California law does not permit a strict 18 liability claim in such circumstances. See *Higginbottom v. Dexcom, Inc.*, 744 F. Supp. 3d 1058, 1084 (S.D. Cal. 19 2024) (finding plaintiff could not proceed with design defect claims relating to his physician- 20 prescribed insulin pump because “California law precludes strict liability claims based on a 21 design defect with respect to manufacturers of prescription medical devices”) (citing *Garrett*, 214 22 Cal. App. 4th at 182); see also *Fussy v. RTI Surgical*, 1:21-cv-01397-DAD-BAK, 2022 WL 23 1122615, at *3 (E.D. Cal. Apr. 14, 2022) (dismissing strict liability design defect claim for a 24 prescribed medical device). Plaintiff’s strict liability claim for design defect should be dismissed. 25 Even if the court were to consider plaintiff’s design defect claim, plaintiff has not alleged 26 that the design of the sensor was defective; rather, he alleges that the sensor itself was defective 27 because it was “atypical, misshapen, and abrasive” and had “an irregular and jagged filament.” 28 1 ECF No. 18 at 3-4. Because plaintiff’s claim lacks the necessary element of a flawed design, his 2 claim fails for this reason too. 3 Negligence: Manufacturing Defect2 4 Defendant argues that the complaint fails to allege that defendant caused the defect. ECF

5 No. 26 at 9. Defendant points out that plaintiff’s conclusory assertion that his injury was “caused 6 by the unusual shape and nature of the Libre 3 filament” fails to connect the alleged defect to 7 defendant’s negligence. Moreover, defendant argues that plaintiff’s theory of *res ipsa loquitur* is 8 misplaced since the complaint is devoid of any allegation that the sensor was under defendant’s 9 exclusive control. *Id.* at 10. Plaintiff again raises no specific arguments in his opposition but 10 argues that his pro se status warrants a more lenient pleading standard and that discovery is 11 necessary for resolution of these claims. ECF No. 28. 12 A negligence claim under California law

requires a plaintiff to allege that the defendant 13 “owed [the plaintiff] a legal duty, breached the duty, and that the breach was a proximate or legal 14 cause of [the plaintiff’s] injury.” *Gonzalez v. Autoliv ASP, Inc.*, 154 Cal. App. 4th 780, 793 15 (2007). “In the context of a products liability lawsuit, [u]nder a negligence theory, a plaintiff 16 must also prove . . . that the defect in the product was due to negligence of the defendant.” *Id.* 17 (quoting *Merrill v. Navegar, Inc.*, 26 Cal. 4th 465, 479 (2001)) (quotations omitted). 18 A manufacturing defect claim is predicated on a theory that “a suitable design is in place, 19 but that the manufacturing process has in some way deviated from that design.” *In re 20 Coordinated Latex Glove Litig.*, 99 Cal. App. 4th 594, 613, 99 (2002). “A product has a 21 manufacturing defect if it differs from the manufacturer’s intended result or from other ostensibly 22 identical units of the same product line.” *Garrett*, 214 Cal. App. 4th at 190, (citing *Barker*, 20 23 Cal. 3d at 429).

24 Here, the complaint fails to explain how the sensor either deviated from defendant’s 25 intended design or how the sensor deviated from other seemingly identical sensors. *Lucas*, 726 F. 26 2 Although plaintiff does not explicitly state that his negligence claim is premised on a 27 theory of manufacturing defect, the court interprets plaintiff’s reference to the manufacturing process as an indication of how the claim is pled. Should plaintiff seek to premise his negligence 28 claim on an alternate basis, he should take care to say so in his amended complaint.

1 Supp. 2d at 1155. Plaintiff alleges that the sensor was shaped differently than the prior model 2 (the Libre 2 sensor), but he does not explain whether the sensor he used was different from other 3 Libre 3 sensors or whether it worked differently than defendant intended. A bare allegation that 4 the sensor had a defective shape is insufficient. *Iqbal*, 556 U.S. at 678. 5 In terms of plaintiff’s *res ipsa loquitur* argument; it too does not aid his claim. *Res ipsa 6 loquitur* may in some circumstances permit an inference of negligence to be drawn from a set of 7 proven facts. *Wilson v. United States*, 645 F.2d 728, 730 (9th Cir. 1981). The traditional 8 elements needed to invoke *res ipsa loquitur* are (1) an injury-producing event occurs of a kind 9 that ordinarily does not occur in the absence of someone’s negligence; (2) the event must be 10 caused by an agency or instrumentality within the exclusive control of the defendant; and (3) the 11 event must not have been due to any voluntary action or contribution on the part of the plaintiff. 12 *Olsen v. States Line*, 378 F.2d 217, 220 (9th Cir. 1967). As defendant argues, plaintiff has not 13 alleged that the sensor was under defendant’s exclusive control. 14 Breach of Implied Warranty of Fitness for a Particular Purpose 15 Defendant argues that plaintiff’s breach of implied warranty claim fails for three 16 independent reasons. First, defendant argues that the product’s disclaimer bars plaintiff’s claim.³ 17 ECF No. 26 at 11. Second, defendant argues that even it had not disclaimed the implied 18 warranty, plaintiff has failed to allege vertical privity, as required by California law. *Id.* at 12. 19 And finally, defendant argues that plaintiff has not alleged that he used the sensor for anything 20 other than its ordinary purpose. *Id.* 21 “An implied breach of warranty of fitness for a particular purpose arises only where 22 (1) the purchaser at the time of contracting intends to use

the goods for a particular purpose, (2) the seller at the time of contracting has reason to know of this particular purpose, (3) the buyer relies on the seller's skill or judgment to select or furnish goods suitable for the particular purpose, and (4) the seller at the time of contracting has reason to know that the buyer is relying on such skill and judgment." Keith v. Buchanan, 173 Cal. App. 3d 13, 25 (1985). "A particular purpose differs from the ordinary purpose for which the goods are used in that it envisages a specific use by the buyer which is peculiar to the nature of his business whereas the ordinary purposes for which goods are used are those envisaged in the concept of merchantability and go to uses which are customarily made of the goods in question." Am. Suzuki Motor Corp. v. Superior Ct., 37 Cal. App. 4th 1291, 1295 n.2 (1995) (internal quotation marks omitted). To state a claim for breach of an implied warranty for a particular purpose, a plaintiff must identify the particular purpose for which he obtained the product. Plaintiff here alleges that he intended to use the sensor to monitor his diabetes, which, according to the complaint's allegations, is the product's ordinary use. ECF No. 18 at 2 ("The sensor uses a non-injurious filament to penetrate a patient's skin and monitor for diabetic symptoms."). Therefore, plaintiff has not stated a claim for breach of the implied warranty of fitness for a particular purpose. See Smith v. LC Electronics U.S.A, Inc., 2014 WL 989742, at *8 ("[P]laintiff has identified no particular purpose for which she purchased the washing machine. She purchased it to wash her laundry, which is the ordinary purpose of a washing machine.") (internal quotation marks omitted); Kent v. Hewlett-Packard Co., No. 09-cv-05341-JF, 2010 WL 2681767, at *5 (N.D. Cal.

19 July 6, 2010) ("Plaintiffs have not alleged that they used the computers . . . for anything other than their ordinary purpose. Thus, plaintiffs have not stated a claim for breach of an implied warranty for a particular purpose."). Moreover, this claim fails because the complaint does not allege that privity exists between plaintiff and defendant. "Under California law, the general rule is that privity of contract is required in an action for breach of either express or implied warranty and that there is no privity between the original seller and a subsequent purchaser who is in no way a party to the original sale." In re Clorox Consumer Litig., 894 F. Supp. 2d 1224, 1236 (N.D. Cal. 2012) (internal quotation marks omitted) (citing Burr v. Sherwin Williams Co., 42 Cal.2d 682, 695 (1954)). "A buyer and a seller stand in privity if they are in adjoining links of the distribution chain"; "[t]hus, an end consumer . . . who buys from a retailer is not in privity with the manufacturer." Clemens v. DaimlerChrysler Corp., 534 F.3d 1017, 1023 (9th Cir. 2008) (citing Osborne v. Subaru of Am. Inc., 198 Cal. App. 3d 646, 656 n.6, (1988)). Accordingly, it is RECOMMENDED that defendant's motion to dismiss, ECF No. 26, be GRANTED and plaintiff be granted thirty days from the date of any order adopting these findings and recommendations to file an amended complaint. These findings and

recommendations are submitted to the United States District Judge 8 | assigned to the case, pursuant to the provisions of 28 U.S.C. § 636(b)(). Within fourteen days of 9 | service of these findings and recommendations, any party may file written objections with the 10 | court and serve a copy on all parties. Any such document should be captioned “Objections to 11 | Magistrate Judge’s Findings and Recommendations,” and any response shall be served and filed 12 | within fourteen days of service of the objections. The parties are advised that failure to file 13 | objections within the specified time may waive the right to appeal the District Court’s order. See 14 | Turner v. Duncan, 158 F.3d 449, 455 (9th Cir. 1998); Martinez v. Yist, 951 F.2d 1153 (9th Cir. 15 1991). 16 7 IT IS SO ORDERED. 18 (1 Oy — Dated: _ March 21, 2025 Q———

19 JEREMY D. PETERSON

UNITED STATES MAGISTRATE JUDGE

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