



November 17, 2023

MASS TORTS
IDAHO STATE BAR – LITIGATION SECTION
NOVEMBER 17, 2023

ENRIQUE G. SERNA

www.serna-associates.com

www.enriquegserna.com

enrique@serna-associates.com

210-865-5800 - cell

YOU TUBE

Social Media - Product

<https://studio.youtube.com/video/cs5UnVqoUCk/>

Hair Relaxer Product

https://studio.youtube.com/video/qQD_MBbcRlk/



IDAHO – MASS TORTS – RULES AND CASE EXAMPLES

1. COMPARISON WITH RULE 23 OF THE FEDERAL RULES OF CIVIL PROCEDURE

The Idaho Rule of Civil Procedure governing class actions was amended July 1, 2016, to move from Rule 23 to Rule 77-79. This was part of a larger renumbering with the result that the class action rules are grouped with other special proceedings. The Idaho Rules are substantially similar, except for their location, to the federal rule though Idaho does not have an analog to Red. R. Ci. 23(f). The Idaho Supreme Court considers “federal authority . . . relevant” because the Idaho rules on class actions were based on the federal rules. *O’Boskey v. First Fed. Sav. & Loan Ass’n. of Boise*, 112 Idaho 1002, 1005 n.2, 739 P.2d 301, 304 n.2 (1987).

2. RULE 77 OF THE IDAHO RULES OF CIVIL PROCEDURE (FORMERLY RULE 23)

Class actions are generally favored where the interests of absent class members can be adequately represented by the named plaintiff and the requirements of the rule are met. *Bush v. Upper Valley Telecable Co.*, 96 Idaho 83, 89, 524 P.2d 1005, 1061 (1974).

Certification of a class action is governed by the Idaho Rules of Civil Procedure, *Bettweiser v. N.Y. Irrigation Dist.*, 154 Idaho 317, 324, 297 P.3d 1124, 1141, and in particular Rule 77, *Zeyen v. Pocatello/Chubbock Scho. Dist. No. 25*, 165 Idaho 690, 697, 451 P.3d 25, 32 (2019). To certify a lawsuit as a class action, the trial court must find that all four factors in Rule 77(a) (at the time Rule 23(a)) exist and that at least one factor in Rule 77(b) (at the time Rule 23(b)) exists. *BHA Invs., Inc. v. City of Boise*, 141 Idaho 168, 171-72, 108 P.3d 315, 318-319 (2004). If the court finds that a necessary factor is absent, the court need not address the other factors. *Id.*

3. RULE 77(A) OF THE IDAHO RULES OF CIVIL PROCEDURE (FORMERLY RULE 23(A))

The first factor in Rule 77(a) (at the time Rule 23(a)) is that “the class is so **numerous** that joinder of all members is impracticable.” *BHA Invs., Inc. v. City of Boise*, 141 Idaho 168, 172, 180 P.3d 315, 319 (2004). In BHS Investments, the Idaho Supreme Court found that the “district court did not abuse its discretion in deciding that seventeen known entities located with the City did not constitute a class that was so numerous that joinder of all members is impracticable.” *Id.*



A plaintiff is not required to file a case as a class action simply because the right he seeks to establish is also held by a number of other individuals, but instead is allowed to choose to bring an individual action. *Camp Easton Forever, Inc. v. Inland Nw. Council BSA*, 156 Idaho 893, 898, 332 P.3d 805 (2014); *Middlekauff v. Lake Cascade, Inc.* 110 Idaho 909, 915, 719 P.2d 1169, 1175 (1986). Rule 77(a) et seq. (at the time Rule 23(a) es seq.) merely establish the conditions under which one “or more members of a class may sue or be sued as representative party on behalf of all.” *Id.* (emphasis in the original: citation omitted).

MASS TORT NEWS

L'Oreal, Revlon Can't Escape Hair Relaxer-Cancer Link MDL

Law360 (November 13, 2023, 11:52 PM EST) -- Consumers who allege L'Oreal, Revlon and other companies' hair relaxers — which are primarily marketed to Black women — cause breast, uterine and other cancers can pursue most of their consolidated claims, an Illinois federal judge ruled Monday.

U.S. District Judge Mary M. Rowland largely denied the companies' bids to dismiss the multidistrict litigation that now comprises more than 8,000 cases, dismissing four fraud-based claims, but keeping alive the bulk of the claims, including negligence, strict liability for design defect and failure to warn, breach of warranty, unjust enrichment and wrongful death, according to the order.

To start, Judge Rowland rejected the companies' argument that eight of the plaintiffs' 15 claims — including claims for negligent misrepresentation, fraudulent misrepresentation and unjust enrichment — are preempted by the Food Drug and Cosmetic Act, saying the companies haven't shown that the plaintiffs want to impose labeling requirements that are different from those imposed by the federal law, the order states.



Turning to the plaintiffs' negligence and strict liability claims, the judge rejected the companies' argument that the plaintiffs haven't sufficiently stated a general negligence claim with their allegations that the defendants had "a duty to exercise reasonable care in the manufacturing, designing ... selling and distributing of their hair relaxer products," according to the order.

"The allegations read as a whole give rise to the inference that defendants' conduct proximately caused plaintiffs' injuries: more is not needed at this stage," Judge Rowland said, noting that the consumers also alleged that the companies "knew or should have known that phthalates and other [endocrine-disrupting chemicals] in their hair relaxer products significantly increase the risk of cancers and other negative health conditions."

The judge also rejected the companies' contention that the consumers' allegations that the hair relaxer products "were defectively designed because they caused serious injuries and death" are conclusory, the order states. Rather, the consumers have specifically alleged the products contained toxic chemicals that significantly increase the risk of cancers, the judge said.

The lawsuits, which were consolidated in Illinois in February, generally accuse L'Oreal and other companies of selling hair relaxation, or lanthionization, products that caused or increased the risk of women developing uterine, ovarian or breast cancer, endometriosis, uterine fibroids or other injuries to the reproductive system.

The hair relaxer products at issue can be applied by cosmetologists or at home, and they temporarily straighten and smooth hair in a process that can cause burns and lesions on the scalp and allow hair relaxer chemicals to enter the body, according to the plaintiffs.

Judge Rowland in March tapped Ben Crump Law Firm, Motley Rice LLC, DiCello Levitt and Douglas & London attorneys to serve as co-lead counsel.

Following the publication last October of a major report from the National Institutes of Health that found women who frequently used hair relaxer products are more than twice as likely to develop uterine cancer as those who don't, suits were launched across the country, including in New York, California, Georgia, Tennessee and Ohio.

The Tennessee lawsuit refers to the history of hair relaxers in America, saying the practice of straightening African American hair began during slavery with the use of hot combs to press the hair straight. Chemical relaxers were created in the early 1900s, according to the suit. Today, millions of Black women use hair relaxers, potentially exposing them to chemicals through scalp lesions and burns, the suit asserts.



The companies moved to dismiss the consolidated litigation in July, according to court records.

In her order Monday, Judge Rowland largely kept the consumers' claims intact, but she did dismiss their fraud-based claims — including negligent misrepresentation, fraud or fraudulent misrepresentation and fraudulent concealment — finding that they hadn't met the heightened pleading standard of stating "with particularity the circumstances constituting fraud."

The consumers urged the judge to focus on the defendants' alleged omissions and not reach the issue of affirmative misrepresentations, which they argue can be taken care of in the individual cases, according to the order. But Judge Rowland said the consumers' fraud-based claims are largely based on alleged misrepresentations.

"They do allege that defendants concealed material facts about the dangers of their products, but do not identify specific omissions of material fact or support these allegations with additional detail," the judge said.

"The allegations in the complaint do not show the concealment was 'more than a mere passive omission of facts during a business transaction' that was 'done with the intention to deceive under circumstances creating an opportunity and duty to speak,'" she added.

The judge also dismissed India-based Dabur International Ltd. from the litigation for lack of personal jurisdiction.

"We feel positive about the order, and most importantly, it gives our clients the opportunity to move forward and seek justice from the defendants," Diandra "Fu" Debrosse Zimmermann of DiCello Levitt LLC, an attorney for the consumers, told Law360 late Monday. "We represent a lot of women who have been devastated by these products."

Counsel for the other parties did not immediately respond to requests for comment.

The plaintiffs are represented by Diandra "Fu" Debrosse Zimmermann of DiCello Levitt LLC, Benjamin L. Crump of Ben Crump Law, Fidelma L. Fitzpatrick of Motley Rice LLC, Michael A. London of Douglas & London PC and Edward A. Wallace of Wallace Miller, among others.

L'Oreal is represented by Gordon Rees Scully Mansukhani LLP and Ellis George Cipollone O'Brien LLP.

Revlon is represented by Paul Weiss Rifkind Wharton & Garrison LLP and Hawkins Parnell & Young LLP.



Dabur International Ltd. is represented by Kirkland & Ellis LLP.

The case is In re: Hair Relaxer Marketing Sales Practices and Products Liability Litigation, case number 1:23-cv-00818, in the U.S. District Court for the Northern District of Illinois.

--Additional reporting by Dorothy Atkins and Emily Field. Editing by Jay Jackson Jr.

Read more at: <https://www.law360.com/classaction/articles/1766316/l-oreal-revlon-can-t-escape-hair-relaxer-cancer-link-mdl?copied=1>

Feds Settle 3 Camp Lejeune Water Contamination Cases

By [Travis Bland](#) · [Listen to article](#)

Law360 (November 13, 2023, 5:51 PM EST) -- The U.S. government has made its first payouts to people who were poisoned by contaminated water at Camp LeJeune Marine training base, more than a year after federal legislation provided a pathway for veterans and residents of the base to sue and be compensated.

The Department of the Navy agreed to a total payment of \$850,000 to be split among three claimants, and two had accepted the settlement, according to a [transcript](#) released Thursday of an Oct. 30 federal court hearing.

The claimants were not identified in the transcript of the hearing, but one received \$250,000 and the two others \$300,000 each.

The payments will come from the U.S. Treasury and not the Marine or Navy budget, according to the transcript.

The settlements were part of an elective program offered by the Navy, which allows people suing the department to take an early payout to end their lawsuits, an attorney for the



government said.

In addition to the three settlement offers, the Navy has made 20 other offers, [U.S. Department of Justice](#) attorney Adam Bain told the court.

"That process is just beginning," Bain said about the elective settlement program at the hearing. "The Department of Justice and the Department of the Navy have been working with the plaintiff's leadership committee to get the information that we need to determine eligibility for settlement offers under the program."

As of the end of October, 1,309 cases have been filed, according to Bain. Those lawsuits alleged that Camp Lejeune's contaminated water caused cancer, miscarriages and other health problems. An additional 117,000 administrative claims have been filed with the Navy, Bain said.

Word of the settlements came little more than a week before the Department of Veterans Affairs announced that it would cover Parkinson's disease care costs for veterans' family members who lived on the base.

Trials in litigation over contaminated drinking water on the Camp Lejeune Marine base are expected to begin next year in North Carolina federal court, involving plaintiffs who said the water caused their bladder cancer, kidney cancer, lymphoma and other diseases, according to court filings.

The sprawling litigation surrounding the Camp Lejeune Marine base, in coastal North Carolina, stems from Congress' passage of the Camp Lejeune Justice Act last year. Judges in the cases have said up to a million lawsuits may be filed.

Those suits involve nearly 40 diseases said to be connected to the water at Camp Lejeune, which allegedly contained excessive levels of chlorinated solvents and other contaminants. People living on the base between 1953 and 1987 were potentially exposed to the toxins.



The U.S. government gave no comment.

The government is represented by Adam Bain of the U.S. Department of Justice.

The plaintiffs are represented by the [Bell Legal Group](#), [Keller Postman LLC](#), [Ward & Smith PA](#), the Dowling Firm and [Lieff Cabraser Heimann & Bernstein LLP](#), among others.

The cases are In Re: Camp Lejeune Water Litigation v. United States of America, case number [7:23-cv-00897](#), in the [U.S. District Court for the Eastern District of North Carolina](#).

School Attys Seek Table Seat In Social Media Addiction MDL

By [Tom Lotshaw](#) · [Listen to article](#)

Law360 (November 13, 2023, 10:03 PM EST) -- At least five firms and attorneys are asking a California federal judge overseeing multidistrict litigation accusing tech giants of injuring young people's mental health to appoint them to leadership roles to help represent the interests of school districts or local government entities that have lodged suits against the companies.

Over the past week, plaintiff leadership applications for [the MDL](#) have been filed by Anne Marie Murphy of [Cotchett Pitre & McCarthy LLP](#), Derek W. Loeser of [Keller Rohrback LLP](#), James E. Cecchi of [Carella Byrne Cecchi Brody & Agnello PC](#), Thomas P. Cartmell of [Wagstaff & Cartmell LLP](#), and Aelish M. Baig of [Robbins Geller Rudman & Dowd LLP](#).

In a letter to the court, Cartmell says co-lead counsel has requested that he serve as co-chair for local government entities and play a key role in developing trial strategies. He states his firm has several cases in the MDL, including cases filed on behalf of individual plaintiffs and on behalf of school districts, and expects to file more.



Cartmell says he represented school districts in the [Juul Labs Inc.](#) MDL, serving as government entity liaison counsel and as lead trial counsel in the only bellwether case [to reach trial](#). He adds that he also served as co-lead counsel or co-lead trial counsel in a number of other MDLs, including [In re 3M Combat Arms Earplug Products Liability Litigation](#).

"I believe I will add significant value to the [plaintiffs steering committee] based on my experience and the emphasis I place on efficiency, courtesy, civility and teamwork in any case I pursue," Cartmell tells the court.

Loeser, in his application, says he represents school districts, including Seattle Public Schools, the first district to sue social media companies in an MDL that now includes more than 200 school district plaintiffs.

While the school districts' suits will involve many of the same questions as the suits that hundreds of individual plaintiffs have filed, there will be key differences, Loeser says.

"For example, the school districts assert claims for public nuisance that are not asserted in the personal injury cases, which will require unique evidence and expert testimony, such as with respect to the harms incurred by school districts," Loeser says, noting districts also seek a different remedy to solve problems the social media companies are causing, rather than only damages for harms.

Loeser states he was appointed co-lead counsel for the [In re Facebook Inc. Consumer Privacy User Profile Litigation](#) and represents dozens of local governments in the In re National Prescription Opiate Litigation, as well as school districts and municipalities in the Juul Labs Inc. MDL.

"I believe it is important to also include counsel who exclusively represent school districts in this litigation and are solely focused on and dedicated to pursuing their interests, and I respectfully submit that I am well qualified to fulfill this role," he tells the court.



Murphy, who is helping represent a number of California school districts in litigation, also emphasizes that plaintiff leadership needs members whose attention is focused on school districts and local governments.

"This litigation needs leaders who can zealously represent the interests of school districts, while working side-by-side with plaintiffs in other tracks," Murphy says.

Cecchi said his firm is representing multiple [New Jersey school districts](#) in the litigation, and also represented many other local governments from that state in opioid multidistrict litigation. He adds that he served on steering committees and as settlement class counsel for [In re Volkswagen "Clean Diesel" Marketing, Sales Practices and Products Liability Litigation](#) and [In re Takata Airbag Products Liability Litigation](#), and has already helped this MDL move forward through the initial creation of a government entity committee.

"We believe that establishing a formal organizational framework — fully defining the roles of counsel and creating a dedicated subcommittee — will ensure that all plaintiffs' cases are prosecuted efficiently and effectively," Cecchi tells the court.

Baig says Robbins Geller Rudman & Dowd LLP is helping represent school districts including [Miami-Dade County Public Schools](#) and Broward County Public Schools — the third- and sixth-largest districts in the country, respectively — as well as Bucks County, Pennsylvania, one of the first municipalities to file suit, and the state of Arkansas, one of the first states to file suit.

Baig states she and the firm have played leading roles and represented a number of other local governments and school districts in [opioid](#) and vaping MDLs, many of which are preparing to file or considering filing lawsuits in this MDL.

"The governmental entity cases, including those brought by school districts and municipality plaintiffs, are different from the personal injury cases, with distinct claims and discovery, expert, trial and resolution needs, and they should have a dedicated team," Baig tells the court.



The applicants declined to comment or could not immediately be reached for comment on Monday. Representatives for other parties in the MDL could not immediately be reached for comment.

The plaintiffs are represented by Lexi Hazam of [Lief Cabraser Heimann & Bernstein LLP](#), Previn Warren of [Motley Rice LLC](#) and Chris Seeger of [Seeger Weiss LLP](#).

[Meta Platforms Inc.](#) is represented by Mark W. Mosier, Paul W. Schmidt, Phyllis A. Jones, Ashley Simonsen and Emily Johnson Henn of [Covington & Burling LLP](#).

[TikTok](#) is represented by Geoffrey M. Drake and David Mattern of [King & Spalding LLP](#), and Andrea Roberts Pierson and Amy Fiterman of [Faegre Drinker Biddle & Reath LLP](#).

[Snap Inc.](#) is represented by Jonathan H. Blavin, Rose L. Ehler, Victoria A. Degtyareva, Ariel T. Teshuva and Lauren A. Bell of [Munger Tolles & Olson LLP](#).

[Google LLC](#) is represented by Brian M. Willen, Lauren Gallo White, Andrew Kramer, Carmen Sobczak, Christopher Chiou, Samantha Machock and Matthew K. Donohue of [Wilson Sonsini Goodrich & Rosati PC](#).

The MDL case is In re: Social Media Adolescent Addiction/Personal Injury Products Liability Litigation, case number [4:22-md-03047](#), in the [U.S. District Court for the Northern District of California](#).

**IN THE UNITED STATES DISTRICT COURT
FOR THE NORTHERN DISTRICT OF ILLINOIS
EASTERN DIVISION**

IN RE: HAIR RELAXER
MARKETING SALES PRACTICES
AND PRODUCTS LIABILITY
LITIGATION

MDL No. 3060

Master Docket No. 23-cv-00818

Judge Mary M. Rowland

MEMORANDUM OPINION AND ORDER

In this multidistrict litigation, Plaintiffs claim that Defendants advertised, manufactured and sold toxic hair relaxer products that caused Plaintiffs to develop cancers and other injuries. Before the Court is Defendants’ joint motion to dismiss Plaintiffs’ master long form complaint. For the reasons stated herein, Defendants’ Joint Motion to Dismiss [142] is granted in part and denied in part and Defendant McBride’s motion to dismiss [192] is granted in part and denied in part.

I. Background

A. Procedural Background

In February 2023, the United States Judicial Panel on Multidistrict Litigation (the “Panel”) consolidated individual and putative class actions, then pending in nineteen districts, for pretrial proceedings in this District under 28 U.S.C. § 1407. See Transfer Order [1]¹. “Congress enacted 28 U.S.C. § 1407 in 1968 to manage more effectively complex sets of related lawsuits pending in multiple districts.” *Bell v.*

¹ Bracketed numbers refer to docket entries and referenced page numbers are from the CM/ECF header at the top of filings.

Publix Super Markets, Inc., 982 F.3d 468, 488–89 (7th Cir. 2020). Section 1407(a) gave the Panel “the power to transfer related cases to one district court for ‘coordinated or consolidated pretrial proceedings.’” *Id.*

In its February 2023 Transfer Order, the Panel noted that “[o]n October 17, 2022, a study led by the National Institutes of Health (NIH) reported findings that women who frequently used chemical hair straightening or hair relaxer products were more than twice as likely to develop uterine cancer as women who did not use such products.” *Id.* Shortly after the study was published, lawsuits were filed in different district courts. The Panel found the cases shared “common questions of fact arising from allegations that defendants’ hair relaxer products contain phthalates, including di-2-ethylhexylphthalate, or other endocrine-disrupting chemicals (EDCs), and that the use of such products caused or increased the risk of developing uterine, ovarian, or breast cancer, endometriosis, uterine fibroids, or other injuries to the reproductive system.” *Id.* The cases in this MDL now number more than 8,200.

Plaintiffs allege that their “use of toxic chemical straightening products designed or manufactured by the Defendants was a direct result of Defendants’ wrongful marketing practices.” [106 ¶ 6]. They claim that Defendants “[i]ntentionally target[ed] Black and Brown women, including Black and Brown teenaged girls and children, as customers to purchase and use their unsafe hair relaxer products.” *Id.* ¶ 280. Defendants jointly moved to dismiss the complaint in its entirety. [142]. They moved for dismissal under Rule 12(b)(6) and Rule 9, and also argue that Defendants

Dabur International and Dermoviva should be dismissed for lack of personal jurisdiction.

B. Factual Background

The factual allegations from the master long form complaint ([106], “MLC”) and short form complaint ([175-1], “SFC”) are accepted as true for the purposes of the motion to dismiss. *See Lax v. Mayorkas*, 20 F.4th 1178, 1181 (7th Cir. 2021).²

Plaintiffs allege they were injured by defective hair relaxers designed, manufactured, sold, distributed, and marketed by the Defendants in this case. [106 ¶ 5]. The MLC was filed on behalf of all Plaintiffs whose claims are subsumed within MDL 3060, and who have suffered personal injuries and death as a result of their use of Defendants’ various hair relaxer products. *Id.* ¶ 10. In their complaint Plaintiffs explain that endocrine-disrupting chemicals (“EDCs”) are chemicals, or chemical mixtures, that interfere with the normal activity of the endocrine system. *Id.* ¶ 74. They allege that natural and synthetic EDCs are present in some of Defendants’ hair relaxer products as “fragrance” and “perfumes”, and enter the body when the products are applied to the hair and scalp. *Id.* ¶ 76. One of the EDCs, phthalates, are harmful because they interferes with individuals’ natural hormone production and degradation. *Id.* ¶ 77. Widely used hair relaxers, Plaintiffs claim, are a source of exposure to carcinogens and these endocrine disrupters. *Id.* ¶ 95.

² The Court adopted the short form complaint on August 3, 2023. Pursuant to the Court’s Case Management Order (“CMO”) No. 8, “[f]or each action in the MDL, subject to this Order, the Master Complaint, together with the Short Form Complaint shall be deemed the operative complaint.” [175]. This opinion sometimes refers to the MLC and SFC together as the “complaint.”

In October 2022, the National Institutes of Health (NIH) released a study of approximately 34,000 women, aged 35-74, that was conducted over approximately 11 years. (“Chang Article”). *Id.* ¶ 85. The study revealed significantly higher rates of uterine cancer in women who had used hair relaxers. *Id.* ¶ 86. And a 2021 study funded by NIH and the National Institute on Minority Health Sciences found frequent use of hair relaxers was strongly associated with ovarian cancer (“White Article”). *Id.* ¶ 89. Plaintiffs in this MDL seek relief in the form of compensatory and punitive damages, monetary restitution, medical monitoring and equitable relief, and other remedies “as a result of injuries incurred by Defendants’ defective products and other wrongful practices.” *Id.* ¶ 1.

The master complaint contains fifteen counts: negligence and/or gross negligence (Count 1); negligent misrepresentation (Count 2); negligence *per se* (Count 3); strict liability: design defect (Count 4); strict liability: failure to warn (Count 5); breach of implied warranty of merchantability/fitness for particular use (Count 6); breach of express warranty under state law and the Magnuson-Moss Warranty Act, 15 U.S.C. § 2301 (Count 7); fraud/fraudulent misrepresentation (Count 8); fraudulent concealment (Count 9); U.S. state and territory statutory consumer protection and unfair or deceptive trade practices claims (Count 10); unjust enrichment (Count 11); wrongful death (Count 12); survival action (Count 13); loss of consortium (Count 14); and punitive damages (Count 15). The wrongful death, survival, and loss of consortium are derivative claims.

II. Standard

“To survive a motion to dismiss under Rule 12(b)(6), the complaint must provide enough factual information to state a claim to relief that is plausible on its face and raise a right to relief above the speculative level.” *Haywood v. Massage Envy Franchising, LLC*, 887 F.3d 329, 333 (7th Cir. 2018) (quoting *Camasta v. Jos. A. Bank Clothiers, Inc.*, 761 F.3d 732, 736 (7th Cir. 2014)); *see also* Fed. R. Civ. P. 8(a)(2) (requiring a complaint to contain a “short and plain statement of the claim showing that the pleader is entitled to relief”). A court deciding a Rule 12(b)(6) motion “construe[s] the complaint in the light most favorable to the plaintiff, accept[s] all well-pleaded facts as true, and draw[s] all reasonable inferences in the plaintiff’s favor.” *Lax*, 20 F.4th at 1181. However, the court need not accept as true “statements of law or unsupported conclusory factual allegations.” *Id.* (quoting *Bilek v. Fed. Ins. Co.*, 8 F.4th 581, 586 (7th Cir. 2021)). “While detailed factual allegations are not necessary to survive a motion to dismiss, [the standard] does require ‘more than mere labels and conclusions or a formulaic recitation of the elements of a cause of action to be considered adequate.’” *Sevugan v. Direct Energy Servs., LLC*, 931 F.3d 610, 614 (7th Cir. 2019) (quoting *Bell v. City of Chicago*, 835 F.3d 736, 738 (7th Cir. 2016)).

Dismissal for failure to state a claim is proper “when the allegations in a complaint, however true, could not raise a claim of entitlement to relief.” *Bell Atl. Corp. v. Twombly*, 550 U.S. 544, 558 (2007). Deciding the plausibility of the claim is “a context-specific task that requires the reviewing court to draw on its judicial experience and common sense.” *McCauley v. City of Chicago*, 671 F.3d 611, 616 (7th Cir. 2011) (quoting *Ashcroft v. Iqbal*, 556 U.S. 662, 679 (2009)).

III. Analysis

Defendants have moved to dismiss the complaint on a number of grounds. The Court begins with Defendants' preemption argument.

A. Preemption

Defendants argue that certain claims of Plaintiffs are expressly preempted by the Food, Drug, and Cosmetic Act ("FDCA"). The FDCA expressly preempts any state law "requirement for labeling or packaging of a cosmetic that is different from or in addition to, or that is otherwise not identical with, a requirement specifically applicable to a particular cosmetic or class of cosmetics under this chapter." 21 U.S.C. § 379s(a). Express preemption means that Congress has defined "explicitly the extent to which its enactments pre-empt state law." *English v. Gen. Elec. Co.*, 496 U.S. 72, 78 (1990).

Defendants concede that FDCA preemption does not apply to product liability claims. *See* 21 U.S.C. § 379s(d) ("Nothing in this section shall be construed to modify or otherwise affect any action or the liability of any person under the product liability law of any State."). They argue, however, that the non-product liability claims are expressly preempted, namely: negligent misrepresentation, breach of warranty, fraud, state consumer protection and unfair or deceptive trade practices, unjust enrichment and punitive damages (Counts 2, 6, 7, 8, 9, 10, 11 and 15). According to Defendants, these claims seek to impose labeling requirements not identical to those imposed by the FDCA and its regulations. Defendants argue they are not required to identify fragrance ingredients or identify "ingredients that are nowhere in their

products.” [142 at 24]. Plaintiffs counter that their claims are state law product liability claims and therefore exempt under 21 U.S.C. § 379s(d). In addition, Plaintiffs maintain they do not seek to impose any labeling requirement that is different from or in addition to an FDCA requirement applicable to Defendants’ products.

In this circuit, it is settled that preemption is an affirmative defense, meaning Defendants bear the burden of proof.³ *Benson v. Fannie May Confections Brands, Inc.*, 944 F.3d 639, 645 (7th Cir. 2019); *Russian Media Grp., LLC v. Cable Am., Inc.*, 598 F.3d 302, 309 (7th Cir. 2010). A plaintiff’s pleading “need not anticipate or attempt to circumvent affirmative defenses.” *Bausch v. Stryker Corp.*, 630 F.3d 546, 561 (7th Cir. 2010) (internal citation omitted). Dismissal is only appropriate when the “plaintiff pleads himself out of court by alleging facts sufficient to establish” the defense. *Sidney Hillman Health Ctr. v. Abbott Labs., Inc.*, 782 F.3d 922, 928 (7th Cir. 2015) (cleaned up). Affirmative defenses “typically turn on facts not before the court at [the dismissal] stage.” *Brownmark Films, LLC v. Comedy Partners*, 682 F.3d 687, 690 (7th Cir. 2012).

As Plaintiffs point out, a “products liability action” is defined in Black’s Law Dictionary as a lawsuit “brought against a manufacturer, seller, or lessor of a product-
-regardless of the substantive legal theory or theories on which the lawsuit is brought-
-for personal injury, death, or property damage caused by the manufacture,

³ An MDL transferee court “applies the laws of the circuit in which it sits to decide issues of federal procedure.” *In re Testosterone Replacement Therapy Prod. Liab. Litig. Coordinated Pretrial Proc.*, 164 F. Supp. 3d 1040, 1044 (N.D. Ill. 2016). *See also U.S. ex rel. Hockett v. Columbia/HCA Healthcare Corp.*, 498 F. Supp. 2d 25, 40 (D.D.C. 2007) (transferee court “follow[s] its own circuit on questions of federal law.”).

construction, design, formulation, installation, preparation, or assembly of a product. (11th ed. 2019) (emphasis added). In their reply, Defendants broadly argue that not every claim in a personal injury action is a product liability claim. They rely on *In re Acetaminophen - ASD-ADHD Prod. Liab. Litig.*, No. 22CV9011 (DLC), 2023 WL 3045802 (S.D.N.Y. Apr. 21, 2023), in which the court found Tennessee’s consumer protection claim preempted by the FDCA. *In re Acetaminophen* analyzed only Tennessee’s law, which “authorize[d] plaintiffs to bring claims based solely on economic loss.” *Id.* at *5. In Count 10 of the master complaint, Plaintiffs claim they suffered “serious injury” and “economic loss, pecuniary loss, personal injury, loss of companionship and society, mental anguish and/or other compensable injuries.” [106 ¶¶ 241, 247]. In addition, the court in *In re Acetaminophen* did not analyze any of the other claims that Defendants argue are preempted here such as negligent misrepresentation and breach of warranty.⁴ Also guiding this Court are the U.S. Supreme Court’s statements that “the historic police powers of the States were not to be superseded by the Federal Act unless that was the clear and manifest purpose of Congress.” *In Medtronic, Inc. v. Lohr*, 518 U.S. 470, 485 (1996) (cleaned up); *see also Wyeth v. Levine*, 555 U.S. 555, 579 (2009) (“the FDA long maintained that state law

⁴ Two other cases cited by Defendants are *Turek v. Gen. Mills, Inc.*, 662 F.3d 423 (7th Cir. 2011) and *Critcher v. L’Oreal USA, Inc.*, 959 F.3d 31 (2d Cir. 2020). *Turek* did not involve product liability claims or address 21 U.S.C. § 379s(d). In *Critcher* plaintiffs alleged they were injured when certain L’Oréal product labels omitted information that the creams could not be fully dispensed from their containers. *Id.* at 36. Plaintiffs did not claim, as here, that the product contained toxic chemicals or ingredients, and the Court did not analyze whether plaintiffs’ claims were exempt under Section 379s(d). The Court also finds Defendants’ implicit exemption argument underdeveloped. [142] at 24. *See M.G. Skinner & Assocs. Ins. Agency, Inc. v. Norman-Spencer Agency, Inc.*, 845 F.3d 313, 321 (7th Cir. 2017).

offers an additional, and important, layer of consumer protection that complements FDA regulation.”); *Bausch*, 630 F.3d at 557.

Thus the Court finds that Defendants have not met their burden to show that Plaintiffs’ Counts 2, 6, 7, 8, 9, 10, 11 and 15 are preempted.

B. Negligence, Design Defect and Failure to Warn Claims

Next, Defendants argue that Plaintiff’s negligence and strict liability (Counts 1, 3, 4 and 5) claims fail. Defendants’ initial contention that Plaintiffs did not identify the products they used or relevant time period for the use is not convincing. As Plaintiffs argue, the SFC is the place where each individual plaintiff identifies the particular products they used and other factual details to support their complaint. In addition to the fact that Defendants concede that Plaintiffs identify products in their SFC [142 at 24], this Court’s CMO No. 8 expressly states that for each action in this MDL, “the Master Complaint, together with the Short Form Complaint shall be deemed the operative complaint.” [175]. Moreover, the cases Defendants rely on to argue that product identification is required at this stage did not rule on motions to dismiss, *Tragarz v. Keene Corp.*, 980 F.2d 411 (7th Cir. 1992), *Zimmer v. Celotex Corp.*, 192 Ill. App. 3d 1088 (Ill. App. Ct. 1989), or applied a state-law fact-pleading standard rather than the federal notice pleading standard applicable here. *See Kozak v. Armstrong World Indus., Inc.*, 572 N.E.2d 279, 282 (Ill. App. Ct. 1991).

1. Negligence Claims

In Count 1, Plaintiffs allege that Defendants breached their duty of care to Plaintiffs by manufacturing, marketing, and selling their hair relaxer products

negligently or recklessly and by failing to adequately warn of these products' risks and dangers. Defendants argue that some jurisdictions do not recognize gross negligence or negligence *per se* as separate causes of action. The Court does not find variation in how states treat negligence claims to be a basis for dismissal at this pleading stage.

Defendants also contend that Plaintiffs have not sufficiently stated a general negligence claim. The elements of a negligence claim are: “the existence of a duty owed by the defendant to the plaintiff, a breach of that duty, and injury proximately resulting from the breach.” *O'Connor*, 477 F. Supp. 3d at 721 (cleaned up). Plaintiffs allege that Defendants had “a duty to exercise reasonable care in the manufacturing, designing, researching, testing, producing, supplying, inspecting, marketing, labeling, packaging, selling, and distributing of their hair relaxer products.” [106 ¶ 116]. According to Plaintiffs, Defendants had a duty to exercise reasonable care in the advertising and sale of their hair relaxer products, including a duty to warn of risks associated with the products, and also owed a continuing duty to Plaintiffs to remove, recall, or retrofit the unsafe and/or defective hair relaxer products. *Id.* ¶¶ 117–18. Plaintiffs claim Defendants breached these duties in several ways including by distributing their hair relaxer products negligently and recklessly and by failing to adequately warn of the products' risks and dangers. *Id.* ¶ 120. Finally, they allege that Defendants' negligence “was a direct and proximate cause of the injuries, harm, and economic losses that Plaintiffs have suffered, and will continue to suffer.” *Id.* ¶ 125. The allegations read as a whole give rise to the inference that Defendants'

conduct proximately caused Plaintiffs' injuries; more is not' needed at this stage. *See Swearingen v. Momentive Specialty Chemicals, Inc.*, 662 F.3d 969, 972 (7th Cir. 2011) ("breach of duty and proximate cause present questions of fact"). In short Plaintiffs state a negligence claim. *See* Fed. R. Civ. P. 8(a)(2) (requiring "a short and plain statement of the claim showing that the pleader is entitled to relief"); *Engel v. Buchan*, 710 F.3d 698, 709 (7th Cir. 2013) (reading complaint as a whole on a motion to dismiss).

Defendants also argue that Plaintiffs did not plead sufficient knowledge. *See e.g. Happel v. Wal-Mart Stores, Inc.*, 766 N.E.2d 1118, 1123 (Ill. 2002) (cleaned up). But Plaintiffs specifically alleged that "Defendants knew or should have known that phthalates and other EDCs in their hair relaxer products significantly increase the risk of cancers and other negative health conditions." [106 ¶ 118]. Plaintiffs claimed they did not know about the products' risks, and Defendants knew or should have known ordinary consumers like plaintiffs would not realize the dangers. *See id.* ¶¶ 119, 153, 185. Defendants criticize Plaintiffs' reliance on the 2022 Chang Article and argue they have not "establish[ed]" Defendants' breach. But these are not issues capable of being resolved on a motion to dismiss.⁵ Accepting Plaintiffs' factual allegations as true and reading the complaint as a whole shows Plaintiffs have plead sufficient facts to avoid dismissal. Defendants' motion to dismiss Counts 1 and 3 is denied.

⁵ Defendants rely on cases that are not dispositive at this point, *Cornstubble v. Ford Motor Co.*, 532 N.E.2d 884 (Ill. App. Ct. 1988) (decided following a bench trial), and *O'Connor*, 477 F. Supp. 3d at 721 (decision turned on the economic loss doctrine).

2. Design Defect

In Count 4, Plaintiffs allege that Defendants' hair relaxer products "were defectively designed because they caused serious injuries and death, including but not limited to uterine cancer and ovarian cancer." [106 ¶ 152]. A strict-liability design defect claim requires: "(1) a condition of the product as a result of design, (2) that made the product unreasonably dangerous, (3) and that existed at the time the product left the defendant's control, and (4) an injury to the plaintiff (5) that was proximately caused by the condition." *Clark v. River Metals Recycling, LLC*, 929 F.3d 434, 439 (7th Cir. 2019) (cleaned up).

Defendants contend that Plaintiffs' allegations are conclusory and they have not identified specific products or defects in those products. The Court does not agree. Plaintiffs allege that Defendants' products contained toxic chemicals. [106 ¶ 55]. Specifically, they allege harmful and carcinogenic ingredients in Defendants' hair relaxer products "are known to disrupt and/or harm a woman's endocrine system." *Id.* ¶ 71. "Such harmful, toxic and carcinogenic ingredients have included over time, but are not limited to, phthalates, parabens, cyclosiloxanes, di-(2-ethylhexyl), octamethylcyclotetrasiloxane, lye, formaldehyde, and other toxic chemicals." *Id.* The phthalates and other EDCs in Defendants' hair relaxer products, Plaintiffs claim, "significantly increase the risk of cancers and other negative health conditions." *Id.* ¶ 118.

Defendants rely on *Griffin v. Medtronic, Inc.*, No. 17 CV 927, 2017 WL 4417821 (N.D. Ill. Oct. 5, 2017), where the complaint was "silent as to what was wrong with

the device”, and plaintiff forfeited any responsive arguments by failing to respond to the motion to dismiss. Here, Plaintiffs have not failed to respond, and the MLC is not silent about the alleged defect. Plaintiffs claim that “[n]atural and synthetic EDCs are present in some of Defendants’ hair relaxer products under the guise of ‘fragrance’ and ‘perfumes’, and thus enter the body when these products are applied to the hair and scalp.” [106 ¶ 76]. They allege that the 2022 NIH study “revealed that there were significantly higher rates of uterine cancer in women who had used hair relaxers,” and the 2021 study found “frequent use of hair relaxers [] strongly associated with ovarian cancer.” *Id.* ¶ 86. In their MLC Plaintiffs rely on a number of scientific studies and secondary sources. At a later stage of the litigation, Defendants will be able to challenge the studies. For now, Defendants’ critique is premature. Plaintiffs have sufficiently stated a design defect claim.

3. Failure to Warn

Plaintiffs claim that Defendants failed to give an appropriate and adequate warning of the risks of uterine cancer, ovarian cancer, and endometrial cancer. [106 ¶ 166]. A duty to warn under Illinois law exists “when there is unequal knowledge and the defendant, possessed of such knowledge, knows or should know that harm might occur if no warning is given.” *Proctor v. Davis*, 682 N.E.2d 1203, 1211 (1st Dist. 1997) (cleaned up). Plaintiffs allege that Defendants did not satisfy their duty to warn of the products’ “dangerous adverse side effects,” “potential or known toxic chemicals and carcinogens,” or that Defendants “had not properly tested the safety of their hair relaxer products.” [106 ¶¶ 58, 110, 133]. Plaintiffs specifically describe EDCs and

identify the EDCs alleged to be in the products. *Id.* ¶¶ 71–81. They claim they would not have purchased or used Defendants’ hair relaxer products had they known the true facts about the products. *Id.* at ¶ 138.

To argue Plaintiffs’ allegations are not adequate, Defendants contend that Plaintiffs do not identify specific products or timeframes. The Court has already addressed that argument. And again, the case law Defendants cite does not require dismissal. In *N. Tr. Co. v. Upjohn Co.*, 572 N.E.2d 1030 (Ill. App. Ct. 1991), the court explained that plaintiff was required to *show at trial* “that the omission of such information made the warning inadequate and the drug ‘defective’ and that this defect was the proximate cause of plaintiff’s injuries.” *Id.* at 1037. Contrary to Defendants’ characterization, the complaint does not allege or raise the inference that the first time Defendants were aware of the possible association between hair straightening products and cancer was in 2021 and 2022 when the Chang and White Articles were published. Instead, the complaint describes the long history of Black and Brown women being compelled to conform to Eurocentric beauty standards of straight hair, leading up to the manufacture of the first lye relaxer in 1971. [106 ¶¶ 38–47, 53–55]. Plaintiffs explain that particular defendants began marketing their first hair relaxer products in the 1970s and 1980s, and by the 1990s were making representations and omissions about chemicals in their products in advertisements and packaging. *Id.* ¶¶ 57–59. The complaint states that “[f]or decades and to present....[Defendants] marketed their hair relaxer products without ever disclosing known health risks of the toxic chemicals contained in these products.” *Id.* ¶ 55.

Plaintiffs allege that Defendants “were aware or should have been aware of both the potential for harm and the increased risk of developing uterine and ovarian cancer from the use of the hair relaxer products based on the evolving scientific studies, ongoing research, and various government standards and regulations.” *Id.* ¶ 70.

Taking the factual allegations as true and drawing reasonable inferences in Plaintiffs’ favor at this stage, the Court finds that Plaintiffs have stated a failure to warn claim.

C. Fraud-Based Claims

Defendants challenge Plaintiffs’ fraud-based claims (Counts 2, 8, 9, and 10) under Rule 9(b)’s heightened pleading standard. That rule requires a plaintiff alleging fraud to “state with particularity the circumstances constituting fraud.” Fed. R. Civ. P. 9(b). Further, “[a] claim that ‘sounds in fraud’—in other words, one that is premised upon a course of fraudulent conduct—can implicate Rule 9(b)’s heightened pleading requirements.” *Borsellino v. Goldman Sachs Grp., Inc.*, 477 F.3d 502, 507 (7th Cir. 2007). Rule 9(b) requires a plaintiff to provide “precision and some measure of substantiation to each fraud allegation.” *Menzies v. Seyfarth Shaw LLP*, 943 F.3d 328, 338 (7th Cir. 2019). Plaintiffs do not dispute that Rule 9(b) applies (though they point out that some jurisdictions consider consumer protection claims under Rule 8).

Plaintiffs ask the Court to focus on Defendants’ alleged omissions, and not reach the issue of affirmative misrepresentations, which Plaintiffs say can be handled in the individual cases. [176 at 31]. But Plaintiff’s fraud-based claims are largely based on alleged misrepresentations. *See* [106 ¶¶ 131–32, 137, 218]. They do allege that

Defendants concealed material facts about the dangers of their products, but do not identify specific omissions of material fact or support these allegations with additional detail. *See Squires-Cannon v. Forest Pres. Dist. of Cook Cnty.*, 897 F.3d 797, 805 (7th Cir. 2018) (allegations must be made with particularity). “To constitute fraud, an omission must be of a material fact that a consumer would rely on in making her decision, or that would have caused her to act differently had she known of it.” *Siegal v. GEICO Cas. Co.*, 523 F. Supp. 3d 1032, 1042 (N.D. Ill. 2021). Plaintiffs are correct that courts consider omission-based claims under a more relaxed standard. *See Fed. Deposit Ins. Corp. v. Patel*, No. 19-cv-6917, 2020 WL 6681348, at *2 (N.D. Ill. Nov. 12, 2020) But Rule 9(b) still applies. *See Pirelli Armstrong Tire Corp. Retiree Med. Benefits Tr. v. Walgreen Co.*, 631 F.3d 436, 446–47 (7th Cir. 2011). “[F]raudulent omission claims are subject to the heightened pleading standard under Rule 9(b), requiring them to be plead with particularity.” *Rodriguez v. Ford Motor Co.*, 596 F. Supp. 3d 1050, 1058 (N.D. Ill. 2022).

In addition, as Defendants point out, a fraudulent omission claim requires that Defendants “intentionally omitted or concealed a material fact that [they were] under a duty to disclose” to plaintiffs. *Wigod v. Wells Fargo Bank, N.A.*, 673 F.3d 547, 571 (7th Cir. 2012). *See also Rodriguez*, 596 F. Supp. 3d at 1057–58. A fraud claim based on an alleged omission of material fact requires the parties to have a special or fiduciary relationship. *See Cohen v. Am. Sec. Ins. Co.*, 735 F.3d 601, 614 (7th Cir. 2013); *Walker v. Bank of Am.*, No. 21-CV-03589, 2022 WL 910585, at *5 (N.D. Ill. Mar. 29, 2022). The allegations in the complaint do not show the concealment was

“more than a mere passive omission of facts during a business transaction” that was “done with the intention to deceive under circumstances creating an opportunity and duty to speak.” *Rodriguez*, 596 F. Supp. 3d at 1057–58.

For these reasons the Court finds Plaintiffs have not met the heightened pleading standard as to Counts 2, 8, and 9. As to Count 10, Plaintiffs argue their unfairness claims should at least survive. The Court finds that Plaintiff’s claims of deceptive acts or practices under the consumer fraud statutes do not meet the heightened pleading standard of Rule 9(b). *See Vanzant v. Hill's Pet Nutrition, Inc.*, 934 F.3d 730, 738 (7th Cir. 2019). However, Plaintiffs’ claims of unfair conduct survive. “The pleading standards for deceptive and unfair claims differ...[b]ecause fraud is not an element of an unfair conduct claim, Rule 9(b)’s heightened standard does not apply, but rather the notice pleading standard under Rule 8(a) does.” *Siegel*, 523 F. Supp. 3d at 1041. Plaintiffs allege that Defendants misled consumers about the safety risks associated with use of their hair relaxer products by among other things misrepresenting that the goods were of a particular standard, quality, or grade. [160 ¶¶ 241, 243]. Reading the complaint as a whole and accepting Plaintiffs’ factual allegations as true, the Court finds Plaintiffs’ claims of unfair conduct survive under Rule 8.

In sum, the Court grants Defendants’ motion to dismiss Counts 2, 8, and 9, and grants Defendants’ motion as to Count 10 but only as to fraud-based claims; Plaintiffs’ claims of unfair conduct survive.

D. Warranty Claims

In Counts 6 and 7, Plaintiffs claim breach of implied warranty and breach of express warranty under state law and the Magnuson-Moss Warranty Act, 15 U.S.C. § 2301 (“MMWA”). An express warranty claim requires that a seller: “(1) made an affirmation of fact or promise; (2) relating to the goods; (3) which was part of the basis for the bargain; and (4) guaranteed that the goods would conform to the affirmation or promise.” *O’Connor*, 477 F. Supp. 3d at 714. To state a claim for breach of express warranty, a plaintiff “must show that [defendant] breached an affirmation of fact or promise that was made a part of the basis of the bargain.” *Bakopoulos v. Mars Petcare US, Inc.*, 592 F. Supp. 3d 759, 756 (N.D. Ill. 2022) (cleaned up). Illinois law, for example, requires that express warranties be created through an “affirmation of fact or promise made by the seller to the buyer.” *See* 810 ILCS 5/2-313(1)(a). To state a claim for breach of the implied warranty of merchantability, “a plaintiff must allege that (1) the defendant sold goods that were not merchantable at the time of sale; (2) the plaintiff suffered damages as a result of the defective goods; and (3) the plaintiff gave the defendant notice of the defect.” *Corwin v. Connecticut Valley Arms, Inc.*, 74 F. Supp. 3d 883, 891 (N.D. Ill. 2014) (cleaned up).

Defendants first argue that Plaintiffs fail to allege the requisite privity for these claims. As Plaintiffs point out, some states have exceptions to the privity requirement. This Court agrees with the approach in other MDL cases in this district declining to rule on these state-specific issues at this stage:

Resolution of those latter issues would require a detailed analysis of the allegations in each individual complaint and whether they are sufficient to state a claim under the law of each plaintiffs’ home state. This

category of challenges is not well-suited to resolution in an omnibus fashion, and the Court declines to resolve them at this stage of the case.

In re Recalled Abbott Infant Formula Prod. Liab. Litig., No. 22 C 4148, 2023 WL 3585639, at *3 (N.D. Ill. May 22, 2023); *see also In re Testosterone Replacement Therapy Prod. Liab. Litig.*, No. 14 C 1748, 2014 WL 7365872 (N.D. Ill. Dec. 23, 2014). This Court therefore declines to rule at this point on specific states' privity requirements for plaintiffs' warranty claims.

Here the Complaint adequately pleads Plaintiffs' warranty claims under Rule 8. Plaintiffs allege that Defendants made affirmations of fact "that their hair relaxer products are safe, healthy, protective, and/or natural." [106] ¶ 197.⁶ Plaintiffs allege Defendants made a number of misrepresentations, including that their products are "gentle," "natural," "healthy," do not contain toxic or "harsh chemicals," "protect the skin and scalp," do not "hurt[] your scalp," and can be used "safely" by women and children. *Id.* at ¶¶ 58–59. Plaintiffs further allege that they would not have purchased Defendants' products had they "known the truth about the misrepresentations." *Id.* at ¶ 204. And Plaintiffs also claim that Defendants' products did not perform as intended: harmful EDCs "enter the body when these products are applied to the hair and scalp," "[c]hronic exposure to phthalates will adversely influence the endocrine system and functioning of multiple organs," and use of these hair relaxer products resulted in serious injuries or death to plaintiffs from certain cancers. *See id.* at ¶¶ 74–79, 112, 120, 152, 153, 177–79.

⁶ The Court disagrees with Defendants' characterization that Plaintiffs concede there were no affirmative misrepresentations by Defendants.

As for Defendants' argument that Plaintiffs cannot demonstrate that Defendants' products were not "fit for a particular purpose," Plaintiffs respond that this argument would only bar this theory for plaintiffs whose claims are governed by Illinois law, and other states' laws differ. This Court will again not parse state law variations at this stage of the case. *See In re Recalled Abbott Infant Formula Prod. Liab. Litig.*, 2023 WL 3585639.

Finally, Plaintiffs bring a claim under the MMWA, the federal warranty statute. *See* 15 U.S.C. § 2301. *See Schimmer v. Jaguar Cars, Inc.*, 384 F.3d 402, 405 (7th Cir. 2004) ("[T]he MMWA allows consumers to enforce [limited] written and implied warranties in federal court [...] borrowing state law causes of action."). Because Plaintiffs' express and implied warranty claims survive, so does their MMWA claim. In short, Plaintiffs' warranty claims need only meet Rule 8's standards, and they do.

E. Unjust Enrichment, Punitive Damages, and Derivative Claims

In their unjust enrichment claim, Plaintiffs allege that Defendants have profited and benefited from payments Plaintiffs and other consumers made for their hair relaxer products. [106 ¶ 252]. Under Illinois law an unjust enrichment claim requires allegations that the defendant "unjustly retained a benefit to the plaintiff's detriment, and that the retention of the benefit violates fundamental principles of justice, equity, and good conscience." *Cleary v. Philip Morris Inc.*, 656 F.3d 511, 516 (7th Cir. 2011). The parties acknowledge that different jurisdictions treat unjust enrichment claims differently. However the Court does not find it appropriate at this

stage to rule on a case wide basis in an MDL that no plaintiff may pursue an unjust enrichment claim.

Plaintiffs also seek punitive damages, alleging: “[t]he acts and omissions of Defendants as alleged throughout this Complaint were willful, wanton, and malicious. Defendants committed these acts with a conscious disregard for the rights, health, and safety of Plaintiffs and other consumers/users of Defendants’ hair relaxer products, for the primary purpose of increasing Defendants’ profits from the sale and distribution of their hair relaxer products.” [106] ¶ 279. Defendants contend that punitive damages are a remedy, not an independent cause of action, and Plaintiffs have not alleged facts sufficient to seek punitive damages. The Court, however, finds the factual allegations supporting the claim for punitive damages adequate. And the Court does not agree that dismissing Plaintiffs’ punitive damages claim, whether viewed as a cause of action or remedy, is warranted now. *See Schramm v. Peregrine Transportation Co., LLC*, No. 3:22-CV-161-NJR, 2023 WL 2349346, at *4 (S.D. Ill. Mar. 3, 2023); *Eliason v. Superior Ref. Co. LLC*, No. 19-CV-829-WMC, 2021 WL 1227607, at *4 (W.D. Wis. Mar. 31, 2021).

Finally, Defendants’ only argument for dismissing the derivate claims of wrongful death, survival action, and loss of consortium is that Plaintiffs’ other claims are not viable. As discussed, however, most of Plaintiffs’ claims have survived dismissal. For these reasons, Plaintiffs’ Counts 11, 12, 13, 14, and 15 survive Defendants’ motions.⁷

⁷ McBride’s motion to dismiss incorporates arguments raised in Defendants’ Omnibus Motion to Dismiss. [192] McBride’s motion largely overlaps with the omnibus motion and raises the same challenges. The Court’s analysis herein applies equally to McBride’s motion, which the Court also grants in part and denies in part.

F. Personal Jurisdiction Challenge

Defendants argue that Dabur International and Dermoviva must be dismissed because Plaintiffs cannot establish that the Court has general or specific personal jurisdiction over those defendants. Plaintiffs respond that Defendants' motion should be denied without prejudice as premature. Plaintiffs further contend that in any event, Dabur would be subject to general jurisdiction here.

1. Standard

Under Rule 12(b)(2), a court may dismiss a claim for lack of personal jurisdiction over the defendant. Fed. R. Civ. P. 12(b)(2). A plaintiff need not allege facts about personal jurisdiction in his or her complaint, but in the face of a Rule 12(b)(2) motion, “the plaintiff bears the burden of demonstrating the existence of jurisdiction.” *Curry v. Revolution Labs., LLC*, 949 F.3d 385, 392 (7th Cir. 2020) (quoting *Purdue Research Found. v. Sanofi-Synthelabo, S.A.*, 338 F.3d 773, 782 (7th Cir. 2003)). When a court rules on a Rule 12(b)(2) motion only based on written submissions, a plaintiff need only establish a prima facie case of personal jurisdiction. *GCIU-Employer Ret. Fund v. Goldfarb Corp.*, 565 F.3d 1018, 1023 (7th Cir. 2009). Where, as here, a defendant submits an affidavit regarding personal jurisdiction, this Court accepts as true any facts in the affidavit that do not conflict with the complaint or the plaintiff's submissions. *Curry*, 949 F.3d at 393. Further, where a defendant challenges by declaration a fact alleged in the plaintiff's complaint, the plaintiff must go beyond the pleadings and submit affirmative evidence supporting the exercise of jurisdiction. *Purdue Research Foundation*, 338 F.3d at 783. If the plaintiff “fails to refute a fact

contained in the defendant's affidavit, that fact is accepted as true." *Mold-A-Rama Inc. v. Collector-Concierge-Int'l*, No. 18-CV-08261, 2020 WL 1530749, at *2 (N.D. Ill. Mar. 31, 2020). A transferee court in an MDL can "exercise personal jurisdiction to the same extent that the transferor court could." *In re Delta Dental Antitrust Litig.*, 509 F. Supp. 3d 1377, 1379 (U.S. Jud. Pan. Mult. Lit. 2020).

2. Analysis

First, the Court grants Dermoviva's motion and dismisses Dermoviva from this MDL. Plaintiffs assert that Dermoviva's motion is premature but state that the three individual plaintiffs who have named this defendant are voluntarily dismissing those claims so that motion is moot. Because Plaintiffs do not put forth any argument to establish this Court's personal jurisdiction over Dermoviva, Dermoviva is dismissed. *See M.G. Skinner & Assocs. Ins. Agency*, 845 F.3d at 321; *Curry*, 949 F.3d at 392.

As for Dabur, Defendants assert, and provide a sworn affidavit of Dabur's CEO, that Dabur is incorporated in the Isle of Man and principally located in Dubai, UAE. [142 at 41; 142-1 (Agrawal Affidavit)]. It has a sole United States office in New Jersey where it conducts limited business related to its own products, none of which are hair relaxers. *Id.* According to Dabur's CEO, the company has no involvement with any hair-relaxer products sold anywhere in the United States. *Id.* Plaintiffs respond that this motion is premature because personal jurisdiction questions cannot be decided on an MDL-wide basis.

Defendants' filing and affidavit show that this Court does not have personal jurisdiction over Dabur, and Plaintiffs have not rebutted those assertions. Dabur is

not “at home” in Illinois, for purposes of general jurisdiction, and for purposes of specific jurisdiction, Dabur did not manufacture, market, distribute, sell, or make representations about any hair relaxer products underlying Plaintiffs’ claims anywhere in the United States, let alone in Illinois. Agrawal Affidavit ¶ 4. Plaintiffs’ argument that Dabur’s motion is premature and cannot be decided on an MDL-wide basis is not supported by authority. As to general jurisdiction, Plaintiffs argue that Dabur’s registration in Illinois from 2012 to 2021 is sufficient to establish personal jurisdiction.

Mallory v. Norfolk S. Ry. Co., 600 U.S. 122 (2023), cited by Plaintiffs, addressed *Pennsylvania’s* requirement that an out-of-state corporation *consent to* personal jurisdiction to do business there. The Illinois long-arm statute’s “doing business” standard for example is “virtually identical to the federal requirement for general jurisdiction that a party maintain continuous and systematic business contacts with the forum.” *Landwer v. Sodhi*, 2018 WL 6000868, at *3 (N.D. Ill. Nov. 15, 2018). Plaintiffs do not argue that any particular state requires a company to consent to suit in the forum; if that is Plaintiffs’ suggestion, they have not developed the argument or offered an analysis of particular state statutes to permit the Court to find that personal jurisdiction is proper. Given Plaintiffs’ burden in the face of Defendants’ Rule 12(b)(2) motion and attached affidavit, the Court finds personal jurisdiction over Dabur lacking.

Therefore the Court grants Defendants' Rule 12(b)(2) motion as to Defendants Dermoviva and Darbur. Dermoviva Skin Essentials, Inc., Darbur International Limited and "Darbur International USA Ltd." are dismissed.

IV. Conclusion

For the stated reasons, Defendants' Joint Motion to Dismiss [142] and Defendant McBride's motion to dismiss [192] are granted in part and denied in part. Defendants Keratin Complex, Keratin Holdings, LLC's unopposed motion for joinder [263] is granted. Dubar International Limited and Dermovia Skin Essentials, Inc. are dismissed as defendants from this matter. Counts 2, 8 and 9 are dismissed. Count 10 is dismissed as to the fraud-based claims only. The motions to dismiss the remaining claims are denied.

E N T E R:

Dated: November 13, 2023



MARY M. ROWLAND
United States District Judge