IN THE STATE COURT OF DEKALB COUNTY STATE OF GEORGIA

LAFAYETTE ARMSTRONG, Individually and as the Personal Representative of the Estate of Barbara Armstrong

Plaintiff,

v.

MCBRIDE RESEARCH LABORATORIES, INC., L'ORÉAL USA, INC., L'ORÉAL USA PRODUCTS, INC., SOFTSHEEN-CARSON LLC., SOFTSHEEN-CARSON (W.I.), INC.,

Defendants.

	23A00243
Civil Action	

JURY TRIAL DEMAND

COMPLAINT

The Plaintiff, Lafayette Armstrong, individually and on behalf of the Estate of Barbara Armstrong, through his undersigned counsel, makes the following Complaint against Defendants McBride Research Laboratories, L'Oréal USA, Inc., USA L'Oréal Products, Inc., SoftSheen-Carson LLC., and SoftSheen-Carson (W.I.), Inc. ("Soft Sheen") (collectively, "Defendants"), and alleges the following:

NATURE OF THE ACTION

1. This action arises out of Mrs. Armstrong's gynecologic injuries caused by Defendants. Specifically, Mrs. Armstrong's uterine cancer and endometriosis were directly and proximately caused by her regular and prolonged exposure to phthalates and other endocrine disrupting chemicals found in Defendants' hair care Products.

2. Plaintiff brings this action against Defendants for claims arising from the direct and proximate result of Defendants', their directors', agents', heirs' and assigns', and/or their corporate predecessors' negligent, willful, and wrongful conduct in connection with the design, development, manufacture, testing, packaging, promoting, marketing, distribution, labeling, regulation, and/or sale of the Products, including those known as Design Essentials and Dark & Lovely.

I. PARTIES

- 3. Plaintiff Lafayette Armstrong, individually and on behalf of the Estate of Barbara Armstrong, is, and at all times relevant to this action was, a citizen and resident of the state of South Carolina, with their place of residence being Sumter County, South Carolina.
- 4. Plaintiff Lafayette Armstrong is the husband and duly appointed personal representative of the Estate of Barbara Armstrong.
- 5. Defendants McBride Research Laboratories, Inc. is, and at all times relevant to this action was, a corporation with its principal place of business at 2272 Park Central Blvd, Decatur, GA 30035. Process may be served upon its registered agent James Hunter at 3343 Peachtree Road #1600, Atlanta, GA, 30326.
- 6. Defendant L'Oréal USA, Inc. is, and at all times relevant to this action was, a corporation with its principal place of business and headquarters located at 575 Fifth Avenue, New York, New York 10017. Process may be served upon its registered agent, Corporation Service Company, 80 State Street, Albany, NY 12207.
- 7. Defendant L'Oréal USA Products, Inc. is, and at all times relevant to this action was, a corporation with its principal place of business and headquarters located at 10 Hudson

Yards, 347 10th Avenue, New York, New York 10001. Process may be served upon its registered agent, Corporation Service Company, 80 State Street, Albany, NY 12207.

- 8. Defendant SoftSheen-Carson LLC., is, and at times relevant to this action was, a corporation with its principal place of business and headquarters located in Savannah, Georgia. Upon information and belief, Defendant SoftSheen-Carson LLC is presently headquartered at 10 Hudson Yards, 347 10th Avenue, New York, New York 10001. Process may be served upon its registered agent, Corporation Service Company, 80 State Street, Albany, NY 12207.
- 9. Defendant SoftSheen-Carson LLC., is and at all relevant times has been, a wholly owned subsidiary of Defendant L'Oréal USA, under the complete dominion and control of Defendant L'Oréal.
- 10. Defendant SoftSheen-Carson (W.I.), Inc., is, and at all times relevant to this action was, a corporation. Process may be served upon its registered agent, Corporate Service Company 251 Little Falls Drive, Wilmington, Delaware 19808.
- 11. Defendant SoftSheen-Carson (W.I.), Inc., is, and at all relevant times has been, a wholly owned subsidiary of Defendant L'Oréal USA, under the complete dominion of and control of Defendant L'Oréal.
- 12. Defendants L'Oréal USA, Inc., L'Oréal USA Products, Inc., SoftSheen-Carson LLC. and SoftSheen-Carson (W.I.), Inc., are collectively referred to herein as the "L'Oréal Defendants."
- 13. Upon information and belief, at all pertinent times, all Defendants regularly transacted, solicited, and conducted business in all fifty states of the United States, with full knowledge that their Hair Straighteners and Relaxers were being placed into interstate commerce and being sold in Georgia.

14. At all times material hereto, Defendants developed, tested, assembled, manufactured, packaged, labeled, prepared, distributed, marketed, supplied, and/or sold hair straightener and relaxer products, hereinafter referred to as the "Products," including but not limited to:

- a. Design Essentials;
- b. Dark & Lovely;
- c. Optimum;
- d. Care Free;
- e. Frizz Free;
- f. Alternatives; and
- g. Soft & Beautiful.
- 15. Defendants' defective hair Products were placed into the stream of interstate commerce and were used by the Plaintiff in or around 1970 through approximately 2018.
- 16. Plaintiff Barbara Armstrong used the Products continually throughout her life every six to eight weeks both in beauty salons and personally.
- 17. Plaintiff Barbara Armstrong also used hair relaxer and straightener Products during the period described in paragraph 15 manufactured by Revlon Inc. and/or Revlon Consumer Products Corporation ("Revlon"), both Delaware corporations headquartered in New York. Revlon and Revlon products used by Plaintiff are intended to be included in the descriptive terms "Defendants" and "Products" herein, although Revlon is not a party to this lawsuit at this time because of ongoing bankruptcy proceedings.
- 18. On or about May 28, 2019, Plaintiff was diagnosed with uterine cancer, a diagnosis caused by Plaintiff's exposure to chemicals in the Defendants' aforementioned hair relaxer

Products.

19. On January 28, 2020, Plaintiff passed away from complications related to her uterine cancer and chemotherapy, caused by the use of Defendants' hair relaxer Products.

II. JURISDICTION AND VENUE

- 20. This Court has personal jurisdiction over McBride Research Laboratories. McBride's principal place of business is located in DeKalb County, Georgia. McBride regularly and systematically transacts business within the State of Georgia, has a registered agent within the State of Georgia, and has committed tortious acts both inside and outside the State of Georgia while regularly doing business in Georgia and engaging in a persistent course of conduct, and/or deriving substantial revenue from goods used and/or consumed and/or services rendered in Georgia. These acts included manufacturing, supplying, packaging, formulating, distributing, shipping, testing, and otherwise selling the Products used by Plaintiff.
- 21. This Court has personal jurisdiction over the L'Oréal Defendants pursuant to O.C.G.A. § 9-10-91. The L'Oréal Defendants regularly and systematically transact business within the State of Georgia, had a subsidiary with a registered agent in the State of Georgia, and have committed tortious acts and omissions within the State of Georgia. Upon information and belief, for many years the SoftSheen-Carson subsidiary entity had a principal place of business in Georgia and upon information and belief they still have a large manufacturing presence and headquarters in Atlanta, Georgia. The L'Oréal Defendants committed tortious acts both inside and outside the State of Georgia while regularly doing business in Georgia and engaging in a persistent course of conduct, and/or deriving substantial revenue from goods used and/or consumed and/or services rendered in Georgia. These acts included manufacturing, formulating, packaging, supplying, distributing, shipping, testing, and otherwise selling the Products used by Plaintiff.

22. Venue is proper in this Court pursuant to O.C.G.A. § 9-10-93 as Defendant McBride Research Laboratories, Inc.'s principal place of business is in DeKalb County, Georgia and Plaintiff has used Products that were created, tested, processed, formulated, packaged, shipped, distributed, and otherwise manufactured in DeKalb County, Georgia, and because a substantial part of the actions or omissions giving rise to Plaintiffs' claims occurred in this judicial district.

III. <u>BACKGROUND</u>

- 23. This is a wrongful death and survival action arising out of the death of Barbara Armstrong.
 - 24. At the time of her death, Barbara Armstrong was 69 (sixty-nine) years old.
- 25. At the time of her death, Barbara Armstrong was happily married to her husband Lafayette Armstrong.

IV. FACTUAL ALLEGATIONS

A. Hair Straighteners and Relaxers

- a. Market for Hair Straightening and Relaxing Products
- 26. Black people make up about 13 percent of the U.S. population, but by one estimate, African American spending accounts for as much as 22 percent of the \$42 billion-a-year personal care products market, suggesting that they buy and use more of such products including those with potentially harmful ingredients than Americans as a whole.

¹ Thandisizwe Chimurenga, *How Toxic is Black Hair Care?*, New America Media, Feb. 2, 2012, www.americamedia.org/2012/02/skin-deep-in-more-ways-than-one.php; *Personal Care Products Manufacturing Industry Profile*, Dun & Bradstreet First Research, August 2016, www.firetresearch.com/Industry-Research/Personal-Care-Products-Manufacturing.html (this report uses "Black" to describe not only people who identify as African-American, but Black people in the U.S. who come from the Caribbean or other areas. "African-American" is used only when a cited source specifies that term).

- 27. In an analysis of ingredients in 1,177 beauty and personal care products marketed to Black women, about one in twelve was ranked highly hazardous on the scoring system of EWG's Skin Deep® Cosmetics Database, a free online resource for finding less-hazardous alternatives to personal care products. Maintained by the Environmental Working Group, this database provides information on the safety and potential harms of ingredients in a wide range of skincare products, cosmetics, and other personal care and beauty products. The database provides information on cancer risk, reproductive toxicities, and allergies.² The worst scoring products marketed for Black women were hair relaxers, hair colors and bleaching products. Each of these categories had an average product score indicating a high potential hazard.
- 28. In the U.S. alone, Black consumers spend over \$1 trillion each year on consumer goods, with an outsized spending on hair care products compared to other ethnicities.
- 29. In 2020, the global Black hair care market was estimated at \$2.5 billion, with the hair relaxer market alone estimated at \$718 million in 2021, with the expectation of growth to reach or exceed \$854 million annually by 2028.

b. History of Hair Relaxers in America

- 30. In its natural or virgin state, afro-hair texture is characterized by coiled, springing, zigzag, and s-curve curl patterns; as well as by its density, fullness, texture, and feel. ³
 - 31. Afro-texture hair "naturally grows up and out."⁴
- 32. In Africa, hair was seen as a source of personal and spiritual power. As the highest point of the body and "most elevated part of the body, some communities believe(d) [their hair]

² <u>https://www.Ewg.org/skindeep/learn_more/about/</u>

³ Patrick Obukowcho, *Hair Relaxers: Science, Design, and Application*, 26, 14 (2018).

⁴ Ayana Byrd & Lori Tharps, *When Black Hair Is Against the Rules*, The New York Times, April 30, 2014,

connected them with the divine."⁵ For some, hair was the "conduit for spiritual interaction with God."⁶

- African hairstyles were status symbols reflecting one's "marital status, age, religion, and rank in society" and tribe. Warriors, kings, and queens were braids to show their ranking in society. The Wolof tribe in West Africa were braided styles when they went to war. 9
- 34. Most styling was extremely intricate and involved days of labor. "Only the mad and mourning did not do their hair." ¹⁰
- 35. Once African slaves arrived in America, their hair was cut off. This was a way to "break their spirit and make slaves easier to control." What was once a sign of pride and symbolism became a tool for subordination and degradation. As such, hair cutting was also an ordinary form of punishment.
- 36. The very nature of slavery involved working long hours in dire conditions. Enslaved people had "no time to care about one's appearance or one's hair." "Hair that was once a source of pride and expression of identity was often tucked away beneath cloth to cover rough,

⁵ Nicki Fox, *6 Things Everyone Should Know About Black Hair History*, Odele, Feb. 22, 2021. https://www.odelebeauty.com/blogs/the-rings/black-hair-history-facts

⁶ Rumeana Jahangir, *How Does Black Hair Reflect Black History?*, BBC News, May 31, 2015. https://www.bbc.com/news/uk-england-merseyside-31438273.amp

⁷ *History of Braids: More Than Just a Hairstyle*, Genesis Career College, https://www.genesiscareer.edu/history-of-braids-more-than-just-a-hairstyle/

⁸ *Id*.

⁹ *Id*.

¹⁰ Hlonipha, Mokoena, *From Slavery to Colonialism and School Rules, Navigating the History of Myths about Black Hair*, Quartz Africa, Fe., 24, 2018, https://qz.com/africa/1215070/black-hair-myths-from-slavery-to-colonialism-school-rules-and-good-hair/

¹¹ Brenda A. Randle, *I Am Not My Hair*, Race, Gender and Class, Volume 22, Number 1-2, 114-121 (2015).

¹² Nikki Fox, 6 Things Everyone Should Know About Black Hair History, Odele, Feb. 22, 2021.

tangled tresses and shield them from hours spent toiling under the sun."¹³ The hair that was once an important spiritual and cultural symbol became tangled and matted.

- 37. Historically, kinky and/or coarse textured hair did not meet the European standards of beauty. Instead, as noted in a New York Times article, British colonist described African hair as "closer to sheep wool than human hair." African hair that was once considered an attractive feature became a source of shame, to be covered or cut.
- 38. In 1786, Governor Don Esteban Miro of Louisiana passed the "Tignon Law" requiring Black women to wear a tignon (scarf) over their hair as a way of signifying they were members of the slave class, *even if they were free*. ¹⁵
- 39. "By requiring free Black women to wear the same hair covering, the governor was marking them as related to enslaved women rather than white women." ¹⁶
- 40. This law sent a direct signal to Black people that their hair held a symbol of inequality and was a sign of poverty regardless of their actual social status.
- 41. Because afro-textured hair was kinky and reflected African heritage rather than European ancestry, afro-textured hair was stigmatized as low social status. ¹⁷

¹³ *Id*.

Ayana Byrd & Lori Tharps, When Black Hair Is Against the Rules, The New York Times, April 30, 2014,

https://www.nytimes.com/2014/05/01/opinion/when-black-hair-is-against-the-rules.html

¹⁵ Nicki Fox, *6 Things Everyone Should Know About Black Hair History*, Odele, Feb. 22, 2021. https://www.odelebeauty.com/blogs/the-rings/black-hair-history-facts

¹⁶ Fashionable Rebellion, Women and the American Story, New York Historical Society Museum and Library, https://www.wams.nyhistory.org/settler-colonialism-and-revolution/settler-colonialism/fashionable-rebellion/

¹⁷ Brenda A. Randle, *I Am Not My Hair*, Race, Gender and Class, Volume 22, Number 1-2, 114-121 (2015).

- 42. Slaves with lighter skin and less coarse hair were favored to work in the home, a far less strenuous position than in the plantation fields. ¹⁸
- 43. Texturism, the idea that "good hair" is equated with a straighter hair texture, was cemented into American culture during its period of chattel slavery. "Eurocentric beauty standards dictated that coily hair and dark skin were unattractive and inferior." "(L)ighter skinned and straighter haired slaves were favored and selected for more desirable positions in the house" as opposed to the fields. ¹⁹ Thus, "the texture of an enslaved person's hair could determine their value and working conditions, which in turn might impact their overall health, comfort and chances for freedom[.] ²⁰ Naturally, Black women strived for a better life in America and were taught that the straighter and less kinky their hair was, the better life they could have. ²¹ This fueled the desire for tools and products that could straighten Black hair texture.
- 44. Gone were the days of African pride in African hairstyles. "The goal of grooming the hair had morphed from the elaborate and symbolic designs of Africa into an imitation of White styles adapted to Black kinks and curls."²²
- 45. In an effort to obtain a better life, many slaves would go to "dangerous lengths to straighten their hair."²³

¹⁸ *Id*.

¹⁹ *Id*.

²⁰ *Id*.

²¹ Chanel Donaldson, *Hair Alteration Practices Amongst Black Women and the Assumption of Self-Hatred*, Applied Psychology Opus, https://www.wp.nyu.edu/steinhardt-appsych_opus/hair-alteration-practices-amongst-black-women-and-the-assumption-of-self-hatred/

²² Brenda A. Randle, *I Am Not My Hair*, Race, Gender and Class, Volume 22, Number 1-2, 114-121 (2015).

Nicki Fox, 6 Things Everyone Should Know About Black Hair History, Odele, Feb. 22, 2021. https://www.odelebeauty.com/blogs/the-rings/black-hair-history-facts

- 46. Black or afro-textured hair can be manipulated into a straightened state with the use of hair tools and hair products. Prior to the invention of the chemical relaxer in the 1900s, individuals would "press" afro-textured hair with metal hair tools such as the "hot comb." Pressing combs or hot combs are metal hair tools that are first heated on a stove or in a ceramic heater, then pressed into hair strands to temporarily straighten them.²⁴
- 47. The hot comb was first invented by Frenchman Marcel Grateau who popularized the hair styling tool in Europe in the 1870s in part by advertising in catalogs of major department stores like Sears and Bloomingdales.²⁵ The hot comb was later modified by Madam C.J. Walker, a trailblazer in the development of Black hair products, to be manufactured with wider comb teeth.²⁶ With Walker's system, once the comb was heated, a softening ointment was then applied for easier manipulation of Black hair.²⁷
- 48. Today, afro-textured hair is still often straightened with alternative heat mechanisms rather than with chemicals. However, pressed hair remains susceptible to "shrinkage." Shrinkage is the process by which curly, kinky hair that has been temporarily straightened coils back into its natural state once the hair interacts with water, humidity, or perspiration, ²⁸ creating a shorter or fuller appearance.

²⁴ Jaclyn Peterson, *The Price of Beauty*, CTI Charlotte Teachers Institute Curriculum (2021).

Henry Louis Gates, *Madam Walker, the First Black American Woman to Be a Self-Made Millionaire*, PBS 100 Amazing Facts About the Negro, https://www.pbs.org/wnet/african-americans-many-rivers-to-cross/history/100-amazing-facts/madam-walker-the-first-black-american-woman-to-be-a-self-made-millionaire

²⁶ Cookie Lommel, Madam C.J. Walker 60 (1993)

²⁷ *Id.* at 62.

²⁸ *Id*.



- 49. African American inventor Garrett Augustus Morgan discovered and created a system that would permanently straighten afro-textured hair and eliminate the issue of "shrinkage."
- 50. In addition to being an inventor, Morgan was also a tailor. In the early 1900s, Morgan was repairing his sewing machines and wanted to find a way to polish the needles to stich fabrics more smoothly.²⁹ He applied a chemical solution to the needles and wiped the solution off with a rag. Morgan soon noticed that the "curly" fibers in the rag were straightened after exposure to the chemical. ³⁰
- 51. Morgan further tested the chemical on a dog with curly hair and eventually on his own hair. The chemical solution successfully straightened curly hair. He turned his formula into a gel-hair product, creating the G.A. Morgan Hair Refining Cream which was marketed in 1913.

²⁹ Patrick Obukowcho, *Hair Relaxers: Science, Design, and Application*, 27 (2018).

Mary N. Oluonye, Garrett Augustus Morgan: Businessman, Inventor, Good Citizen, 28 (2008).





52. Morgan's invention paved the way for the alkaline relaxer and later development of additional chemical-based permanent hair straightening products in the Black hair care market.³¹

³¹ Patrick Obukowcho, *Hair Relaxers: Science, Design, and Application*, 27 (2018).

c. Defendants' Marketing Efforts

- 53. In 1971, Carson, since acquired by L'Oreal and merged with SoftSheen, manufactured the first lye relaxer. The formula consisted of sodium hydroxide, water, petroleum jelly, mineral oils, and emulsifiers.³²
- 54. In the 1970s, lye relaxer users and manufacturers noticed that the lye formula stripped proteins from the hair strand, resulting in the hair thinning and breaking.³³ As a result, Johnson & Johnson marked the first "gentle" hair relaxer in 1981, which used milder chemicals such as potassium hydroxide and lithium hydroxide.³⁴
- 55. Over time, Soft & Beautiful and other chemical relaxer manufacturers developed herbal and botanical hair relaxer formulas. ³⁵
- 56. Today, Defendants market their hair relaxer products to Black customers across the United States, and the world, reinforcing the same historical Eurocentric standards of beauty. Defendants' marketing schemes rely heavily on branding and slogans that reinforce straight hair as the preferred standard. ³⁶
- 57. Defendant McBride Research Laboratories under its Design Essentials line depicts a woman going into a salon with curly hair and coming out with strait hair after using the product.

³² Cicely A. Richard, *This History of Hair Relaxers*, September 29, 2017, https://www.classrooms.synonym.com/the-history-of-hair -relaxers-12078983.html

³³ *Id*.

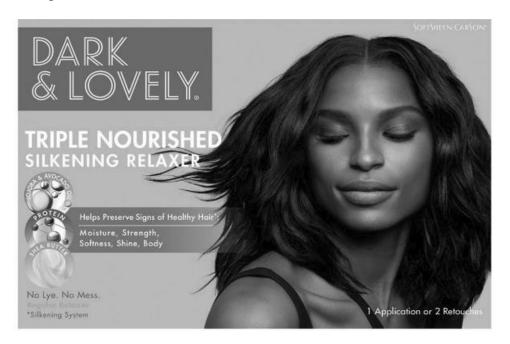
³⁴ *Id*.

³⁵ *Id*.

³⁶ *Id*.



58. Defendant L'Oreal depicts a Black woman with strait hair on its Dark & Lovely brand of relaxer product.



d. Chemical Relaxer Use

- 59. Hair relaxers are classified as creams or lotions which are specifically marketed to Black and Brown women to "tame their ethnic hair by making it smoother, straighter, and easier to manage on a daily basis."
- 60. Hair relaxing, or lanthionization, can be performed by a professional cosmetologist in a salon or barbershop, or at home with at-home relaxer kits designed for individual use. These home kits are sold in grocery, drug, beauty, and other retail supply stores in urban and rural cities throughout the United States.
- 61. Relaxers are applied to the base of the hair shaft and left in place for a "cooking" interval, during which the relaxer alters the hair's texture by purposefully damaging the hair's natural protein structure. The effect of this protein damage straightens and smooths the hair. After a period of weeks (4-8 weeks on average), depending on the hair's natural growth rate, the treated portion of the hair grows away from the scalp as new growth sprouts from the roots, requiring additional relaxer treatment to smooth the roots. These additional treatments are colloquially referred to in the community as "re-touches," resulting in women relaxing their new growth every four to eight weeks on average, typically for decades.
- 62. Hair relaxers can, and often do, cause burns and lesions on the scalp, facilitating introduction of hair relaxer chemicals into the body. The main ingredient of "lye" relaxers is sodium hydroxide; no-lye relaxers contain calcium hydroxide and guanidine carbonate, and "thio" relaxers contain thioglycolic acid salts. No-lye relaxers are advertised to cause fewer scalp lesions and burns than lye relaxers, but there is little evidence to support this claim.
- 63. Some studies show that up to 90% of Black and Brown women have used hair relaxants and straighteners. Hair relaxers contain hormonally active and carcinogenic compounds,

such as phthalates, known to cause endocrine disruption, which are not required to be listed separately as ingredients and are often broadly lumped into the "fragrance" or "perfume" categories of ingredients. Relaxer habits usually begin in the formative childhood years, and adolescence is a period of enhanced susceptibility to debilitating conditions resulting from exposure to these chemicals. ³⁷

- 64. In the 1990s, the first relaxer product for young Black girls, Just for Me TM, hit the market with a catchy advertising jingle that captured consumer attention. ³⁸ It soon became one of the most popular straightening treatments, touting a no-lye formula designed to be gentler for children's sensitive scalps.
- 65. Once relaxer use begins in childhood, it usually becomes a lifetime habit. The frequency of scalp burns with relaxer application can increase the risk of permanent and debilitating diseases associated with long-term exposure to endocrine-disrupting chemicals.
- 66. The reasons for Black women's use and dependence upon hair straightening products are associated with a range of factors, including (1) slavery and internalization of acceptable beauty norms, (2) media and advertisements, (3) assimilation and economic security, (4) ease of hair maintenance, and (5) culture. ³⁹
- 67. In a culture where Black women are often reduced to a lower standard of beauty, these factors impact women of color's decisions to begin and continue using products to alter the natural state of their hair, many times as a protective mechanism against racial discrimination. In

³⁷ Patrick Obukowcho, Hair Relaxers: Science, Design, and Application 27 (2018)

³⁸ Dana Oliver, *The 90s Just For Me Hair Relaxer Commercial Song Is Stuck In Our Heads*, HuffPost, Feb. 1, 2014. https://www.huffpost.com/entry/just-for-me-hair-relaxer-commercial-song n 4689981

³⁹ Chanel Donaldson, *Hair Alteration Practices Amongst Black Women and the Assumption of Self-Hatred*, Applied Psychology Opus, https://www.wp.nyu.edu/steinhardt-appsych_opus/hair-alteration-practices-amongst-black-women-and-the-assumption-of-self-hatred/

the Dove CROWN Study for girls (2021) conducted by JOY Collective, the following statistics were discovered⁴⁰:

- a. 100% of Black elementary school girls in majority-white schools who report experiencing hair discrimination state they experienced the discrimination by the age of (10)
- b. 86% of Black teens who experience discrimination state that they have experienced discrimination based on their hair by the age of twelve (12).
- c. 66% of Black girls in majority-white schools report experiencing hair discrimination compared to 45% of Black girls in all school environments.
- d. 53% of Black mothers, whose daughters have experienced hair discrimination, say their daughters experienced the discrimination as early as five (5) years old.
- e. 47% of Black mothers report having experienced discrimination related to their hair.
- f. Trauma from these experiences cause girls to miss days from school; teenage Black girls are missing a week of school per year due to hair dissatisfaction.
- g. While 90% of Black girls believe their hair is beautiful, the microaggressions and discrimination she endures has an impact on how she sees herself.
- h. Black women are 1.5 times more likely to be sent home from the workplace because of their hair.

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⁴⁰ The CROWN Act was created in 2019 by Dove and the CROWN Coalition, in partnership with then State Senator Holly J. Mitchell of California, to ensure protection against discrimination based on race-based hairstyles by extending statutory protection to hair texture and protective styles such as braids, locks, twists, and knots in the workplace and public schools. https://www.thecrownact.com

- i. Black women are 89% more likely than white women to agree with this statement, "I have to change my hair from its natural state to fit in at the office."
- 68. The CROWN Act of 2021 is a legislative bill introduced in both houses of Congress to address discrimination against protective hair styles worn predominantly by women of color. While the bill has not yet passed fully on a federal level, eighteen states have signed a version of the bill into state law. Unless and until the CROWN Act makes hair discrimination illegal in every state, teenagers and women of color continue to face discriminatory practices related to their hair choices, with relaxing and straightening their hair being a defensive, yet dangerous and toxic option.

e. Regulatory Framework

- 69. The law does not require cosmetic products and ingredients, other than color additives, to have FDA approval before they go to market. But there are laws and regulations that apply to cosmetics. The two most important laws pertaining to cosmetics marketed in the United States are the Federal Food Drug and Cosmetic Act (FD&C Act") and the Fair Packaging and Labeling Act ("FPLA").
- 70. The FD&C Act expressly prohibits the marketing of "adulterated" or "misbranded" cosmetics in interstate commerce.
- 71. Adulteration refers to a violation involving product composition whether it results from ingredients, contaminants, processing, packaging, shipping, or handling.
- 72. Under the FD&C Act a cosmetic is adulterated if: 1) it bears or contains any poisonous or deleterious substance causing injury to the product user, and 2) if its container is composed in whole or in part, of any poisonous or deleterious substance which may render the contents injurious to health.

- 73. Misbranding refers to violations involving improperly labeled or deceptively packaged products.
- Under the FD&C Act, a cosmetic is misbranded if 1) labeling is false or misleading;

 2) the label does not include all required information; 3) required information is not prominent and conspicuous; or 4) the packaging and labeling is in violation of an applicable regulation issued pursuant to Sections 3 and 4 of the Poison Prevention Packaging Act of 1970.⁴¹
- 75. Under U.S. law, cosmetic manufacturers are not required to submit their safety data to the FDA. However, it is against the law to put an ingredient in a cosmetic that makes the cosmetic harmful when used as intended. ⁴² An example is methylene chloride because it causes cancer in animals and is likely to be harmful to human health too. ⁴³
- 76. On May 19, 2022, the FDA issued a rule to amend its food additive regulations to no longer provide for most previously authorized phthalates to be used as food additives, and these uses have purportedly been abandoned by industry.⁴⁴ The FDA revoked authorizations for food contact use of 23 phthalates and two other substances used as plasticizers, adhesives, defoaming agents, lubricants, resins, and slimicides. ⁴⁵
- 77. Companies and/or individuals who manufacture or market cosmetics have a legal responsibility and duty to ensure the safety of their own products. Neither the law nor FDA regulations require specific tests to demonstrate the safety of individual products or ingredients,

⁴¹ Food and Drug Administration Cosmetic Act § 602 (1938)

⁴² Prohibited & Restricted Ingredients in Cosmetics, U.S. Food and Drug Administration, https://www.fda.gov/cosmetics/cosmetics-law-regulations/prohibited-restricted-ingredients-cosmetics

^{43 21} Code of Federal Regulations § 700.19

⁴⁴ § 87FR 31080

⁴⁵ *Phthalates in Food Packages and Food Contact Applications*, U.S. Food and Drug administration, https://www.fda.gov/food/food-ingredients-packaging/phthalates-food-packaging-and-food-contact-applications

and the law also does not require cosmetic companies to share their safety information with the FDA.

- 78. The FDA has consistently advised manufacturers to use whatever testing is necessary to ensure the safety of products and ingredients, which may be substantiated through (a) reliance on already available toxicological test data on individual ingredients and on product formulations that are similar in composition to the particular cosmetic, and (b) performance of any additional toxicological and other tests that are appropriate in light of such existing data and information. ⁴⁶
- 79. Except for color additives and ingredients prohibited or restricted by regulation, a manufacturer may use any ingredient in the formulation of a cosmetic, provided that (1) the ingredient and the finished cosmetic are safe under labeled or customary conditions of use, (2) the product is properly labeled, and (3) the use of the ingredient does not otherwise cause the cosmetic to be adulterated or misbranded under the laws the FDA enforces. 47
- 80. With respect to whether the product is properly labeled, Title 21 of the Code of Federal Regulations defines the manufacturer's duty to warn consumers of potential health hazards. Section 740.1 states that "[t]he label of a cosmetic product <u>shall</u> bear a warning statement whenever necessary or appropriate to prevent a health hazard that <u>may</u> be associated with the product." (Emphasis added). This warning mandate directly correlates with the broad authority of manufacturers over their own cosmetic products to ensure that products are safe under labeled or customary conditions of use, properly labeled, and not adulterated or misbranded under FDA laws.

⁴⁷ *Id*.

⁴⁶ FDA Authority Over Cosmetics: How Cosmetics Are Not FDA-Approved, but are FDA-Regulated, U.S. Food and Drug Administration, Mar. 3, 2005, https://www.fda.gov/cosmetics/cosmetics-laws-regulations/fda-authority-over-cosmetics-how-cosmetics-are-not-fda-approved-are-fda-regulated

In short, under the current regulatory framework in the United States, it is incumbent upon the manufacturers of cosmetic products, and them alone, to assess the safety and efficacy of their products, and to warn consumers any time a health hazard may be associated with their products. Here, a wealth of scientific information is available regarding long-term use of hair relaxers, straighteners and hair dyes as containing certain endocrine-disrupting chemicals, which should have alerted manufacturers of these products to the specific and dangerous harms associated with their products when used as intended, particularly in women of color. Defendants' Products contain no such warnings.

f. Endocrine-Disrupting Chemicals

- 82. The endocrine system is vital for life and influences nearly every cell, organ, and process in the body.⁴⁸ The endocrine system helps regulate all biological processes in the body from conception through adulthood, including the development of the brain and nervous system, the growth and function of the reproductive system, as well as metabolism and blood sugar maintenance.⁴⁹
- 83. The endocrine system is a tightly regulated system made up of glands that produce and release precise amounts of hormones that bind to receptors located on specific target cells throughout the body. ⁵⁰
- 84. Hormones, such as estrogen, testosterone, progesterone, and androgen, control or regulate critical biological processes.⁵¹

⁴⁸ Endocrine System: The Endocrine System Includes the Thyroid, Adrenals, and the Pituitary Gland, Science Direct, https://www.sciencedirect.com/topics/psychology/endocrine-system

⁴⁹ Endocrine Disruption, United States Environmental Protection Agency, Mar. 7, 2022, https://www.epa.gov/endocrine-disruption/what-endocrine-disruption

⁵⁰ *Id*.

⁵¹ *Id*.

- 85. When a hormone binds to a target cell's receptor, the receptor carries out the hormone's instructions, the stimulus, and either switches on or switches off specific biological processes in cells, tissues, and organs. ⁵²
- 86. The precise functioning of the endocrine system is vital to maintain hormonal homeostasis within the body. A slight variation in hormone levels can lead to significant adverse-health effects, including reproductive impairment and infertility, cancer, cognitive deficits, immune disorders, and metabolic syndrome. ⁵³
- 87. Endocrine disrupting chemicals ("EDCs") are chemicals, or chemical mixtures, which interfere with the normal activity of the endocrine system.
- 88. EDCs can act directly on hormone receptors as mimics or antagonists, or on proteins that control hormone delivery. 54
- 89. EDCs disrupt the endocrine system and interfere with the body's hormonal regulation and homeostasis in several ways.
- 90. EDCs can cause the body to operate as if there were a proliferation of a hormone and thus over-respond to the stimulus or respond when it was not supposed to by mimicking a natural hormone.
- 91. EDCs can increase or decrease the levels of the body's hormones by affecting the production, degradation, and storage of hormones.

⁵² *Id*.

⁵³ *Id.*; Michele La Merrill, et al., *Consensus on the Key Characteristics of Endocrine-Disrupting Chemicals as a Basis for Hazard Identification*, Nature Reviews Endocrinol, Nov. 12,2019, https://www.nature.com/aricles/s41574-019-0273-8

⁵⁴ Evanthia Diamanti-Kandarakis, et al., *Endocrine-Disrupting Chemicals: An Endocrine Society Scientific Statement*, Endocrine Reviews, June 30, 2009, https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2726844

- 92. EDCs can block the hormone's stimulus through inducing epigenetic changes, modifications to DNA that regulate whether genes are turned on or off or altering the structure of target cells' receptors. ⁵⁵
- 93. EDCs are known to cause numerous adverse human health outcomes including endometriosis, impaired sperm quality, abnormalities in reproductive organs, various cancers, altered nervous system and immune function, respiratory problems, metabolic issues, diabetes, obesity, cardiovascular problems, growth, and neurological and learning disabilities.⁵⁶
- 94. EDCs that mimic the effects of estrogen in the body may contribute to disease risk because exposure to estrogen, both endogenously and exogenously, is associated with uterine and ovarian cancers, and a woman's lifetime risk of developing the diseases increases with greater duration and cumulative exposure.
- 95. Natural and synthetic EDCs are present in hair products under the guise of "fragrance" and "perfumes," and thus enter the body when these products are applied to the hair and scalp. Studies exploring this issue have thus far classified EDCs as estrogens, phthalates, and parabens.
- 96. Indeed, numerous studies spanning more than two decades have demonstrated the adverse impact EDCs including Di-2ethyhexylphthalate have on the male and female reproductive

⁵⁵ Luis Daniel Martinez-Razo, et al., *The impact of Di-(2-ethylhexyl) Phthalate and Mono(2-ethylhexyl) Phthalate in placental development, function, and pathophysiology,* Environment International, January 2021,

https://www.sciencedirect.com/science/article/pii/S0160412020321838?via%3Dihub

⁵⁶ Endocrine Disrupting Chemicals (EDCs), Endocrine Society, Jan. 24, 2022, https://www.endocrine.org/patient-engagement/endocrine-library/edcs

system, such as inducing endometriosis, abnormal reproductive trace formation, decreased sperm counts and viability, pregnancy loss, and abnormal puberty onset.⁵⁷

g. Phthalates

- 97. Phthalates are used in a variety of cosmetics and personal care products. Phthalates are chemical compounds developed in the last century that are used to make plastics more durable. These colorless, odorless, oily liquids are also referred to as "plasticizers" based on their most common uses.
- 98. Phthalates also function as solvents and stabilizers in perfumes and other fragrance preparations. Cosmetics that may contain phthalates include nail polishes, hair sprays, aftershave lotions, cleansers, and shampoos.
- 99. Upon information and belief, at all relevant times herein, phthalates were used in Defendants' Products.
- 100. Phthalates are chemicals used to improve the stability and retention of fragrances and to help topical products stick to and penetrate the skin and hair. 58
- 101. Phthalates are known EDCs which interfere with natural hormone production, regulation, and degradation, and are detrimental to human health. ⁵⁹
- 102. Phthalates are commonly used by cosmetics and hair care product manufacturers to make fragrances and colors last longer, and to make hair more flexible after a product is applied, among other uses.

⁵⁷ Hee-Su Kim, et al., *Hershberger Assays for Di-2-ethylhexyl Phthalate and Its Substitute Candidates*, Dev Reproduction, Mar. 22, 2018, https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5915764/

Olivia Koski & Sheila Hu, Fighting Phthalates, National Resources Defense Council, April 20, 2022, https://www.nrdc.org/stories/fighting-phthalates

⁵⁹ Yufei Wang & Haifeng Qian, *Phthalates and Their Impacts on Human Health*, Healthcare (Basel) 9, 603, May 9, 2021, https://www.ncbi.nlm.nih.gov/pmc/articles/PMC8157593

- 103. Phthalates can be found in most products that have contact with plastics during production, packaging, or delivery. Despite short half-lives in tissues, chronic exposure to phthalates adversely influences the endocrine system and functioning of multiple organs, have negative long-term impacts on the success of pregnancy, child growth and development, and reproductive systems in both young children and adolescents. Several countries have imposed restrictions and regulations on some types of phthalates. ⁶⁰
 - 104. Phthalates have been shown to disrupt the endocrine system. ⁶¹
- 105. Defendants' Products referenced herein contain phthalates, including Di-2-ethylhexylphthalate. Di-2-ethylhexylphthalate⁶² ("DEHP") is a highly toxic manufactured chemical⁶³ that does not occur naturally in the environment. ⁶⁴
 - 106. DEHP is a phthalate. 65
- 107. Under the authority of the Fair Packaging and Labeling Act ("FPLA"), the FDA requires an ingredient declaration on cosmetic products sold at the retail level to consumers.
- 108. However, the regulations do not require the listing of the individual fragrance or flavor ingredients, meaning phthalates evade listing when combined with a fragrance. As a result, consumers, including Plaintiff, are not able to determine from the ingredient declaration on the

⁶⁰ *Id*.

⁶¹ *Id*.

⁶² Also known as Bis(2-ethylhexyl) phthalate

⁶³ Sai Rowdhwal & Jiaxing Chen, *Toxic Effects of DI-2-ethylhexyl Phthalate: An Overview*, Biomed Research International, Feb. 22, 2018,

 $[\]frac{https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5842715/\#:\sim:text=DEHP\%20is\%20noncovalent}{ly\%20bound\%20to,and\%20plastic\%20waste\%20disposal\%20sites}$

Toxicological Profile for Di(2_ethylhexyl) Phthalate (DEHP), U.S. Dept of Health and Human Services, January 2022, https://www.atsdr.cdc.gov/ToxProfiles/tp9.pdf (DEHP are listed as hazardous pollutants under the Clean Air Act.; DEHP is on the Proposition 65 list because it can cause cancer and birth defects or other reproductive harm).

⁶⁵ Di(2-ethylhexyl) phthalate (DEHP), Proposition 65, California. Gov, https://www.p65warnings.ca.gov/fact-sheets/di2-ethylhexylphthalate-dehp

label if phthalates were present in a fragrance used in the herein-referenced hair Products placed into the stream of commerce by Defendants and used by the Plaintiff.

- 109. Since 1999, the Centers for Disease Control and Prevention ("CDC") has found phthalates in individuals studied for chemical exposure. ⁶⁶ The National Toxicology Program (NTP) evaluated DEHP in the Fifteenth Report on Carcinogens and found it reasonably anticipated to be a human carcinogen based on sufficient evidence of carcinogenicity from studies in experimental animals, however the data available from cancer studies on humans was found to be inadequate to fully evaluate the relationship between human cancers and DEHP exposure. ⁶⁷ .
- 110. DEHP was first used in 1949 in the United States and was the most abundantly used phthalate derivative in the 20th century. ⁶⁸
- 111. DEHP does not covalently bind to its parent material. Non-covalent bonds are weak and, as a result, DEHP readily leaches from its products, increasing human exposure.⁶⁹
- 112. Humans can be exposed to DEHP through ingestion, inhalation, and dermal contact for their lifetimes, including intrauterine life. ⁷⁰

https://www.cdc.gov/exposurereport/pdf/Biomarker_Groups_Infographic-508.pdf

⁶⁶ *Biomarkers Groups*, National Report on Human Exposure to Environmental Chemicals, Center for Disease Control.

⁶⁷ NTP (National Toxicology Program). 2021. Report on Carcinogens, Fifteenth Edition. Research Triangle Park, NC: U.S. Department of Health and Human Services, Public Health Service. https://ntp.niehs.nih.gov/go/roc15 DOI: https://doi.org/10.22427/NTP-OTHER-1003

⁶⁸ Pinar Erkekoglu & Belma Kocer-Gumusel, *Environmental Effects of Endocrine-Disrupting Chemicals: A Special Focus on Phthalates and Bisphenol A*, Environmental Health Risk, June 16, 2016, https://www.intechopen.com/chapters/50234

⁶⁹ Katelyn H. Wong & Timur Durrani, *Exposures to Endocrine Disrupting Chemicals in Consumer Products – A Guide for Pediatricians*, Current Problems in Pediatric and Adolescent Health Care, Science Direct, May 2017,

https://www.sciencedirect.com/science/article/pii/S1538544217300822?via%3Dihub

⁷⁰ Schmidt, Juliane-Susanne, et al., *Effects of Di(2-ethylhexyl) Phthalate (DEHP) on Female Fertility and Adipoenesis in C3H/N Mice*, Environmental Health Perspective, May 15, 2012, http://www.ncbi.nlm.nih.gov/pmc/articles/PMC3440070

- 113. The Agency for Toxic Substances and Disease Registry ("ATSDR") estimates that the range of daily human exposure to DEHP is 3-30 µg/kg/day.⁷¹
- 114. The no-observed-adverse-effect level for DEHP to humans is 4.8 mg/kg bodyweight/day and the tolerated daily intake (TDI) is 48 μ g/kg bodyweight.⁷²

Endpoint	Cancer (NSRL)		Developmental and Reproductive Toxicity (MADL)	
Route of Exposure	Oral	Inhalation	Oral	Inhalation
DEHP	310 µg/day	N.C.	410 μg/day	N.C.

Source: OEHHA's safe harbor levels for TDCIPP, DBP, DEHP, benzene, and formaldehyde. N.C. = not calculated by OEHHA as of August 2020.⁷³

115. When DEHP enters in the human body, it breaks down into specific metabolites. The toxicity of DEHP is mainly attributed to its unique metabolites which include the primary metabolite, mono-(2-ethylhexyl) phthalate (MEHP) and secondary metabolites, mono-(2-ethyl-5-hydroxyhexyl) phthalate (MEHHP), and mono-(2-ethyl-5-oxohexyl) phthalate (MEOHP).⁷⁴

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Hannon, Patrick et al., *Daily Exposure to Di*(2-ethylhexl) *Phthalate Alters Estous Cyclicity and Accelerates Primordial Follicle Recruitment Potentially Via Dysregulation of the Phosphatidylinositiol 3-Kinase Signaling Pathway in Adult Mice*, Biology of Reproduction Volume 90, Issue 6, June 2014, 136, 1-11, https://academic.oup.com/biolreprod/article/90/6/136,%201-11/2514356

Yufei Wang & Haifeng Qian, *Phthalates and Their Impacts on Human Health*, Healthcare (Basel) 9(5):603, May 18, 2021, https://www.ncbi.nlm.nih.gov/pmc/articles/PMC8157593/
Aalekhya Reddam & David Volz, *Inhalation of Two Prop 65-listed Chemicals Within Vehicles May Be Associated with Increased Cancer Risk*, Environmental International Volume 149, April 2021, https://www.sciencedirect.com/science/article/pii/SO16041202100026X
A Focus on Di-and Mono-(2-ethylhexyl) *Phthalates and Their Metabolites in Hospitalized Patients: A Focus on Di-and Mono-(2-ethylhexyl) Phthalates Exposure from Intravenous Plastic Bags*. Toxics, 10(7), 357, https://pubmed.ncbi.nlm.nih.gov/35878262/; Istaf Sheikh, et. Al., Endocrine disruption: In silico perspectives of interactions of di-(2-ethylhexyl) phthalate and its five major metabolites with progesterone receptor. BMC Structural Biology Volume 16, Suppl 1, 16, Sept. 30, 2016, https://bmcstructbiol.biomedcentral.com/articles/10.1186/s12900-016-0066-4

- DEHP and its metabolites are known to cause significant adverse health effects including, but not limited to, endometriosis, developmental abnormalities, reproductive dysfunction, and infertility, various cancers, and metabolic syndrome within the human population and their future children.
- 117. Studies on the health effects of DEHP in laboratory animals used oral inhalation, and dermal exposures. ⁷⁷
- 118. The results of the studies indicate potential associations between DEHP exposure and the following negative health effects:
 - a) Reproductive effects: Epidemiological studies suggest a potential association between DEHP exposure and decreased serum testosterone and altered sperm parameters in males. Available studies on fertility effects in humans do not indicate an association between DEHP exposure and infertility. In animals, the available oral and inhalation studies provide evidence that the male reproductive system, particularly the testes, is susceptible to DEHP toxicity. Evidence from animal studies indicates decreased male and female fertility at high oral doses.
 - **b) Developmental effects.** Epidemiological studies suggest a potential association between reduced AGD (Anogenital distance) and testicular descent in male infants and prenatal DEHP exposure. In addition, human epidemiological studies provide mixed results for potential relationships between exposure to DEHP and preterm birth, early puberty, and delayed mental and psychomotor development in children. Studies in animals indicate that altered glucose homeostasis and the development of the reproductive system following early life exposure is a particularly sensitive target of DEHP toxicity.

⁽Other secondary metabolites include mono (2-ethyl-5-carboxypentyl) phthalate (5-cx-MEPP) and mono[2-[carboxymethyl)hexylphthalate (2-cx-MMHP)).

⁷⁵ Richardson, Kadeem et. Al., *Di*(2-ethylhexyl) Phthalate (*DEHP*) Alters Proliferation and Uterine Gland Numbers in the Uterine of Adult Expose Mice, Reproductive Toxicology, 77, 70-79, https://pubmed.ncbi.nlm.nih.gov/29458081/

Yufei Wang & Haifeng Qian, *Phthalates and Their Impacts on Human Health*, Healthcare (Basel) 9, 603, May 9, 2021, https://www.ncbi.nlm.nih.gov/pmc/articles/PMC8157593

Chapter 2: *Health Effects*, Toxicological profile for Di(2-ethylheyxl) phthalate (DEHP) (2001), https://www.atsdr.cdc.gov/ToxProfiles/tp9-c2.pdf

- 119. The global consumption of DEHP in 2020 was estimated at 3.07 million tons, and global demand for plasticizers continues to rise. The estimated global market of phthalates in 2020 was expected to reach \$10 Billion and would still be widely used in plasticizers.⁷⁸
- 120. Human epidemiology studies have shown a significant association between phthalates exposures and adverse reproductive health outcomes in both women and men.⁷⁹
- 121. DEHP exposure is also significantly related to insulin resistance, consistently higher systolic blood pressure, and other reproduction system problems including early menopause, low birth weight, pregnancy loss, and preterm birth. ⁸⁰
- 122. When it comes to the impacts on children, epidemiology studies on phthalate toxicity have focused primarily on pregnancy outcomes, genital development, semen quality, precocious puberty, thyroid function, respiratory systems, and neurodevelopment. ⁸¹
- 123. Since the turn of the century, restrictions on phthalates have been proposed in many Asian and western countries. In 2008, the US Congress announced the Consumer Protection Safety Act (CPSA) that permanently banned the products, especially children's toys and childcare articles, containing DEHP, DBP, and BBP at levels >0.1% by weight.⁸²

⁷⁸ *Id*.

⁷⁹ *Id*.

N.M. Grindler, et al., *Exposure to Phthalate, an Endocrine Disrupting Chemical, Alters the First Trimester Placental Methylome and Transcriptome in Women*, Scientific Reports Volume 8, April 17, 2018, https://doi.org/10.1038/s41598-018-24505-w

⁸² Consumer Product Safety Improvement Act of 2008, H.R. 4040, 110th Cong. (2008), https://www.congress.gov/110/plaws/publ1314/PLAW-110publ314.pdf

B. Injuries Associated with Exposure to Endocrine Disrupting Chemicals

a. Uterine Cancer

- 124. Uterine cancer is associated with phthalate metabolites found in chemical hair straightening and hair relaxer products.
- 125. Uterine cancer⁸³ is among the fourth most common cancer in women in developed countries, ⁸⁴ accounting for about 3% of all new cancer cases each year.⁸⁵
- 126. Every year around 65,000 women develop uterine cancer in the United States, out of which more than 90% are of endometrial origin. The mean age at diagnosis is 61 years. ⁸⁶
- 127. The incidence in Black women is twice that of White women. ⁸⁷ In addition, Black women with uterine cancer carry a poorer prognosis as compared to White women. ⁸⁸
- 128. Though death rates from other cancers in women have declined in recent years, death rates for uterine cancer have increased by more than 100% in the last 20 years, ⁸⁹
- 129. Indeed, new cases of uterine cancer have increased by 0.6 percent per year from 2010 to 2019, and death rates have risen an average of 1.7 percent per year during the same period.⁹⁰

⁸³ Otherwise known as endometrial carcinoma.

⁸⁴ Unaiza Faizan & Vijayadershan Muppidi, *Uterine Cancer*, In: StatPearls, National Library of Medicine, Jan 2022, https://www.ncbi.nlm.nih.gov.books/NBK562313

⁸⁵ Cancer State Facts: Uterine Cancer, National Cancer Institute, https://seer.cancer.gov/statfacts/html/corp.html

⁸⁶ *Id*.

⁸⁷ *Id*.

⁸⁸ Joel Sorosky, *Endometrial Cancer*, Obstetrics & Gynecology Volume 120, 383-97, Aug. 2012, https://pubmed.ncbi.nlm.nih.gov/22825101/

⁸⁹ Linda Duska, et al., *Treatment of Older Women With Endometrial Cancer: Improving Outcomes with Personalized Care*, American Society Clinical Oncology Educational Book, 35:164-74, 2016, https://pubmed.ncbi.nlm.nih.gov/27249697/

⁹⁰ Jack J. Lee, Rising Endometrial Cancer Rate Spur New Approaches to Prevention, National Cancer Institute: Division of Cancer Prevention, June 28, 2022, https://prevention.cancer.gov/news-and-events/blog/rising-endometrial-cancer

- 130. A groundbreaking study by the National Cancer Institute recently found that women who use chemical hair straightening or relaxing products have a considerably higher risk of contracting uterine cancer. 91
- 131. The study found that an estimated 1.64% of women who never used chemical hair straighteners or relaxers would go on to develop uterine cancer by the age of seventy; but for frequent users, that risk more than doubles, increasing to 4.05%. 92
- 132. These risks are more substantial among Black women, who use hair straightening and hair relaxing products the most, including Defendants' Products.

b. Ovarian Cancer

- 133. Ovarian Cancer is a rare disease, making up approximately 1% of new cancer cases, with around 20,000 new cases diagnosed in the United States in 2022. Approximately 1.1% of all women will be diagnosed with ovarian cancer. Of the 10.6 per 100,000 women per year who will be diagnosed with ovarian cancer, the death rate is 6.3 per 100,000 women a 49.7% survival rate.⁹³
- 134. While overall rates of ovarian cancer are slowly declining in the U.S., attributable in part to increased use of oral contraceptives, Black women continue to have the poorest survival rate across all ovarian cancer subtypes.⁹⁴

⁹¹ Che-Jung Chang, et al., *Use of Straighteners and Other Hair Products and Incident Uterine Cancer*, Journal of the National Cancer Institute, Oct. 17, 2022, https://pubmed.ncbi.nlm.nih.gov/36245087/

⁹² *Id*.

⁹³ Cancer State Facts; Ovarian Cancer, National Cancer Institute, https://seer.cancer.gov/statfacts/html/ovary.html.

⁹⁴ Park H.K., et al. *Recent Trends in Ovarian Cancer Incidence and Relative Survival in the United States by Race/Ethnicity and Histologic Subtypes*. Cancer Epidemiol. Biomarkers Prev. 2017;26 (10): 1511-1518. doi: 10.1158/1055-9965.EPI-17-0290.

Another recent publication from the researchers of the Sister Study found the risk of ovarian cancer doubled with frequent use (defined as greater than four times per year) of chemical hair straighteners/relaxers in the previous year as opposed to never use. (HR = 2.19)⁹⁵

136. While the study was not powered to detect differences based on race/ethnicity, among Black women the hazard ratios were elevated for ever use of straighteners (HR = 1.28) or perms (HR = 1.80). Further, the researchers noted that "given the much higher prevalence of use of these products, the impact of these results is more relevant for African American/Black women."

c. Endometriosis

- 137. Endometriosis is associated with phthalate metabolites found in chemical hair straightening and hair relaxer products.
- 138. For Black women in the U.S., endometriosis is one of the most common indications for major gynecological surgery such as a hysterectomy. It is associated with long hospital stays and high hospital costs. ⁹⁷
- 139. One study demonstrated Phthalate metabolites promote increased endometrial volume, often an indication of fibroids on ultrasound. ⁹⁸ DEHP increased volume risk by 33% and the sum of androgenic phthalates increased risk by 27%. ⁹⁹

⁹⁵ White A.J., et al. *Use of hair products in relation to ovarian cancer risk*. Carcinogenesis 2021; 42(9):1189-1195

⁹⁶ *Id*.

⁹⁷ M.C. Kyama, *The prevalence of endometriosis among African-American and African-indigenous women*. Gynecologic and obstetric investigation, Vol. 57(1) (2004), https://pubmed.ncbi.nlm.nih.gov/14974452

Amir R. Zota et al., *Phthalates exposure and uterine fibroid burden among women undergoing surgical treatment for fibroids: a preliminary study*, Fertility and sterility, Vol. 111(1) (2019), https://www.ncbi.nlm.nih.gov/pmc/articles/PMC6321778/

- 140. The function of the uterine lining, the endometrium, is based on cell-cell interactions under the instruction of steroid hormones. ¹⁰⁰ Endometriosis, a common cause of female infertility, occurs almost exclusively in menstruating women of reproductive age and often results from disruptions of this well-balanced hormonal cellular equilibrium. ¹⁰¹
- 141. It is estimated that 20% to 50% of women being treated for infertility have endometriosis. 102
- 142. Endometriosis is a painful, estrogen dependent disease resulting from the growth of endometrial glands and stroma outside the uterus that causes a chronic inflammatory reaction. ¹⁰³
- 143. During the follicular phase of the menstrual cycle, estrogen, working through estrogen receptors¹⁰⁴, induces growth of the endometrium. ¹⁰⁵

¹⁰⁰ L. Cobellis et al., *High plasma concentrations of di-(2-ethylhexl)-phthalate in women with endometriosis*, Human Reproduction, Vol. 18, Issue 7 (2003), 1512-1515, https://doi.org/10.1093/humrep/deg254

¹⁰¹ D.L. Olive and L. B. Schwartz, *Endometriosis*, The New England J. of Med., Vol. 328(24):1759-69 (1993), https://pubmed.ncbi.nlm.nih.gov/8110213/; K.G. Osteen and E. Sierra-Rivera, *Does disruption of immune and endocrine systems by environmental toxins contribute to development of endometriosis?*, Seminars in Reproductive Endocrinology, Vol. 15 (3):301-8 (1997) https://pubmed.ncbi.nlm.nih.gov/938389/

¹⁰² *Endometriosis*, World Health Organization (March 31, 2021), https://www.who.int/news-room/fact-sheets/detail/endometriosis

 $^{^{103}}$ Id

¹⁰⁴ Illaria Paterni et al., *Estrogen receptors alpha (ERα) and beta (Erβ): subtype-selective ligands and clinical potential,* Steroids, Vol. 90:13-29 (2014), https://pubmed.ncbi.nlm.nih.gov/24971815/

Kun Yu et al., *Estrogen Receptor Function: Impact on the Human Endometrium*, Frontiers in endocrinology, Vol. 13 (2022), https://pubmed.ncbi.nlm.nih.gov/35285981/

- 144. A developing fetus and the female reproductive tract are particularly susceptible to EDCs. ¹⁰⁶ EDCs are known to interfere with hormonal homeostasis, leading to alteration of estrogen signaling. ¹⁰⁷ Specifically, DEHP is known to cause enhanced estrogenic activity. ¹⁰⁸
- 145. DEHP exposure has been demonstrated to promote atypical cell proliferation and increased uterine volume (weight) in female mice. ¹⁰⁹
- 146. Numerous studies, spanning decades, establish that DEHP can lead to the development of endometriosis, as it is known to increase the viability, activity, proliferation, and migration of endometrial stromal cells, a required precondition of endometriosis. ¹¹⁰
- 147. Studies have shown that endometriotic women have significantly higher plasma DEHP concentrations than those without the disease.¹¹¹ A study that included a sample size of

Saniya Rattan et al., *Di*(2-Ethylhexyl) Phthalate Exposure During Prenatal Development Causes Adverse Transgenerational Effects on Female Fertility in Mice, Toxicol Sci., Vol. 163 (2) (2018), https://www.nci.nlm.nih.gov/pmc/articles/PMC5974785/

¹⁰⁷ Xueping Chen et al., *Toxicity and Estrogenic Endocrine Disrupting Activity of Phthalates and Their Mixtures*, Int'l J. Envitl. Res. And Pub. Health, 1(3(:3156-3168 (2014) https://doi.org/10.3390/ijerph110303156

¹⁰⁸ Chou CK, Huang HW, Yang CF, Dahms HU, Liang SS, Wang TN, Kuo PL, Hsi E, Tsai EM, Chiu CC. *Reduced camptothecin sensitivity of estrogen receptor-positive human breast cancer cells following exposure to di(2-ethylhexyl)phthalate (DEHP) is associated with DNA methylation changes*. Environ Toxicol. 2019 Apr;34(4):401-414. doi: 10.1002/tox.22694. Epub 2019 Feb 5. PMID: 30720231.

Juhye Kim, et al., *Chronic Low-Dose Nonylphenol or Di-(2-ethylhexyl) Phthalate has a Different Estrogen-like Response in Mouse Uterus, Development & Reproduction, Vol.* 22(4):379-391(2018), https://pubmed.ncbi.nlm.nih.gov/30680337/

L. Cobellis et al., *High plasma concentrations of di-(2-ethylhexl)-phthalate in women with endometriosis*, Human Reproduction, Vol. 18, Issue 7 (2003), 1512-1515, https://doi.org/10.1093/humrep/deg254 (Concluded that 92.6% of women with endometriosis tested had detectable levels of DEHP and/or its metabolite, MEHP.

approximately five hundred women living in various states observed that DEHP's metabolite MEHP was consistently associated with endometriosis. 112

C. Mrs. Armstrong's Use of Hair Relaxing Products

- 148. Mrs. Armstrong was first exposed to Defendants' EDCs and phthalate-laden Products in the early 1960's, at or around her teenage years, when she began using Defendants' Products.
- 149. Mrs. Armstrong used Defendants' Products by applying the product herself or with family following Defendants instructions, or by going to a professional salon with friends and having them apply Defendants' Products as instructed by Defendants.
- 150. Mrs. Armstrong continued using Defendants' Products until around 2018, using the Products every six to eight weeks throughout this time, resulting in around 50 years of consistent exposure.
- 151. Mrs. Armstrong would keep the Defendants' Products on her hair for the time recommended in the instructions.
- 152. There was never any indication on any of the Defendants' Products' packaging or otherwise that this normal use could and would cause her to develop uterine cancer and endometriosis, ultimately leading to her death.
 - 153. Mrs. Armstrong was diagnosed with uterine cancer on or about May 28, 2019.
- 154. On November 4th, 2019, Mrs. Armstrong underwent surgical treatment for her cancer including a total abdominal hysterectomy and bilateral salpingo-oophorectomy.

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Buck Louis G.M. et al., *Bisphenol A and phthalates and endometriosis: the Endometriosis: Natural History, Diagnosis and Outcomes Study*, Fertility and sterility, Vol. 100(1):162-9.e1-2 (2013), https://pubmed.ncbi.nlm.nih.gov/23579005/

- 155. Following this surgery Mrs. Armstrong underwent aggressive chemotherapy treatment for her cancer. However, the treatment was ultimately unsuccessful, and she succumbed to the disease less than three short months later, on January 28, 2020.
- 156. As a result of Defendant's acts and/or omissions, Mrs. Armstrong suffered extreme pain and suffering, emotional distress, and eventual death not even one year from her original diagnosis, from the metastasis of her uterine cancer and chemotherapy.

COUNT ONE – STRICT LIABILITY (FAILURE TO WARN)

- 157. Plaintiff repeats and realleges the allegations in Paragraphs 1-156 above, as if fully set forth herein.
- 158. At all pertinent times, the Defendants were manufacturing, marketing, testing, promoting, selling and/or distributing the Products in the regular course of business.
- 159. At all pertinent times, Plaintiff used the Products on her hair and scalp area, which is a reasonably foreseeable and intended use.
- At all pertinent times, Defendants in this action knew or should have known that the use of phthalates and other EDCs in their hair Products significantly increases the risk of diseases including but not limited to cancer and endometriosis, based upon scientific knowledge dating back for decades.
- At all pertinent times, including the time of sale and consumption, the Products, when put to reasonably foreseeable use, were in an unreasonably dangerous and defective condition because they failed to contain adequate and proper warnings and/or instructions regarding the increased risk of diseases, including but not limited to cancer and endometriosis associated with the use of Defendants' hair Products. Defendants failed to properly and adequately warn and instruct Plaintiff as to the inherent risks of these debilitating and life-altering conditions.

- 162. Had Plaintiff received a warning that the use of the Products would significantly increase her risk of developing uterine cancer and endometriosis, she would never have used them. As a proximate result of Defendants' design, manufacture, marketing, sale, and distribution of the Products, Plaintiff was injured catastrophically, and was caused severe pain, suffering, disability, impairment, loss of enjoyment of life, loss of care, comfort, economic damages and ultimately death.
- 163. The development of uterine cancer and endometriosis by Plaintiff was the direct and proximate result of the unreasonably dangerous and defective condition of the Products at the time of sale and consumption, including their lack of warnings; Plaintiff suffered injuries and damages including, but not limited to, physical and mental pain and suffering, medical expenses, and death.
- 164. Defendants' Products were defective because they failed to contain warnings and/or instructions, and breached express warranties, and/or failed to conform to express factual representations upon which Plaintiff justifiably relied in electing to use the Products. The defect or defects made the Products unreasonably dangerous to persons such as Plaintiff, who could reasonably be expected to use and rely upon such Products. As a result, the defect or defects were a producing cause of Plaintiff's injuries and damages.
- 165. Defendants' Products failed to contain, and to this day fail to contain, adequate warnings and/or instructions regarding the increased risk of diseases, including but not limited to cancer and endometriosis, with the use of their Products by women. Defendants continue to market, advertise, and expressly represent to the general public that it is safe for women to use their relaxer Products. These Defendants continue with these marketing and advertising campaigns despite having scientific knowledge that their Products contained chemicals and other ingredients

that could and would increase women's risk of developing these debilitating, life-altering, and potentially fatal diseases.

- 166. Plaintiff sustained the following damages as a foreseeable, direct, and proximate result of the Defendants' acts and/or omissions:
 - a. Economic losses including medical care and lost earnings; and
 - b. Noneconomic losses including physical and mental pain and suffering, emotional distress, inconvenience, loss of enjoyment and impairment of past quality of life.
 - c. At all times material to this action, Plaintiff Lafayette Armstrong is the surviving spouse of Barbara Armstrong. As a result of the negligence of Defendants, Plaintiff is entitled to recover for the wrongful death of Barbara Armstrong, an amount equal to the full value of the life of Barbara Armstrong without deduction for expenses or necessaries had she lived.
 - d. Plaintiff Lafayette Armstrong, individually and as the Personal Representative of the Estate of Barbara Armstrong, as evidenced by the copy of the Certificate of Appointment attached hereto as Exhibit "A", is entitled to recover on behalf of the Estate of Barbara Armstrong for the conscious pain and suffering of Barbara Armstrong, as well as any medical, funeral, or other related expenses incurred on her behalf as a direct and proximate result of the negligence of Defendants.

COUNT TWO – STRICT LIABILITY (DESIGN AND/OR MANUFACTURING DEFECT)

167. Plaintiff repeats and realleges the allegations in Paragraphs 1-156 above, as if fully set forth herein.

- 168. Defendants engaged in the design, development, manufacture, marketing, sale, and distribution of the Products in a defective and unreasonably dangerous condition to consumers, including Plaintiff.
- 169. Defendants caused the Products to enter the stream of commerce and to be sold through various retailers, where Plaintiff purchased the Products.
- 170. The Products were expected to, and did, reach consumers, including Plaintiff, without change in the condition in which they were manufactured and sold by Defendants and/or otherwise placed into the stream of commerce.
- 171. Plaintiff used the Products in a manner normally intended, recommended, promoted, and marketed by Defendants.
- 172. Products failed to perform safely when used by Plaintiff in a reasonably foreseeable manner, specifically increasing her risk of developing uterine cancer and endometriosis.
- 173. The propensity of phthalates and other endocrine disrupting chemicals to trigger and promote tumors and cancerous growths in women, thereby substantially increasing the risk of diseases, including but not limited to cancer and endometriosis, renders the Products unreasonably dangerous when used in the manner intended, and to an extent beyond what would be contemplated by the ordinary consumer.
- 174. Importantly, the Products are an inessential cosmetic product that do not treat or cure any disease or health condition. Further, safer alternatives, including fragrance free products, have been available for decades.
- 175. Defendants have known, or should have known, that the Products are unreasonably dangerous, but have continued to design, manufacture, sell, distribute, market, promote, and supply

the Products so as to maximize sales and profits all at the expense of public health and safety, in conscious disregard of the foreseeable harm to the consuming public, including Plaintiff.

- 176. Plaintiff sustained the following damages as a foreseeable, direct, and proximate result of the Defendants' acts and/or omissions:
 - a. Economic losses including medical care and lost earnings; and
 - b. Noneconomic losses including physical and mental pain and suffering, emotional distress, inconvenience, loss of enjoyment and impairment of past quality of life, and ultimately death.
 - c. At all times material to this action, Plaintiff Lafayette Armstrong is the surviving spouse of Barbara Armstrong. As a result of the negligence of Defendants, Plaintiff is entitled to recover for the wrongful death of Barbara Armstrong, an amount equal to the full value of the life of Barbara Armstrong without deduction for expenses or necessaries had she lived.
 - d. Plaintiff Lafayette Armstrong, individually and as the Personal Representative of the Estate of Barbara Armstrong, as evidenced by the copy of the Certificate of Appointment attached hereto as Exhibit "A", is entitled to recover on behalf of the Estate of Barbara Armstrong for the conscious pain and suffering of Barbara Armstrong, as well as any medical, funeral, or other related expenses incurred on her behalf as a direct and proximate result of the negligence of Defendants.

COUNT THREE – STRICT LIABILITY (DESIGN AND/OR MANUFACTURING DEFECT)

177. Plaintiff repeats and realleges the allegations in Paragraphs 1-156 above as if fully set forth herein.

- 178. At all relevant times Defendants engaged in the design, development, manufacture, marketing, sale, and distribution of the Products in a defective and unreasonably dangerous condition to consumers, including Plaintiff.
- 179. Defendants caused the Products to enter the stream of commerce and be sold through various retailers, where Plaintiff purchased the Products.
- 180. The Products were expected to, and did, reach consumers, including Plaintiff, without change in the condition in which they were manufactured and sold by Defendants and/or otherwise placed into the stream of commerce.
- 181. Plaintiff used the Products in a manner normally intended, recommended, promoted, and marketed by Defendants.
- 182. Products failed to perform safely when used by Plaintiff in a reasonably foreseeable manner, specifically increasing her risk of developing uterine cancer and endometriosis.
- 183. The propensity of phthalates and other endocrine disrupting chemicals to trigger and promote cancerous growths in women, thereby substantially increasing the risk of diseases including, but not limited to, cancer and endometriosis, renders the Products unreasonably dangerous when used in the manner intended, and to an extent beyond what would be contemplated by the ordinary consumer.
- 184. Importantly, the Products are an inessential cosmetic that do not treat or cure any disease or health condition. Further, safer alternatives, including fragrance free products, have been readily available for decades.
- 185. Defendants knew, or by the exercise of reasonable care should have known, that the Products are unreasonably dangerous. However, they have continued to design, manufacture, sell, distribute, market, promote, and supply the Products to maximize sales and profits at the

expense of public health and safety, in conscious disregard of the foreseeable harm to the consuming public, including Plaintiff.

- 186. Defendants owed a duty to all reasonably foreseeable users to design a safe product.
- 187. Defendants breached their duty by failing to use reasonable care in the design and/or manufacturing of their Products because the Products were unreasonably dangerous in that they increase the risks of tumors and cancerous growths in women, thereby substantially increasing the risk of diseases including but not limited to cancer and endometriosis. This renders the Products unreasonably dangerous when used in a manner that was intended by the Defendants, and to an extent beyond that which would be contemplated by the ordinary consumer.
- 188. Defendants also breached their duty by failing to use reasonable care by failing to use cost-effective, safer, and reasonably feasible alternative designs in the design and/or manufacturing of their Products.
- 189. A reasonable company under the same or similar circumstances would have designed a safer product.
- 190. Plaintiff sustained the following damages as a foreseeable, direct, and proximate result of the Defendants' acts and/or omissions:
 - a. Economic losses including medical care and lost earnings; and
 - b. Noneconomic losses including physical and mental pain and suffering, emotional distress, inconvenience, loss of enjoyment and impairment of past quality of life, and ultimately death.
 - c. At all times material to this action, Plaintiff Lafayette Armstrong is the surviving spouse of Barbara Armstrong. As a result of the negligence of Defendants, Plaintiff is entitled to recover for the wrongful death of Barbara Armstrong, an amount equal to the

full value of the life of Barbara Armstrong without deduction for expenses or necessaries had she lived.

d. Plaintiff Lafayette Armstrong, individually and as the Personal Representative of the Estate of Barbara Armstrong, as evidenced by the copy of the Certificate of Appointment attached hereto as Exhibit "A", is entitled to recover on behalf of the Estate of Barbara Armstrong for the conscious pain and suffering of Barbara Armstrong, as well as any medical, funeral, or other related expenses incurred on her behalf as a direct and proximate result of the negligence of Defendants.

COUNT FOUR – PRODUCTS LIABILITY (NEGLIGENT FAILURE TO WARN)

- 191. Plaintiff repeats and realleges the allegations in Paragraphs 1-156 above, as if fully set forth herein.
- 192. At all relevant times, Defendants engaged in the design, development, manufacture, marketing, sale, and distribution of the Products in a defective and unreasonably dangerous condition to consumers, including Plaintiff.
- 193. Defendants knew, or by the exercise of reasonable care, should have known the use of their Products was dangerous, harmful, and injurious when used by Plaintiff in a reasonably foreseeable manner.
- 194. Defendants knew, or by the exercise of reasonable care, should have known ordinary consumers such as Plaintiff would not have realized the potential risks and dangers of their Products, and that Products were likely to increase the risks of tumors and cancerous growths in women, thereby substantially increasing the risk of cancer and/or endometriosis, when used in the manner intended, and to an extent beyond that which would be contemplated by the ordinary consumer.

- 195. Defendants owed a duty to all reasonably foreseeable consumers to disclose the risks associated with the use of their Products.
- 196. Defendants breached their duty of care by failing to use reasonable care in providing adequate warnings on their Products, including that Products were likely to increase the risks of tumors and cancerous growths in women, thereby substantially increasing the risk of diseases including but not limited to cancer and endometriosis, when used in the manner intended and to an extent beyond that which would be contemplated by the ordinary consumer.
- 197. The failure of Defendants to adequately warn about their defective Products, and their efforts to misleadingly advertise through conventional avenues, created a danger of injuries described herein that were reasonably foreseeable at the time of design and/or manufacture and distribution.
- 198. At all relevant times, Defendants could have provided adequate warnings and instructions to prevent the harms and injuries set forth herein, such as providing full and accurate information about the Products in advertising.
- 199. A reasonable company under the same or similar circumstances would have warned and instructed of the dangers.
- 200. Plaintiff was injured as a direct and proximate result of Defendants' failure to warn and instruct because she would not have used the Products had she received adequate warnings and instructions that the Products could increase the risks of tumors and cancerous growths in women, thereby substantially increasing the risk of diseases including but not limited to cancer and/or endometriosis when used in the manner intended, and to an extent beyond that would be contemplated by the ordinary consumer.

- 201. Defendants' lack of adequate and sufficient warnings and instructions, and their inadequate and misleading advertising, was a substantial contributing factor in causing harm to Plaintiff.
- 202. Plaintiff sustained the following damages as a foreseeable, direct, and proximate result of the Defendants' acts and/or omissions:
 - a. Economic losses including medical care and lost earnings; and
 - b. Noneconomic losses including physical and mental pain and suffering, emotional distress, inconvenience, loss of enjoyment and impairment of past quality of life, and ultimately death.
 - c. At all times material to this action, Plaintiff Lafayette Armstrong is the surviving spouse of Barbara Armstrong. As a result of the negligence of Defendants, Plaintiff is entitled to recover for the wrongful death of Barbara Armstrong, an amount equal to the full value of the life of Barbara Armstrong without deduction for expenses or necessaries had she lived.
 - d. Plaintiff Lafayette Armstrong, individually and as the Personal Representative of the Estate of Barbara Armstrong, as evidenced by the copy of the Certificate of Appointment attached hereto as Exhibit "A", is entitled to recover on behalf of the Estate of Barbara Armstrong for the conscious pain and suffering of Barbara Armstrong, as well as any medical, funeral, or other related expenses incurred on her behalf as a direct and proximate result of the negligence of Defendants.

COUNT FIVE – NEGLIEGENCE (DESIGN AND/OR MANUFACTURING DEFECT)

203. Plaintiff repeats and realleges the allegations in Paragraphs 1-156 above, as if fully set forth herein.

- 204. At all relevant times, Defendants engaged in the design, development, manufacture, marketing, sale, and distribution of the Products in a defective and unreasonably dangerous condition to consumers, including Plaintiff.
- 205. Defendants caused the Products to enter the stream of commerce and to be sold through various retailers, where Plaintiff purchased the Products.
- 206. The Products were expected to, and did, reach consumers, including Plaintiff, without change in the condition in which they were manufactured and sold by Defendants and/or otherwise placed into the stream of commerce.
- 207. Plaintiff used the Products in a manner intended, recommended, promoted, and marketed by Defendants.
- 208. The Products failed to perform safely when used by Plaintiff in a reasonably foreseeable manner, specifically increasing her risk of developing uterine cancer and endometriosis.
- 209. The propensity of phthalates and other endocrine disrupting chemicals to trigger tumors and cancerous growths in women, thereby substantially increasing the risk of diseases including but not limited to cancer, and/or endometriosis, renders the Products unreasonably dangerous when used in the manner they were intended, and to an extent beyond that which would be contemplated by the ordinary consumer.
- 210. Importantly, the Products are an inessential cosmetic product that do not treat or cure any disease or health condition. Further, safer alternatives, including fragrance free products, have been readily available for decades.
- 211. Defendants knew, or by the exercise of reasonable care should have known, that the Products are unreasonably dangerous. However, they have continued to design, manufacture,

sell, distribute, market, promote, and supply the Products to maximize sales and profits at the expense of public health and safety in conscious disregard of the foreseeable harm to the consuming public, including Plaintiff.

- 212. Defendants owed a duty to all reasonably foreseeable users to design a safe product.
- 213. Defendants breached their duty by failing to use reasonable care in the design and/or manufacturing of their Products because Products were unreasonably dangerous in that they increase the risks of tumors and cancerous growths in women, thereby substantially increasing the risk of diseases including but not limited to cancer and/or endometriosis, rendering the Products unreasonably dangerous when used in the manner intended, and to an extent beyond that which would be contemplated by the ordinary consumer.
- 214. Defendants also breached their duty by failing to use reasonable care by failing to use cost-effective, safer, reasonably feasible alternative designs in the design and/or manufacturing of their Products.
- 215. A reasonable company under the same or similar circumstances would have designed a safer product.
- 216. Plaintiff sustained the following damages as a foreseeable, direct, and proximate result of the Defendants' acts and/or omissions:
 - a. Economic losses including medical care and lost earnings; and
 - b. Noneconomic losses including physical and mental pain and suffering, emotional distress, inconvenience, loss of enjoyment and impairment of past quality of life, and ultimately death.
 - c. At all times material to this action, Plaintiff Lafayette Armstrong is the surviving spouse of Barbara Armstrong. As a result of the negligence of Defendants, Plaintiff is

entitled to recover for the wrongful death of Barbara Armstrong, an amount equal to the full value of the life of Barbara Armstrong without deduction for expenses or necessaries had she lived.

d. Plaintiff Lafayette Armstrong, individually and as the Personal Representative of the Estate of Barbara Armstrong, as evidenced by the copy of the Certificate of Appointment attached hereto as Exhibit "A", is entitled to recover on behalf of the Estate of Barbara Armstrong for the conscious pain and suffering of Barbara Armstrong, as well as any medical, funeral, or other related expenses incurred on her behalf as a direct and proximate result of the negligence of Defendants.

COUNT SIX – NEGLIGENCE (NEGLIGENCE AND/OR GROSS NEGLIGENCE)

- 217. Plaintiff repeats and realleges the allegations in Paragraphs 1-156 above, as if fully set forth herein.
- 218. The Defendants' negligence and extreme carelessness includes, but is not limited to, their marketing, designing, manufacturing, producing, supplying, inspecting, testing, selling and/or distributing the Products in one or more of the following respects:
 - a. In failing to warn Plaintiff of the hazards associated with the use of the Products;
 - In failing to properly test their Products to determine adequacy and effectiveness of safety measures, if any, prior to releasing the Products for consumer use;
 - c. In failing to properly test their Products to determine the increased risk of diseases such as uterine cancer, ovarian cancer, and endometriosis during the normal and/or intended use of the Products;

- d. In failing to inform ultimate users such as Plaintiff as to the safe and proper methods of handling and using the Products;
- e. In failing to remove the Products from the market when Defendants knew or should have known the Products were defective;
- f. In failing to instruct the ultimate users, such as Plaintiff, as to the methods for reducing the type of exposure to the Products which caused increased risk of diseases such as uterine cancer, ovarian cancer, and endometriosis;
- g. In failing to inform the public in general and Plaintiff in particular of the known dangers of using the Products;
- h. In failing to advise users how to prevent or reduce exposure that caused increased risk for diseases such as uterine cancer, ovarian cancer, and endometriosis;
- In marketing and labeling the Products as safe for all uses despite knowledge to the contrary;
- j. In failing to act like a reasonably prudent company under similar circumstances. Each and all of these acts and omissions, taken singularly or in combination, were a proximate cause of the injuries and damages sustained by Plaintiff.
- 219. At all pertinent times, the Defendants knew or should have known that the Products were unreasonably dangerous and defective when put to their reasonably anticipated use.
- 220. Defendants' acts and/or omissions constitute gross negligence because they constitute a total lack of care and extreme departure from what a reasonably careful company would do in the same situation to prevent foreseeable harm to Plaintiff.

- 221. Defendants acted and/or failed to act willfully, and with conscious and reckless disregard for the rights and interests of Plaintiff, and their acts and omissions had a great probability of causing significant harm and in fact resulted in such harm to Plaintiff.
- 222. Plaintiff was injured as a direct and proximate result of negligence and/or gross negligence as described herein.
- 223. Defendants' negligence and/or gross negligence were a substantial factor in causing and/or contributing to Plaintiff's injuries.
- 224. Plaintiff sustained the following damages as a foreseeable, direct, and proximate result of the Defendants' acts and/or omissions:
 - a. Economic losses including medical care and lost earnings; and
 - b. Noneconomic losses including physical and mental pain and suffering, emotional distress, inconvenience, loss of enjoyment and impairment of past quality of life, and ultimately death.
 - c. At all times material to this action, Plaintiff Lafayette Armstrong is the surviving spouse of Barbara Armstrong. As a result of the negligence of Defendants, Plaintiff is entitled to recover for the wrongful death of Barbara Armstrong, an amount equal to the full value of the life of Barbara Armstrong without deduction for expenses or necessaries had she lived.
 - d. Plaintiff Lafayette Armstrong, individually and as the Personal Representative of the Estate of Barbara Armstrong, as evidenced by the copy of the Certificate of Appointment attached hereto as Exhibit "A", is entitled to recover on behalf of the Estate of Barbara Armstrong for the conscious pain and suffering of Barbara Armstrong, as well

as any medical, funeral, or other related expenses incurred on her behalf as a direct and proximate result of the negligence of Defendants.

COUNT SEVEN – NEGLIGENCE (NEGLIGENT MISREPRESENTATION)

- 225. Plaintiff repeats and realleges the allegations in Paragraphs 1-156 above, as if fully set forth herein.
- 226. Defendants had a duty to accurately and truthfully represent to consumers, Plaintiff, and the public that the Products had been tested and found to be safe and effective for use. The representations made by Defendants were in fact false.
- 227. Defendants failed to exercise ordinary care in the representations concerning the Products while they were involved in their manufacture, sale, testing, quality, assurance, quality control, and distribution in interstate commerce, because Defendants negligently misrepresented the Products' substantial risk of unreasonable, dangerous, and adverse side effects.
- 228. Defendants breached their duty in representing that the Products have no serious side effects.
- 229. As a foreseeable, direct and proximate result of the negligent misrepresentation of Defendants as set forth herein, Defendants knew, and had reason to know, that the Products had been insufficiently tested, or had not been tested at all, and that they lacked adequate and accurate warnings, and that they created a high risk, and/or higher than acceptable risk, and/or higher than reported and represented risk, of adverse side effects, including but not limited to uterine cancer and/or endometriosis.
- 230. Plaintiff sustained the following damages as a foreseeable, direct, and proximate result of the Defendants' acts and/or omissions:
 - a. Economic losses including medical care and lost earnings; and

- b. Noneconomic losses including physical and mental pain and suffering, emotional distress, inconvenience, loss of enjoyment and impairment of past quality of life, and ultimately death.
- c. At all times material to this action, Plaintiff Lafayette Armstrong is the surviving spouse of Barbara Armstrong. As a result of the negligence of Defendants, Plaintiff is entitled to recover for the wrongful death of Barbara Armstrong, an amount equal to the full value of the life of Barbara Armstrong without deduction for expenses or necessaries had she lived.
- d. Plaintiff Lafayette Armstrong, individually and as the Personal Representative of the Estate of Barbara Armstrong, as evidenced by the copy of the Certificate of Appointment attached hereto as Exhibit "A", is entitled to recover on behalf of the Estate of Barbara Armstrong for the conscious pain and suffering of Barbara Armstrong, as well as any medical, funeral, or other related expenses incurred on her behalf as a direct and proximate result of the negligence of Defendants.

COUNT EIGHT – VIOLATION OF THE GEORGIA FAIR BUSINESS PRACTICES ACT OF 1975 (O.C.G.A. § 10-1-390 eq seq.)

- 231. Plaintiff repeats and realleges the allegations in Paragraphs 1-156 above, as if fully set forth herein.
- 232. The Georgia Fair Business Practices Act of 1975 (O.C.G.A. § 10-1-390 eq seq.), states that, "Unfair or deceptive acts or practices in the conduct of consumer transactions and consumers acts or practices in trade or commerce are declared unlawful."
- 233. By the conduct described in detail above and incorporated herein, Defendant engaged in unfair or deceptive acts in violation of Georgia consumer protection laws.

- 234. Plaintiff purchased and used Defendants' Products primarily for personal use and thereby suffered ascertainable losses because of Defendants' actions in violation of the consumer protection laws.
- 235. Had Defendants not engaged in deceptive conduct described herein, Plaintiff would not have purchased and/or paid for Defendants' product and would not have incurred related injuries and damages.
- 236. Defendants engaged in wrongful conduct while at the same time obtaining, under false pretenses, monetary gain from Plaintiff for the Products that would not have been paid had Defendants not engaged in unfair and deceptive conduct.
- 237. Defendants engaged in unfair methods of competition and deceptive acts or practices that were proscribed by law, including the following:
 - a. Representing that goods or services have characteristics, ingredients, uses, benefits, or quantities that they do not have;
 - b. Advertising goods or services with the intent not to sell them as advertised; and
 - Engaging in fraudulent or deceptive conduct that created a likelihood of confusion or misunderstanding.
- 238. Defendants intended for Plaintiff to rely on their representations and advertisements regarding the Products in order to achieve monetary gain from Plaintiff through her purchase of the Products.
- 239. Plaintiff was injured by the cumulative and indivisible nature of Defendants' conduct. The cumulative effect of Defendants' conduct directed at Plaintiff and other consumers was to create demand for and sell the Products. Each aspect of Defendants' conduct combined to artificially create sales of the product.

- 240. Defendants have a statutory duty to refrain from unfair or deceptive acts or trade practices in the design, labeling, development, manufacture, promotion, and sale of the Products.
- 241. Had Defendants not engaged in the deceptive conduct described above, Plaintiff would not have purchased and/or paid for the Products and would not have incurred related injuries and damages.
- 242. Defendants' intentional, deceptive, unconscionable, and fraudulent representations and material omissions to Plaintiff, cosmetologists, physicians, and consumers, constituted unfair and deceptive acts and trade practices in violation of Georgia consumer protection laws.
- 243. Defendants' actions, as complained of herein, constitute unfair competition or unfair, unconscionable, deceptive, or fraudulent acts, or trade practices in violation of Georgia consumer protection laws.
- 244. Defendants have engaged in unfair competition or unfair or deceptive acts or trade practices or have made false representations in violation of Georgia consumer protection laws.
- 245. Under these statutes, Defendants are the suppliers, manufacturers, advertisers, and sellers, who are subject to liability under such legislation for unfair, deceptive, fraudulent, and unconscionable consumer sales practices.
- 246. Defendants violated the statutes that were enacted to protect consumers against unfair, deceptive, fraudulent, and unconscionable trade and business practices and false advertising, by knowingly and falsely representing that Defendants' Products were fit to be used for the purposes intended, when in fact they were defective and dangerous, and by other acts alleged herein. These representations were made in marketing and promotional materials.

- 247. The actions and omissions of Defendants alleged herein are uncured or incurable deceptive acts under the statutes enacted to protect consumers against unfair, deceptive, fraudulent, and unconscionable trade and business practices and false advertising.
- 248. Defendants had actual knowledge of the defective and dangerous condition of Defendants' Products and failed to take any action to cure such defective and dangerous conditions.
- 249. Plaintiff relied upon Defendants' misrepresentations and omissions in determining which products to use.
- 250. Defendants' deceptive, unconscionable, or fraudulent representations and material omissions to Plaintiff and other consumers constitute deceptive acts and practices.
- 251. By reason of the unlawful acts engaged in by Defendants, and as a direct and proximate result thereof, Plaintiff suffered ascertainable losses and damages.
- 252. Plaintiff sustained the following damages as a foreseeable, direct, and proximate result of the Defendants' acts and/or omissions:
 - a. Economic losses including medical care and lost earnings; and
 - b. Noneconomic losses including physical and mental pain and suffering, emotional distress, inconvenience, loss of enjoyment and impairment of past quality of life, and ultimately death.
 - c. At all times material to this action, Plaintiff Lafayette Armstrong is the surviving spouse of Barbara Armstrong. As a result of the negligence of Defendants, Plaintiff is entitled to recover for the wrongful death of Barbara Armstrong, an amount equal to the full value of the life of Barbara Armstrong without deduction for expenses or necessaries had she lived.

d. Plaintiff Lafayette Armstrong, individually and as the Personal Representative of the Estate of Barbara Armstrong, as evidenced by the copy of the Certificate of Appointment attached hereto as Exhibit "A", is entitled to recover on behalf of the Estate of Barbara Armstrong for the conscious pain and suffering of Barbara Armstrong, as well as any medical, funeral, or other related expenses incurred on her behalf as a direct and proximate result of the negligence of Defendants.

COUNT NINE - FRAUD

- 253. Plaintiff repeats and realleges the allegations in Paragraphs 1-156 above, as if fully set forth herein.
- 254. Defendants, who engaged in the development, manufacture, marketing, sale, and distribution of cosmetic and personal care products, including the Products, owed a duty to provide accurate and complete information regarding their Products.
- 255. Defendants fraudulently misrepresented the use of the Products as safe and effective, specifically:
 - a. Defendant McBride Laboratories Products Design Essentials are intentionally labeled as "Natural," when in fact they contain numerous unnatural chemicals.
 - b. Defendant L'Oreal Product Dark & Lovely is marketed as "Nourishing Care" and with Argan Oil and Vitamin E, implying that it contains natural, beneficial, and healthy ingredients.
- 256. On October 4, 2022, Yahoo, MSN and many other online news websites published an article titled "The Relaxer Box Girls: Where Are They Now", which revealed that many of the young black girls that were featured on relaxer boxes and advertisements throughout the 1980s, 1990s, and 2000s, never actually got their hair relaxed, and never used any of the Products in

question. Instead, they used other straightening tools, technology, and styling products to achieve the look of having used Defendants' Products, without having exposed themselves to the dangerous chemicals contained within Defendants' relaxers. Many of these women are wearing natural hairstyles today.¹¹³

- 257. Defendants knew that these misrepresentations and/or omissions were material, and that they were false, incomplete, misleading, deceptive, and deceitful when they were made.
- 258. Defendants made the misrepresentations and/or omissions for the purpose of deceiving and defrauding consumers, including Plaintiff, with the intention of having them act and rely on such misrepresentations and omissions.
- 259. Plaintiff relied, with reasonable justification, on the misrepresentations by Defendants, which induced her to purchase and use the Products on a regular basis for decades.
- 260. Defendants profited significantly from their unethical and illegal conduct that fraudulently induced Plaintiff and millions of other consumers to purchase a dangerous and defective product.
- 261. Defendants' actions, and Plaintiff's justifiable reliance thereon, were substantial contributing factors in causing injury and incurrence of substantial damage.
- 262. Plaintiff sustained the following damages as a foreseeable, direct, and proximate result of the Defendants' acts and/or omissions:
 - a. Economic losses including medical care and lost earnings; and

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¹¹³ The Relaxer Box Girls: Where Are They Now?, https://news.yahoo.com/relaxer-box-girls-where-now-192747557.html

- b. Noneconomic losses including physical and mental pain and suffering, emotional distress, inconvenience, loss of enjoyment and impairment of past quality of life, and ultimately death.
- c. At all times material to this action, Plaintiff Lafayette Armstrong is the surviving spouse of Barbara Armstrong. As a result of the negligence of Defendants, Plaintiff is entitled to recover for the wrongful death of Barbara Armstrong, an amount equal to the full value of the life of Barbara Armstrong without deduction for expenses or necessaries had she lived.
- d. Plaintiff Lafayette Armstrong, individually and as the Personal Representative of the Estate of Barbara Armstrong, as evidenced by the copy of the Certificate of Appointment attached hereto as Exhibit "A", is entitled to recover on behalf of the Estate of Barbara Armstrong for the conscious pain and suffering of Barbara Armstrong, as well as any medical, funeral, or other related expenses incurred on her behalf as a direct and proximate result of the negligence of Defendants.

COUNT TEN – FRAUDULENT CONCEALMENT

- 263. Plaintiff repeats and realleges the allegations in Paragraphs 1-156 above, as if fully set forth herein.
- 264. Defendants owed consumers, including Plaintiff, a duty to disclose all material facts fully and accurately regarding the Products, not to conceal material defects related thereto, not to place these defective Products into the stream of commerce, and to fully and accurately label product packaging. To the contrary, Defendants explicitly and/or implicitly represented that the Products were safe and effective.

- 265. Defendants actively and intentionally concealed and/or suppressed material facts, in whole or in part, to induce consumers, including Plaintiff, to purchase and use the Products and did so at her expense. Specifically, Defendants have been aware of the positive association between phthalates and other endocrine disrupting chemicals used in their Products and an increased risk of ovarian cancer, uterine cancer, and endometriosis demonstrated by epidemiology studies that showed exposure to phthalates in their Products enhance invasive and proliferative activities of endometrial cells.
- 266. Recent studies have established a statistically significant association between Defendants' Products and uterine fibroids and uterine cancer. The development of fibroids can also be a precursor to and increase the risk of uterine cancer in women.
- 267. Defendants made the misrepresentations and/or omissions for the purpose of deceiving and defrauding Plaintiff and with the intention of having her act and rely on such misrepresentations and/or omissions.
- 268. Defendants knew that their concealments, misrepresentations and/or omissions were material, and that they were false, incomplete, misleading, deceptive, and deceitful at the time they were made. Alternatively, Defendants concealed information, and/or made the representations with such reckless disregard for the truth that knowledge of the falsity can be imputed to them.
- 269. Defendants profited significantly from their unethical and illegal conduct that caused Plaintiff to purchase and habitually use a dangerous and defective product.
- 270. Defendants' actions and representations, and Plaintiff's justifiable reliance thereon, were substantial contributing factors in causing injury and incurrence of substantial damages.

- 271. Plaintiff sustained the following damages as a foreseeable, direct, and proximate result of the Defendants' acts and/or omissions:
 - a. Economic losses including medical care and lost earnings; and
 - b. Noneconomic losses including physical and mental pain and suffering, emotional distress, inconvenience, loss of enjoyment and impairment of past quality of life, and ultimately death.
 - c. At all times material to this action, Plaintiff Lafayette Armstrong is the surviving spouse of Barbara Armstrong. As a result of the negligence of Defendants, Plaintiff is entitled to recover for the wrongful death of Barbara Armstrong, an amount equal to the full value of the life of Barbara Armstrong without deduction for expenses or necessaries had she lived.
 - d. Plaintiff Lafayette Armstrong, individually and as the Personal Representative of the Estate of Barbara Armstrong, as evidenced by the copy of the Certificate of Appointment attached hereto as Exhibit "A", is entitled to recover on behalf of the Estate of Barbara Armstrong for the conscious pain and suffering of Barbara Armstrong, as well as any medical, funeral, or other related expenses incurred on her behalf as a direct and proximate result of the negligence of Defendants.

COUNT ELEVEN – BREACH OF EXPRESS WARRANTY

- 272. Plaintiff repeats and realleges the allegations in Paragraphs 1-156 above, as if fully set forth herein.
- 273. Defendants expressly warranted, through direct-to-consumer marketing, advertisements, and labels, that the Products were safe and effective for reasonably anticipated users.

- 274. The Products did not conform to these express representations because they can cause significant injury, including but not limited to uterine cancer, ovarian cancer, and endometriosis when used in the manner directed by Defendants.
- 275. Plaintiff sustained the following damages as a foreseeable, direct, and proximate result of the Defendants' acts and/or omissions:
 - a. Economic losses including medical care and lost earnings; and
 - b. Noneconomic losses including physical and mental pain and suffering, emotional distress, inconvenience, loss of enjoyment and impairment of past quality of life, and ultimately death.
 - c. At all times material to this action, Plaintiff Lafayette Armstrong is the surviving spouse of Barbara Armstrong. As a result of the negligence of Defendants, Plaintiff is entitled to recover for the wrongful death of Barbara Armstrong, an amount equal to the full value of the life of Barbara Armstrong without deduction for expenses or necessaries had she lived.
 - d. Plaintiff Lafayette Armstrong, individually and as the Personal Representative of the Estate of Barbara Armstrong, as evidenced by the copy of the Certificate of Appointment attached hereto as Exhibit "A", is entitled to recover on behalf of the Estate of Barbara Armstrong for the conscious pain and suffering of Barbara Armstrong, as well as any medical, funeral, or other related expenses incurred on her behalf as a direct and proximate result of the negligence of Defendants.

COUNT TWELVE – BREACH OF IMPLIED WARRANTIES

276. Plaintiff repeats and realleges the allegations in Paragraphs 1-156 above, as if fully set forth herein.

- 277. At the time the Defendants manufactured, marketed, labeled, promoted, distributed and/or sold the Products, the Defendants knew of the uses for which the Products were intended and impliedly warranted the Products to be of merchantable quality and safe for such use.
- 278. Defendants breached their implied warranties of the Products sold to Plaintiff because they were not fit for their common, ordinary, and intended uses.
- 279. Plaintiff sustained the following damages as a foreseeable, direct, and proximate result of the Defendants' acts and/or omissions:
 - a. Economic losses including medical care and lost earnings; and
 - b. Noneconomic losses including physical and mental pain and suffering, emotional distress, inconvenience, loss of enjoyment and impairment of past quality of life, and ultimately death.
 - c. At all times material to this action, Plaintiff Lafayette Armstrong is the surviving spouse of Barbara Armstrong. As a result of the negligence of Defendants, Plaintiff is entitled to recover for the wrongful death of Barbara Armstrong, an amount equal to the full value of the life of Barbara Armstrong without deduction for expenses or necessaries had she lived.
 - d. Plaintiff Lafayette Armstrong, individually and as the Personal Representative of the Estate of Barbara Armstrong, as evidenced by the copy of the Certificate of Appointment attached hereto as Exhibit "A", is entitled to recover on behalf of the Estate of Barbara Armstrong for the conscious pain and suffering of Barbara Armstrong, as well as any medical, funeral, or other related expenses incurred on her behalf as a direct and proximate result of the negligence of Defendants.

COUNT THIRTEEN - NEGLIGENT FAILURE TO RECALL

- 280. Plaintiff repeats and realleges the allegations in Paragraphs 1-156 above, as if fully set forth herein.
- 281. At all relevant times, Defendants designed, developed, managed, operated, inspected, tested (or not), marketed, advertised, promoted, disseminated, made publicly available, and/or benefitted from the Products and therefore, owed a duty of reasonable care to avoid causing harm to those who used the Products, such as Plaintiff.
- 282. Defendants knew or should have known through the exercise of reasonable care, the risks to consumers posed by the Products.
- 283. Defendants knew or, by the exercise of reasonable care, should have known use of the Products was harmful and had the potential to increase the risks of endometriosis and/or cancer in women, and as such, renders the Products unreasonably dangerous when used in the manner intended, and to an extent beyond that which would be contemplated by an ordinary consumer.
- 284. Defendants owed a duty to the users of the Products, including Plaintiff, to exercise reasonable care in conducting their business to properly and reasonably design, research, develop, manufacture, produce, process, assemble, inspect, supply, distribute, deliver, broker, market, warn, maintain, repair, modify, recall, retrofit, engineer, test, recommend, advertise, and/or make available the Products.
- 285. Defendants also owed a continuing duty to Plaintiff to remove, recall, or retrofit the unsafe and/or defective Products across the United States (including in Plaintiff's state and case).
- 286. Defendants knew or reasonably should have known that the Products were dangerous and not safe for use.

- 287. Defendants knew or, in the exercise of reasonable and ordinary care, should have known that the Products were defective and unsafe for Plaintiff, who is a person likely to use the Products for the purpose and in the manner for which the Products were intended to be used and for purposes reasonably foreseeable by Defendants.
- 288. At all times, Defendants negligently breached said duties and unreasonably and negligently allowed the Products to be used by Plaintiff without proper recall or warning.
- 289. Defendants to this day have not made any reasonable effort to remove and/or retrofit the serious safety risk posed by the Products to consumers.
- 290. In failing to properly recall and/or retrofit the Products, or even warn of the serious safety risks the Products pose to consumers and the public, Defendants have failed to act as a reasonable manufacturer, designer, or distributor would under the same or similar circumstances and failed to exercise reasonable care.
- 291. Plaintiff was injured as a direct and proximate result of the negligent conduct as described herein.
- 292. Plaintiff sustained the following damages as a foreseeable, direct, and proximate result of the Defendants' acts and/or omissions:
 - a. Economic losses including medical care and lost earnings; and
 - b. Noneconomic losses including physical and mental pain and suffering, emotional distress, inconvenience, loss of enjoyment and impairment of past quality of life, and ultimately death.
 - c. At all times material to this action, Plaintiff Lafayette Armstrong is the surviving spouse of Barbara Armstrong. As a result of the negligence of Defendants, Plaintiff is entitled to recover for the wrongful death of Barbara Armstrong, an amount equal to the

full value of the life of Barbara Armstrong without deduction for expenses or necessaries had she lived.

d. Plaintiff Lafayette Armstrong, individually and as the Personal Representative of the Estate of Barbara Armstrong, as evidenced by the copy of the Certificate of Appointment attached hereto as Exhibit "A", is entitled to recover on behalf of the Estate of Barbara Armstrong for the conscious pain and suffering of Barbara Armstrong, as well as any medical, funeral, or other related expenses incurred on her behalf as a direct and proximate result of the negligence of Defendants.

PRAYER FOR RELIEF

WHEREFORE, Plaintiff demands judgment against Defendants on each of the above referenced claims and causes of action, as follows:

- a. All damages allowed under Georgia law for the wrongful death of Barbara Armstrong, including, but not limited to the full value of Barbara Armstrong's life as permitted by O.C.G.A. § 51-4-1, et seq.; damages for funeral, medical, and other necessary expenses resulting from the injuries and death of Barbara Armstrong pursuant to O.C.G.A. § 51-4-5; and, damages for Barbara Armstrong's physical and mental pain and suffering, all in an amount to be determined at the trial of this action;
- b. Punitive and/or exemplary damages for the wanton, willful, fraudulent, reckless acts of the Defendants who demonstrated a complete disregard and reckless indifference for the safety and welfare of the general public and Barbara Armstrong in particular in an amount sufficient to punish Defendants and deter future similar conduct;
 - c. Prejudgment interest;
 - d. Post-judgment interest;

- e. Plaintiff's reasonable attorneys' fees;
- f. Plaintiff's costs of these proceedings; and
- g. Such other and further relief as this Court deems just and proper.

DEMAND FOR JURY TRIAL

Plaintiff hereby demands a trial by jury on all Counts and as to all issues and allegations presented herein.

On the ____ day of January, 2023.

Respectfully submitted,

/s/ Kenneth C. Wilson

Kenneth C. Wilson Georgia Bar No. 690235

BEASLEY ALLEN, CROW, METHVIN, PORTIS & MILES, P.C.

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Attorney for Plaintiff

STATE COURT OF DEKALB COUNTY, GA. 1/18/2023 5:49 PM E-FILED BY: Jodessea Allen

Exhibit A

STATE OF SOUTH CAROLINA) IN THE PROBATE COURT
COUNTY OF: <u>SUMTER</u>	CERTIFICATE OF APPOINTMENT
IN THE MATTER OF: BARBARA ARMSTRONG (Decedent)) CASE NUMBER: <u>2020ES4300221</u>)
	This is to certify that
	LAFAYETTE ARMSTRONG
	is/are the duly qualified
	PERSONAL REPRESENTATIVE SUCCESSOR PERSONAL REPRESENTATIVE SPECIAL ADMINISTRATOR
in the above matter and that this appointmental day of May, 2020, is in full force and effective	
RESTRICTIONS:	
Executed th	is 22nd day of May, 2020.

Do not accept a copy of this certificate without the raised seal of the Probate Court.