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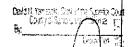
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27 28 Curtis G. Hoke (SBN 282465) THE MILLER FIRM, LLC 108 Railroad Ave. Orange, VA 22960 Telephone: (540) 672-4224 Facsimile: (540) 672-3055 choke@millerfirmllc.com FILED

JUN - 1 2015



Attorneys for Plaintiffs,

SUPERIOR COURT OF THE STATE OF CALIFORNIA

FOR THE COUNTY OF SANTA CLARA

(UNLIMITED JURISDICTION)

LYN KILLIAN ROSEMAN, ALEXANDER HOLCOMB, Individually, and as POA for WILLIE LEE HOLCOMB, Individually, and as Successor in Interest to the Estate of QUEEN ESTHER HOLCOMB, Deceased; CEDRIC HUNT, Individually, and as Executor of the Estate of EVELYN HUNT, Deceased; KARI L. MORRIS, Individually, and as Successor in Interest to the Estate of NANCY GUSTAVSON. Deceased; CARL PFEIFFER, Individually, and as Executor of the Estate of CAROL PFEIFFER, Deceased; VALERIE LOMBARDI; LINDA WONG-FARENBAUGH; MICA BAILEY, Individually, and as Successor in Interest to the Estate of CAROLYN T. JOHNSON, Deceased; PAM BAILEY, Individually, and as Successor in Interest to the Estate of LORENE G. HEPWORTH, Deceased; MALEA DAUGHTON,

Plaintiffs.

& MOZNHOL; MOZNHOL & MOZNHOL

Case No.:

16cvasaaa

COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

- I. STRICT LIABILITY FAILURE TO WARN (AGAINST IMERYS TALC)
- 2. STRICT LIABILITY FAILURE TO WARN (AGAINST JOHNSON & JOHNSON DEFENDANTS)
- 3. STRICT LIABILITY DESIGN DEFECT AND MANUFACTURING DEFECT (AGAINST IMERYS TALC)
- 4. STRICT LIABILITY
 MANUFACTURING DEFECT AND
 DESIGN DEFECT (AGAINST
 JOHNSON & JOHNSON
 DEFENDANTS)
- 5. NEGLIGENCE (AGAINST IMERYS

COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

1 2 3 3 4 4 5 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21	JOHNSON CONSUMER INC. F/K/A JOHNSON & JOHNSON CONSUMER COMPANIES, INC.; and IMERYS TALC AMERICA. INC. F/K/A LUZENAC AMERICA, INC., Defendants.	TALC) 6. NEGLIGENCE (AGAINST JOHNSON & JOHNSON DEFENDANTS) 7. BREACH OF EXPRESS WARRANTY (AGAINST JOHNSON & JOHNSON DEFENDANTS) 8. BREACH OF IMPLIED WARRANTY (AGAINST JOHNSON & JOHNSON DEFENDANTS) 9. CIVIL CONSPIRACY (AGAINST ALL DEFENDANTS) 10. FRAUD, FRAUDULENT MISREPRESENTATION, INTENTIONAL CONCEALMENT (AGAINST JOHNSON & JOHNSON DEFENDANTS) 11. NEGLIGENT MISREPRESENTATION (AGAINST ALL DEFENDANTS) 12. WRONGFUL DEATH (AGAINST ALL DEFENDANTS) 13. PUNYTIVE DAMAGES (AGAINST ALL DEFENDANTS) 14. LOSS OF CONSORTIUM (AGAINST ALL DEFENDANTS)
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COMPLAINT AND DEMAND FOR JURY TRIAL

COMES NOW, the above-captioned Plaintiffs (collectively referred to as "Plaintiffs"), and each of them, bring this Complaint and Demand for Jury Trial by and through their attorneys THE MILLER FIRM, LLC and complain and allege against Defendant JOHNSON & JOHNSON ONSUMER COMPANIES, INC., and IMERYS TALC AMERICA, INC. F/K/A LUZENAC AMERICA, INC. (collectively referred to as "Defendants") as follows:

SUMMARY OF ALLEGATIONS

- 1. This is a products flability action against the Defendants because Plaintiffs and Plaintiffs Decedents have suffered from and have passed away from the severe effects of Ovarian Cancer caused by Johnson & Johnson's baby powder and Shower-to-Shower products which were manufactured, mined, and/or marketed by Defendants (hereinafter, the "PRODUCTS") Defendants' PRODUCTS each contain tale powder, which caused Plaintiffs and Plaintiffs' Decedents to develop Ovarian Cancer after they used the PRODUCTS in their perineal area.
- 2. All Plaintiffs in this action seek recovery for damages as a result of developing ovarian cancer, which was directly and proximately caused by such wrongful conduct by Defendants, the unreasonably dangerous and defective nature of the PRODUCTS and talcum powder, and the attendant effects of developing ovarian cancer. All of the claims involve common legal and medical issues.
- At all relevant times, all Defendants were engaged in the research, development.
 manufacture, design, testing, sale and marketing of PRODUCTS, and introduced such products into interstate commerce with knowledge and intent that such products be sold in all States,

COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

including but not limited to the States of California, Texas, Indiana, Missouri, Kentucky, Alabama, Virginia, New York, Georgia, Michigan, and Pennsylvania.

4. Defendants concealed and continue to conceal their knowledge of tale powder's unreasonably dangerous risks from Plaintiffs, Plaintiffs' Decedents, other consumers, and the medical community. Specifically, Defendants failed to adequately inform Plaintiffs. Plaintiffs Decedents, consumers, and the medical community about the known risks of Ovarian Cancer associated with perincal use of the PRODUCTS.

PARTY PLAINTIFFS

- 5. Plaintiff LYN KILLIAN ROSEMAN is a competent individual over the age of 18 currently residing in California and hereby submits to the jurisdiction of this Court and alleges that Venue in this Court is proper. Ms. Roseman regularly used Defendants' PRODUCTS in her perineal region and suffered from severe physical, economic, and emotional injuries as a result of her use of Defendants' PRODUCTS, including but not limited to Ovarian Cancer diagnosed in 2000.
- 6. Plaintiff ALEXANDER HOLCOMB, INDIVIDUALLY, AND AS POA FOR WILLIE LEE HOLCOMB, INDIVIDUALLY, AND AS SUCCESSOR IN INTEREST TO THE ESTATE OF QUEEN ESTHER HOLCOMB, DECEASED, is a competent individual over the age of 18 currently residing in North Carolina and hereby submits to the jurisdiction of this Court and alleges that Venue in this Court is proper. Decedent QUEEN ESTHER HOLCOMB regularly used Defendants' PRODUCTS in her perincal region, was diagnosed with Ovarian Cancer and subsequently died in June, 2014.

- 7. Plaintiff CEDRIC HUNT, INDIVIDUALLY, AND AS EXECUTOR OF THE ESTATE OF EVELYN HUNT. DECEASED, is a competent individual over the age of 18 currently residing in Texas and hereby submits to the jurisdiction of this Court and alleges that Venue in this Court is proper. Decedent EVELYN HUNT regularly used Defendants' PRODUCTS in her perineal region, was diagnosed with Ovarian Cancer and subsequently died in June, 2014.
- 8. Plaintiff KARI L. MORRIS, INDIVIDUALLY, AND AS SUCCESSOR IN INTEREST TO THE ESTATE OF NANCY GUSTAVSON, DECEASED, is a competent individual over the age of 18 currently residing in Wisconsin and hereby submits to the jurisdiction of this Court and alleges that Venue in this Court is proper. Decedent NANCY GUSTAVSON regularly used Defendants' PRODUCTS in her perineal region, was diagnosed with Ovarian Cancer and subsequently died in June, 2014.
- 9. Plaintiff CARL PFEIFFER, INDIVIDUALLY, AND AS EXECUTOR OF THE ESTATE OF CAROL PFEIFFER, DECEASED, is a competent individual over the age of 18 currently residing in Michigan and hereby submits to the jurisdiction of this Court and alleges that Venue in this Court is proper. Decedent CAROL PFEIFFER regularly used Defendants PRODUCTS in her perincal region, was diagnosed with Overian Cancer and subsequently died in June, 2013.
- 10. Plaintiff VALERIE LOMBARDI is a competent individual over the age of 18 currently residing in Washington and hereby submits to the jurisdiction of this Court and alleges that Venue in this Court is proper. Ms. Lombardi regularly used Defendants' PRODUCTS in her perineal region and suffered from severe physical, economic, and emotional injuries as a result of her use of Defendants' PRODUCTS, including but not limited to Ovarian Cancer diagnosed in 2013.
- 11. Plaintiff LINDA WONG-FARENBAUGH is a competent individual over the age of 18 currently residing in Kansas and hereby submits to the jurisdiction of this Court and alleges that

Venue in this Court is proper. Ms. Wong-Farenbaugh regularly used Defendants' PRODUCTS in her perineal region and suffered from severe physical, economic, and emotional injuries as a result of her use of Defendants' PRODUCTS, including but not limited to Ovarian Cancer diagnosed in July, 2013.

- 12. Plaintiff MICA BAILEY, INDIVIDUALLY, AND AS SUCCESSOR IN INTEREST TO THE ESTATE OF CAROLYN T. JOHNSON, DECEASED, is a competent individual over the age of 18 currently residing in Florida and hereby submits to the jurisdiction of this Court and alleges that Venue in this Court is proper. Decedent CAROLYN T. JOHNSON regularly used Defendants' PRODUCTS in her perincal region, was diagnosed with Ovarian Cancer and subsequently died in July, 2014.
- 13. Plaintiff PAM BAILEY, INDIVIDUALLY, AND AS SUCCESSOR IN INTEREST TO THE ESTATE OF LORENE G. HEPWORTH, DECEASED, is a competent individual over the age of 18 currently residing in Utah and hereby submits to the jurisdiction of this Court and alleges that Venue in this Court is proper. Decedent LORENE G. HEPWORTH regularly used Defendants' PRODUCTS in her perineal region, was diagnosed with Ovarian Cancer and subsequently died in July, 2014.
- 14. Plaintiff MALEA DAUGHTON is a competent individual over the age of 18 currently residing in Maryland and hereby submits to the jurisdiction of this Court and alleges that Venuc in this Court is proper. Ms. Daughton regularly used Defendants' PRODUCTS in her perineal region and suffered from severe physical, economic, and emotional injuries as a result of her use of Defendants' PRODUCTS, including but not limited to Ovarian Cancer diagnosed in June 2013.

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 Defendant, JOHNSON & JOHNSON, is a New Jersey corporation with its principal place of business in the State of New Jersey.

- 16. At all relevant times, JOHNSON & JOHNSON was engaged in the business of manufacturing, marketing, testing, promoting, selling, and/or distributing the PRODUCTS. At all relevant times, Johnson & Johnson regularly transacted, solicited, and conducted business in all States of the United States, including the State of California.
- Defendant JOHNSON & JOHNSON CONSUMER COMPANIES, INC. is a New Jersey corporation with its principal place of business in the State of New Jersey.
- 18. At all relevant times, Johnson & Johnson Consumer Companies, Inc. was engaged in the business of manufacturing, marketing, testing, promoting, selling, and/or distributing the PRODUCTS. At all relevant times, JOHNSON & JOHNSON CONSUMER COMPANIES, INC regularly transacted, solicited, and conducted business in all States of the United States, including the State of California.
- Defendants JOHNSON & JOHNSON and JOHNSON & JOHNSON CONSUMER
 COMPANIES, INC. are collectively referred to herein as the "Johnson & Johnson Defendants".
- 20. Defendant IMERYS TALC AMERICA, INC. I/k/a LUZENAC AMERICA, INC. is a Delaware corporation with its principal place of business in the State of California specifically, its head office and laboratory are located at 1732 North First Street, Suite 450, San Jose, California 95112 (County of Santa Clara).

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27 28 All allegations regarding actions taken by Imerys Tale also include actions taken while that entity was known as Luzenac America, Inc.

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COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

Retailer Wal-Mart lists the lubels for Johnson's Bahy Powder, http://www.walmart.com/ip/Johnson-s-Bahy-

COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

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- 30. Plaintiffs used the PRODUCTS to dust their perineum for feminine hygiene purposes.

 This was an intended and foreseeable use of the PRODUCTS based on the advertising, marketing, and labeling of the PRODUCTS.
- 31. Upon information and belief, in 1971, the first study was conducted that suggested an association between tale and ovarian cancer. This study was conducted by Dr. WJ Henderson and others in Cardiff, Wales.
- 32. Upon information and belief, in 1982, the first epidemiologic study was performed on tale powder use in the female genital area. That study was conducted by Dr. Daniel Cramer and others. This study found a ninety-two percent increased risk of ovarian cancer with women who reported genital tale use. Shortly after this study was published, Dr. Bruce Semple of Johnson & Johnson came and visited Dr. Cramer about his study. Dr. Cramer advised Dr. Semple that Johnson & Johnson should place a warning on its taleum powders about the ovarian cancer risks so that women can make an informed decision about their health.
- 33. Upon information and belief, since approximately 1982, there have been approximately twenty-two additional epidemiologic studies providing data regarding the association of tale and ovarian cancer. Nearly all of these studies have reported an elevated risk of ovarian cancer associated with genital tale use in women.
- 34. Upon information and belief, in or about 1993, the United States National Toxicology Program published a study on the toxicity of non-asbestos form tale and found clear evidence of

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27 28 carcinogenic activity. Tale was found to be a carcinogen, with or without the presence of asbestos-like fibers.³

Upon information and belief, in response to the United States National Toxicology 35. Program's study, the Cosmetic Toiletry and Fragrance Association (CTFA), now known as the PCPC, formed the Tale Interested Party Task Force (TIPTF). Johnson & Johnson, Inc., Johnson & Johnson Consumer Companies, Inc., and Luzenac-now known as Imerys Tale-were members of the CTFA and were the primary actors and contributors of the TIPTF. The stated purpose of TIPTF was to pool financial resources of these companies in order to collectively defend tale use at all costs and to prevent regulation of any type over this industry. TIPTF hired scientists to perform biased research regarding the safety of tale. TIPTF members, including Johnson & Johnson and Luzenac, then edited these scientific reports hired by this group prior to the submissions of these scientific reports to governmental agencies. In addition, members of TIPTF knowingly released false information about the safety of tale to the consuming public and used political and economic influence on regulatory hodies regarding tale. These activities were conducted by these companies and organizations over the past four decades in an effort to prevent regulation of tale and to create confusion to the consuming public about the true hazards of tale and its association to ovarian cancer.

³ Inhalation Toxicology Research Institute Annual Report, 1993 – 1994.

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Upon information and belief, on or about November 19, 1994, the Cancer Prevention 36. Coalition sent a letter to then Johnson & Johnson C.E.O. Ralph Larsen, urging him to substitute cornstarch for talcum powder products and to label its products with a warning on cancer risks.4

Upon information and belief, in or about 1996, the FDA requested that the condom industry stop dusting condoms with tale due to the health concerns that studies linked tale to ovarian cancer. Upon this request, all U.S. manufacturers discontinued the use of tale in its condom manufacturing process to reduce the potential health hazards to women.5

Upon information and belief, in or about 1990, the U.S. Food and Drug Administration (FDA) asked manufacturers to voluntarily stop putting tale on surgical gloves because mounting scientific evidence showed that it caused adhesions in surgical patients.5

Upon information and belief, in or about February 2006, the International Agency for 39, Research on Cancer (IARC), the specialized cancer agency of the World Health Organization published a paper whereby they classified perineal use of tale-based body powder as a "Group 2B" human carcinogen. IARC, which is universally accepted as the international authority on cancer issues, concluded that studies from around the world consistently found an increased risk of ovarian cancer in women who used tale in perincal areas. IARC determined that between 16-

Petition Seeking a Cancer Warning on Cosmetic Tale PRODUCTS, May 13, 2008 hugs (www.persmesseer.com/publications.pdf/EDNA). Citter (sleCr.Ca. may F18.pdf.

"A Women's Campaign Against Tale on Condoms," Philly.com. http://aris.des.phills.com/1996-01-08 tiving 25052370 1 tale condomissorarians another.

⁷ IARC, "Perinent use of tate-based body powder (Group 2B)," available at https//monososubusians.fr/4.NG/Monovombs/PDE/s/v/Lode.pdf.

 52% of women worldwide used tale to dust their perineum and found an increased risk of ovarian cancer in women tale users ranging from 30-60%.

- 40. Upon information and belief, in or about 2006, the Canadian government, under The Hazardous PRODUCTS Act and associated Controlled PRODUCTS Regulations, classified tale as a "D2A," "very toxic," "cancer-causing" substance under its Workplace Hazardous Materials Information System (WHMIS). Ashestos is also classified as "D2A."
- 41. Upon information and belief, in or about 2006, Defendant Imerys Tale began placing a warning on the MSDS it provided to the Johnson & Johnson Defendants regarding the tale it sold to them for use in the PRODUCTS. The MSDSs not only provided the warning information about the IARC classification but also included warning information regarding "States Rights to Know" and warning information about the Canadian Government's D2A classification of tale. Although the Johnson & Johnson Defendants admittedly received these MSDs, they never passed this warning information on to consumers. On September 26, 2012, the corporate representative for Imerys testified in open court that his company exclusively supplied the Johnson & Johnson Defendants with tale used for its baby powder products and that ovarian cancer is a potential hazard associated with women's perincal use of tale-based body powders such as the PRODUCTS. Despite this, the Johnson & Johnson defendants continue to mislead consumers, such as Plaintiffs, maintaining that tale is safe for personal use.

See, e.g., https://www.sadety.outcaregonymispent.com/ingration-infe/order/tale ("tale can be used safety in personal care products"; We want to assure women and caregivers who use our tale products that numerous studies support its safety, and these include assessments by external experts in addition to our company testing. Many research papers and epidemiology studies have specifically evaluated tale and perineal use and these studies have found tale to be safe")

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Presently, the National Cancer Institute¹¹ and the American Cancer Society¹² list genital The Gilda Radner Familial Ovarian Cancer Registry, Roswell Park Center Institute, and Cancer Prevention Coalition "Petition Seeking a Cancur Warning on Cosmetic Tale PRODUCTS" submitted to the 14 COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

 likely to be read and understood by the ordinary individual under customary conditions of purchase and use.

The PRODUCTS, in violation of 21 C.F.R. § 740.10, do not conspicuously state on their principal display panel that the safety of the PRODUCTS have not been determined and/or that the safety of the PRODUCTS' principal ingredients have not been determined.

COUNT 1 - STRICT LIABILITY FAILURE TO WARN

(Against Imerys Tale)

- 54. Plaintiffs incorporate by reference all other paragraphs in this Complaint as if set forth fully herein.
- 55. At all relevant times, Imerys Tale mined and sold tale to the Johnson & Johnson Defendants with full knowledge that the Johnson & Johnson Defendants were then packaging the tale and selling to consumers as the PRODUCTS and consumers of the PRODUCTS were using it to powder their perineal regions.
- 56. At all relevant times, by mining tale and supplying that tale to the Johnson & Johnson Defendants for use in the PRODUCTS, Imerys Tale was knowingly an integral part of the overall manufacture, design, and production of the PRODUCTS and their introduction into the stream of interstate commerce.
- 57. At all relevant times, linerys Tale knew or should have known of the unreasonably dangerous and careinogenic nature of the tale it was selling to the Johnson & Johnson Defendants, especially when applied to a woman's perineal regions, and it knew or should have known that Johnson & Johnson was not warning its consumers of this danger.

58. At all relevant times, Imerys Tale knew or should have known that the use of the PRODUCTS significantly increase the risk of ovarian cancer in women based upon scientific knowledge dating back until at least 1971.

- 59. At all relevant times, the PRODUCTS were defective and unreasonably dangerous when used in a reasonably foreseeable manner because, despite Imerys Tale's knowledge that the PRODUCTS were carcinogenic and could lead to an increased risk of ovarian cancer, Imerys Tale failed to provide adequate warning and/or instruction to consumers, including Plaintiff, regarding the increased risk of ovarian cancer associated with the use of the PRODUCTS when applied to the perincal area.
- 60. Had Plaintiffs received warning or instruction regarding the increased risk of ovarian cancer associated with the PRODUCTS when applied to the perincul area, Plaintiffs would not have used the PRODUCTS in this manner.
- 61. Due to the absence of any warning or instruction by the Defendants as to the significant health and safety risks posed by the PRODUCTS as described herein. Plaintiffs were unaware that the PRODUCTS created an increased risk of ovarian cancer, as this danger was not known to the general public.
- 62. As a direct and proximate result of Imerys Tale's failure to warn Plaintiffs of the increased risk of ovarian cancer associated with the PRODUCTS when applied to the perineal area, despite their actual knowledge of this material fact. Plaintiffs developed ovarian cancer and have been injured catastrophically and have been caused severe and permanent pain, suffering, disability, impairment, loss of enjoyment of life, loss of care, comfort and economic damages.

63. WHEREFORE, Plaintiffs demand judgment against Imerys Tale for compensatory, treble damages pursuant to California Civil Code Section 3345, and punitive damages in excess of the jurisdictional minimum of this Court, together with interest, costs of suit, attorneys' fees, and all such other relief, as this Court deems proper.

COUNT II - STRICT LIABILITY FAILURE TO WARN

(Against Johnson & Johnson Defendants)

- 64. Plaintiffs incorporate by reference all other paragraphs in this Complaint as if set forth fully herein.
- 65. At all relevant times, the Johnson & Johnson Defendants were engaged in the business of manufacturing, marketing, testing, promoting, selling and/or distributing, and otherwise introducing into the stream of interstate commerce, the PRODUCTS.
- 66. At all relevant times, the Johnson & Johnson Defendants knew or should have known that the use of the PRODUCTS in the female perineal area significantly increased the risk of ovarian cancer in women based upon scientific knowledge dating back until at least 1971.
- 67. At all relevant times, the PRODUCTS, manufactured and supplied by the Johnson & Johnson Defendants, were defective and unreasonably dangerous because, despite the Johnson & Johnson Defendants' knowledge that its PRODUCTS were carcinogenic and could lead to an increased risk of ovarian cancer when applied to the female perineal area, a reasonably foresecable use of the PRODUCTS, the Johnson & Johnson Defendants failed to provide adequate warning or instruction to consumers, including Plaintiffs, regarding the increased risk of ovarian cancer when the PRODUCTS are applied to the female perineal area.

68.	At all relevant times, Plaintiffs used the PRODUCTS to powder their perineal areas, a us
that wa	as reasonably foresecable and for which the PRODUCTS were supplied.

- 69. Had Plaintiffs received warning and/or instruction from the Johnson & Johnson Defendants regarding the increased risk of ovarian cancer associated with the PRODUCTS when applied to the perineal area, Plaintiffs would not have used the PRODUCTS in this manner.
- 70. Due to the absence of any warning or instruction by the Johnson & Johnson Defendants as to the significant health and safety risks posed by the PRODUCTS as described herein. Plaintiffs were unaware that the PRODUCTS created an increased risk of ovarian cancer, as this danger was not known to the general public.
- 71. As a direct and proximate result of Johnson & Johnson Defendants' failure to warm Plaintiffs of the increased risk of ovarian cancer associated with the PRODUCTS when applied to the perineal area, despite their actual knowledge of this material fact, Plaintiffs developed ovarian cancer and have been injured catastrophically and have been caused severe and permanent pain, suffering, disability, impairment, loss of enjoyment of life, loss of care, comfort and economic damages.
- 72. WHEREFORE, Plaintiffs demand judgment against the Johnson & Johnson Defendants for compensatory, treble damages pursuant to California Civil Code Section 3345, and punitive damages in excess of the jurisdictional minimum of this Court, together with interest, costs of suit, attorneys' fees, and all such other relief, as this Court deems proper.

(Against Imerys Talc)

- Plaintiffs incorporate by reference all other paragraphs in this Complaint as if set forth fully herein.
- 74. At all relevant times, Defendant imerys Tale was engaged in the business of mining and distributing talcum to Johnson & Johnson Defendants for use in the PRODUCTS, and they were knowingly an integral part of the overall manufacture, design, and production of the PRODUCTS and their introduction into the stream of interstate commerce.
- 75. At all relevant times, the PRODUCTS were expected to and did reach Plaintiffs without a substantial change in their condition.
- 76. At all relevant times, the PRODUCTS were defectively and improperly manufactured and designed by Imerys Tale in that, when Imerys Tale supplied its tale product to Johnson & Johnson with full knowledge that Johnson & Johnson would use its tale in formulating the PRODUCTS and that the tale would be the primary ingredient in the PRODUCTS, the foreseeable risks of the PRODUCTS far outweighed the benefits associated with their design and formulation.
- 77. At all relevant times, the PRODUCTS were defectively manufactured and designed by Imerys Tale in that their design and formulation is more dangerous than an ordinary consumer would expect when used in an intended and reasonably foresecable manner.

COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

78. At all relevant times, the PRODUCTS created significant risks to the health and safety of consumers that far outweigh the risks posed by other products on the market used for the same therapeutic purpose.

79. As a direct and proximate result of the defective design and manufacture of the PRODUCTS. Plaintiffs developed ovarian cancer and have been injured catastrophically and have been caused severe and permanent pain, suffering, disability, impairment, loss of enjoyment of life, loss of care, comfort and economic damages.

80. WHEREFORE, Plaintiffs demand judgment against Imerys Tale for compensatory, treble damages pursuant to California Civil Code Section 3345, and punitive damages in excess of the jurisdictional minimum of this Court, together with interest, costs of suit, attorneys' fees, and all such other relief, as this Court deems proper.

COUNT IV - STRICT LIABILITY MANUFACTURING DEFECT AND DESIGN

DEFECT

(Against Johnson & Johnson Defendants)

- Plaintiffs incorporate by reference all other paragraphs in this Complaint as it set forth fully herein.
- 82. At all relevant times, the Johnson & Johnson Defendants were engaged in the business of manufacturing, formulating, creating, designing, testing, labeling, packaging, supplying, marketing, promoting, selling, advertising, and otherwise introducing the PRODUCTS into the stream of interstate commerce, which they sold and distributed throughout the United States.

COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

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substantial change in condition.

- 84. At all relevant times, the PRODUCTS were defectively and improperly manufactured and designed by the Johnson & Johnson Defendants in that, when the PRODUCTS left the hands of the Johnson & Johnson Defendants, the foreseeable risks of the PRODUCTS far outweighed the benefits associated with their design and formulation.
- 85. At all relevant times, the PRODUCTS were defectively manufactured and designed by the Johnson & Johnson Defendants in that their design and formulation is more dangerous than an ordinary consumer would expect when used in an intended and reasonably foresceable manner.
- 86. At all relevant times, the PRODUCTS created significant risks to the health and safety of consumers that far outweigh the risks posed by other products on the market used for the same therapeutic purpose.
- 87. At all relevant times, a reasonable and safer afternative design existed, which could have feasibly been employed by the Johnson & Johnson Defendants to manufacture a product with the same therapeutic purpose as the PRODUCTS. Despite knowledge of this reasonable and safer alternative design, the Johnson & Johnson Defendants failed to after the PRODUCTS* design and formulation. The magnitude of the danger created by the PRODUCTS far outweighs the costs associated with using an afternative, safer design.
- 88. As a direct and proximate result of the defective design and manufacture of the PRODUCTS, Plaintiff's developed ovarian cancer and have been injured catastrophically and have been caused severe and permanent pain, suffering, disability, impairment, loss of enjoyment of life, loss of care, comfort and economic damages.

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WHEREFORE, Plaintiffs demand judgment against the Johnson & Johnson Derendants 89. for compensatory, treble damages pursuant to California Civil Code Section 3345, and punitive damages in excess of the jurisdictional minimum of this Court, together with interest, costs of suit, attorneys' fees, and all such other relief, as this Court deems proper.

COUNT V-NEGLIGENCE

(Against Imervs Tale)

- Plaintiffs incorporate by reference all other paragraphs in this Complaint as if set forth 90. fully herein.
- At all relevant times, linerys Tale had a duty to exercise reasonable care to consumers including Plaintiffs herein, in the design, development, manufacture, testing, inspection, packaging, promotion, marketing, distribution, labeling and/or sale of the PRODUCTS.
- 92. At all relevant times, Imerys Talc mined and sold talc to the Johnson & Johnson Defendants, which it knew was then being packaged and sold to consumers as the PRODUCTS by the Johnson and Johnson Defendants. Further, Imerys Tale knew that consumers of the PRODUCTS were using it to powder their perineal regions.
- At all relevant times, Imerys Tale knew or should have known that the use of the PRODUCTS in the perineal area significantly increases the risk of ovarian cancer based upon scientific knowledge dating back to the 1971.
- At all relevant times, Imerys Tale knew that Johnson & Johnson Defendants were not providing warnings to consumers of the PRODUCTS of the risk of ovarian cancer posed by tald contained therein.

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COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

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At all relevant times, Imerys Tale was negligent in providing tale to the Johnson & 95. Johnson Defendants. Imerys Talc possessed information on the carcinogenic properties of talc including its risk of causing ovarian cancer, Imerys Tale was negligent because it knew that the tale they provided to Johnson & Johnson Defendants would be used in the PRODUCTS, but they did not adequately take steps to ensure that ultimate consumers of the PRODUCTS, including Plaintiffs, received the information that Imerys Tale possessed on the careinogenic properties of talc.

- As a direct and proximate result of Imerys Tale's negligence, Plaintiffs developed 96. ovarian cancer and have been injured catastrophically and have been caused severe and permanent pain, suffering, disability, impairment, loss of enjoyment of life, loss of care, comfort and economic damages.
- 97. WHEREFORE, Plaintiffs demand judgment against Imerys Talc for compensatory, treble damages pursuant to California Civil Code Section 3345, and punitive damages in excess of the jurisdictional minimum of this Court, together with interest, costs of suit, attorneys' fees, and all such other relief, as this Court deems proper,

COUNT VI - NEGLIGENCE

(Johnson & Johnson Defendants)

- 98. Plaintiffs incorporate by reference all other paragraphs in this Complaint as if set forth fully herein.
- 99. At all relevant times, the Johnson & Johnson Defendants breached their duty to Plaintiffs and were otherwise negligent in 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 Plaintiffs of the hazards associated with the use of the PRODUCTS;
- In failing to properly test their products to determine adequacy and effectiveness or safety measures, if any, prior to releasing the PRODUCTS for consumer use;
- In failing to properly test the PRODUCTS to determine the increased risk of ovarian cancer during the normal and/or intended use of the PRODUCTS;
- d. In failing to inform ultimate users, such as Plaintiffs, as to the safe and proper methods of handling and using the PRODUCTS;
- e. In failing to remove the PRODUCTS from the market when the Defendants knew or should have known the PRODUCTS were defective:
- f. In failing to instruct the ultimate users, such as Plaintiffs, as to the methods for reducing the type of exposure to the PRODUCTS which caused increased risk of ovarian cancer;
- g. In failing to inform the public in general and the Plaintiffs in particular of the known dangers of using the PRODUCTS for dusting the perincum;
- h. In failing to advise users how to prevent or reduce exposure that caused an increased risk for ovarian cancer:
- In marketing and labeling the PRODUCTS as safe for all uses despite knowledge to the contrary;
- In failing to act like a reasonably prudent company under similar circumstances;
- k. In failing to use a safer alternative to tale in the PRODUCTS, such as cornstarch.

Each and all of these acts and omissions, taken singularly or in combination, were a proximate cause of the injuries and damages sustained by Plaintiffs.

that the PRODUCTS were unreasonably dangerous and defective when put to their reasonably anticipated use.

101. As a direct and proximate result of the Johnson & Johnson Defendants' negligence. Plaintiffs purchased and used the PRODUCTS that directly and proximately caused each Plaintiff to develop ovarian cancer. As a direct and proximate result, Plaintiffs were caused to incur medical bills, lost wages, and conscious pain and suffering.

102. WHEREFORE, Plaintiffs demand judgment against the Johnson & Johnson Defendants for compensatory, treble damages pursuant to California Civil Code Section 3345, and punitive damages in excess of the jurisdictional minimum of this Court, together with interest, costs of suit, attorneys' fees, and all such other relief, as this Court deems proper.

COUNT VII - BREACH OF EXPRESS WARRANTY

(Against Johnson & Johnson Defendants)

103. Plaintiffs incorporate by reference all other paragraphs in this Complaint as if set forth fully herein.

104. At all relevant times, the Johnson & Johnson Defendants knew or should have known that the PRODUCTS were unreasonably dangerous and defective when put to their reasonably anticipated use.

105. At all relevant times, the Johnson & Johnson Defendants expressly warranted, through direct-to-consumer marketing, advertisements, and labels, that the PRODUCTS were safe and effective for reasonably anticipated uses, including use by women in their perineal area. Although the label has changed over time, the message has been the same: that the product is

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safe for use on women as well as babies. At least as of 2014, the baby powder label stated that "Johnson's Baby Powder is designed to gently absorb excess moisture helping skin feel comfortable. Our incredibly soft, hypoallergenic, dermatologist and allergy-tested formula glides over skin to leave it feeling delicately soft and dry while providing soothing relief." The Johnson & Johnson Defendants instruct consumers on the product labeling to "Shake powder directly into your hand, away from the face, before smoothing onto the skin."

Defendants similarly encouraged women to use the product daily. Defendants state that Johnson's Baby Powder "keeps skin feeling soft, fresh and comfortable. It's a classic Johnson's Baby Powder helps eliminate friction while keeping skin cool and comfortable. It's made of millions of tiny slippery plates that glide over each other to help reduce the irritation caused by friction." Under a heading "How to Use." "For skin that feels soft, fresh and comfortable, apply Johnson's Baby Powder close to the body, away from the face. Shake powder into your hand and smooth onto skin." Under a heading "When to Use", the Johnson & Johnson Defendants recommend that the consumer "Use anytime you want skin to feel soft, fresh and comfortable. For baby, use after every bath and diaper change." On their website for Johnson's Baby Powder. Defendants also state the product is "Clinically proven to be safe, gentle and mild."

107. Even more recently, in February or March, 2016, after a St. Louis Jury rendered a \$72 million dollar verdict against Johnson & Johnson, including punitive damages, Johnson &

Johnson published a web page directed at consumers misleadingly assuring them of the safety of tale titled "Our Safety & Care Commitment" and touted the safety of tale, stating, inter alia:

- a. "Decades of Safety: Our confidence in using tale reflects more than 30 years of research by independent scientists, review boards and global authorities, which have concluded that tale can be used safely in personal care products. Various government agencies and other bodies also have examined tale to determine the potential for any safety risks, and none have concluded that there are safety risks. In fact, no regulatory agency has ever required a change in labeling to reflect any safety risk from tale powder products."
- b. "Our Position on Tale: At Johnson & Johnson Consumer Inc., our confidence in using tale is based on a long history of safe use and more than 30 years of research by independent researchers, scientific review boards and global regulatory authorities. Various agencies and governmental bodies have examined whether tale is a carcinogen, and none have concluded that it is. With over 100 years of use, few ingredients have the same demonstrated performance, mildness and safety profile as cosmetic tale."
- c. "We want to assure women and caregivers who use our tale products that numerous studies support its safety, and these include assessments by external experts in addition to our company testing. Many research papers and epidemiology studies have specifically evaluated tale and perineal use and these studies have found tale to be safe"
- 108. At all relevant times, even up until present day, the Johnson & Johnson Defendant's representations relating to tale: that the PRODUCTS are safe for personal use, including in the perineal region.
- 109. At all relevant times, the PRODUCTS did not conform to these express representations because they cause serious injury, including ovarian cancer, when used by women in the perincal area.

¹⁴ See, http://www.safetyundcarecommitment.com/ingrodient-info/other/tale

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- 10. As a direct and proximate result of the Defendants' breach of warranty, Plaintiffs purchased and used the PRODUCTS that directly and proximately caused each Plaintiff to develop ovarian cancer. Plaintiffs were caused to incur medical bills, lost wages, and conscious pain and suffering.
- 111. WHEREFORE, Plaintiffs demand judgment against the Johnson & Johnson Defendants for compensatory, treble damages pursuant to California Civil Code Section 3345, and punitive damages in excess of the jurisdictional minimum of this Court, together with interest, costs of suit, attorneys' fees, and all such other relief, as this Court deems proper.

COUNT VIII - BREACH OF IMPLIED WARRANTIES

(Against Johnson & Johnson Defendants)

- 112. Plaintiff's incorporate by reference all other paragraphs in this Complaint as if set forth fully herein.
- 113. At the time the Defendants manufactured, marketed, labeled, promoted, distributed and/or sold the PRODUCTS, the Johnson & Johnson Defendants knew of the uses for which the PRODUCTS were intended, including use by women in the perineal area. With this knowledge, they impliedly warranted the PRODUCTS to be of merchantable quality and safe for such use.
- 114. Defendants breached their implied warranties of the PRODUCTS sold to Plaintiffs because they were not fit for their common, ordinary and intended uses, including use by women in the perineal area.
- 115. As a direct and proximate result of the Johnson & Johnson Defendants' breach of implied warranties, Plaintiff's purchased and used the PRODUCTS that directly and proximately caused

each Plaintiff to develop ovarian cancer. As a result, Plaintiffs were caused to incur medical bills, lost wages, and conscious pain and suffering.

116. WHEREFORE, Plaintiffs demand judgment against the Johnson & Johnson Defendants for compensatory, treble damages pursuant to California Civil Code Section 3345, and punitive damages in excess of the jurisdictional minimum of this Court, together with interest, costs of suit, attorneys' fees, and all such other relief, as this Court deems proper.

COUNT IX - CIVIL CONSPIRACY

(Against All Defendants)

- 117. Plaintiffs incorporate by reference all other paragraphs in this Complaint as if set forth fully herein.
- 118. Defendants and/or their predecessors-in-interest knowingly agreed, contrived, combined, confederated and conspired among themselves to cause Plaintiffs' injuries, diseases, and/or illnesses by exposing the Plaintiffs to harmful and dangerous PRODUCTS. Defendants further knowingly agreed, contrived, confederated and conspired to deprive the Plaintiffs of the opportunity of informed free choice as to whether to use the PRODUCTS or to expose themselves to the stated dangers. Defendants committed the wrongs as described herein by willfully misrepresenting and suppressing the truth as to the risks and dangers associated with the use of and exposure to the PRODUCTS.
- 119. In furtherance of said conspiracies, Defendants performed the following overt acts:
 - a. For many decades, Defendants, individually, jointly, and in conspiracy with each other, have been in possession of medical and scientific data, literature and test reports that clearly indicated that use of their by women resulting from ordinary and

b. Despite the medical and scientific data, literature, and test reports possessed by and available to Defendants, Defendants individually, jointly, and in conspiracy with each other, fraudulently, willfully and maliciously:

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- i. Withheld, concealed and suppressed said medical information regarding the increased risk of ovarian cancer from Plaintiffs, as described above; in addition, on July 27, 2005, Defendants, as part of the TIPTF, corresponded about and agreed to edit and delete portions of scientific papers being submitted on their behalf to the United States Toxicology Program in an attempt to prevent tale from being classified as a carcinogen;
- ii. Instituted a "defense strategy" through the TIPTF to defend tale at all costs. In furtherance of this defense strategy, Defendants, through the TIPTF, used their influence over the National Toxicology Program ("NTP") Subcommittee and the threat of litigation against the NTP to prevent the NTP from classifying tale as a careinogen on its 10th Report on Carcinogens ("RoC");
- iii. Caused to be released, published and disseminated medical and scientific data, literature, and test reports containing information and statements regarding the risks of ovarian cancer which Defendants knew were incorrect incomplete, outdated, and misleading. Specifically, Defendants, through the TIPTF, collectively agreed to release false information to the public regarding the safety of talc on July 1, 1992; July 8, 1992; and November 17, 1994. In a letter dated September 17, 1997, the Defendants were criticized by their own toxicologist consultant for releasing this false information to the public, yet nothing was done by the Defendants to correct or redact this public release of knowingly false information.
- c. By these false and fraudulent representations, omissions, and concealments. Defendants intended to induce and did induce the Plaintiffs to rely upon these false and fraudulent representations, omissions and concealments, and to continue to expose themselves to the dangers inherent in the use of and exposure to the PRODUCTS.
- 120. Plaintiffs reasonably and in good faith relied upon the fraudulent representations, omissions, and concealments made by Defendants regarding the nature of the PRODUCTS.

purchased and used the PRODUCTS in the perineal areas, which directly and proxima	iel;
caused each Plaintiff to develop ovarian cancer. Plaintiffs were caused to incur medical bills,	los
wages, and conscious pain and suffering.	

122. WHEREFORE, Plaintiffs demand judgment against all Defendants for compensatory treble damages pursuant to California Civil Code Section 3345, and punitive damages in excess of the jurisdictional minimum of this Court, together with interest, costs of suit, attorneys' fees, and all such other relief, as this Court deems proper.

COUNT X - FRAUD, FRAUDULENT MISREPRESENTATION, AND INTENTIONAL CONCEALMENT

(Against Johnson & Johnson Defendants)

- 123. Plaintiffs incorporate by reference all other paragraphs in this Complaint as if set forthfully herein.
- 124. At all relevant times, the Johnson & Johnson Defendants intentionally, willfully, and/or recklessly, with the intent to deceive, misrepresented and/or concealed material facts to consumers and users, including Plaintiffs.
- 125. At all relevant times, the Johnson & Johnson Defendants misrepresented and/or concealed material facts concerning the PRODUCTS to consumers, including the Plaintiffs, with knowledge of the falsity of their misrepresentations.

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27 28 126. At all relevant times, upon information and belief, the misrepresentations and concealments concerning the PRODUCTS made by the Johnson & Johnson Defendants include but are not limited to the following:

- a. The Johnson & Johnson Defendants falsely labeled and advertised the PRODUCTS in the following ways, among others: "For you, use every day to help feel soft, fresh, and comfortable," "a sprinkle a day keeps the odor away," "your body perspires in more places than just under your arms," "Use SHOWER to SHOWER to feel dry, fresh, and comfortable throughout the day," and "SHOWER to SHOWER can be used all over your body."
- b. The Johnson & Johnson Defendants falsely advertised the PRODUCT SHOWER to SHOWER to be applied "all over," and in particular, urges women to use it to "Soothe Your Skin: Sprinkle on problem areas to soothe skin that has been irritated from friction. Apply after a bikini wax to help reduce irritation and discomfort."
- c. The Johnson & Johnson Defendants, through the advertisements described above, knowingly misrepresented to Plaintiff and the public that the PRODUCTS were safe for use all over the body, including the perineal areas of women.
- d. The Johnson & Johnson Defendants intentionally failed to disclose that tale and the associated PRODUCTS, when used in the perineal area, increase the risk of ovarian cancer.
- e. The Johnson & Johnson Defendants intentionally failed to include adequate warnings with the PRODUCTS regarding the potential and actual risks of using the PRODUCTS in the perineal area on women and the nature, scope, severity, and duration of any serious injuries resulting therefrom. ¹⁵
- Despite knowing about the carcinogenic nature of tale and its likelihood to increase the risk of ovarian cancer in women, the Johnson & Johnson Defendants falsely marketed, advertised, labeled and sold the PRODUCTS as safe for public consumption and usage, including for use by women to powder their perineal areas.

¹⁵ Household PRODUCTS Database, Label for Johnson's Baby Powder, Original, http://household/products.nlm.nih.gov/egi-bin/household/brands/tbl=brands&id=10001040

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127. At all relevant times, the Johnson & Johnson Defendants actively, knowingly, and intentionally concealed and misrepresented these material facts to the consuming public with the intent to deceive the public and Plaintiffs, and with the intent that the consumers would purchase and use the PRODUCTS in the female perineal area.

128. At all relevant times, the consuming public, including Plaintiffs, would not otherwise have purchased the PRODUCTS and/or applied the PRODUCTS in the perineal area if they had been informed of the risks associated with the use of the PRODUCTS in the perineal area.

129. At all relevant times, Plaintiffs relied on the Johnson & Johnson Defendants' misrepresentations concerning the safety of the PRODUCTS when purchasing the PRODUCTS and using them in her perineal area, and her reliance was reasonable and justified.

130. As a direct, foreseeable and proximate result of the Johnson & Johnson Defendants' fraudulent conduct, Plaintiffs purchased and used the PRODUCTS in their perineal areas. As a direct and proximate result of such use, each Plaintiff developed ovarian cancer, and Plaintiffs were caused to incur medical bills, lost wages, and conscious pain and suffering.

131. WHEREFORE, Plaintiffs demand judgment against the Johnson & Johnson Defendants for compensatory, treble damages pursuant to California Civil Code Section 3345, and punitive damages in excess of the jurisdictional minimum of this Court, together with interest, costs of suit, attorneys' fees, and all such other relief, as this Court deems proper.

COUNT XI - NEGLIGENT MISREPRESENTATION

(Against All Defendants)

132. Plaintiffs incorporate by reference all other paragraphs in this Complaint as if set forth fully herein.

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133. As a direct, foresceable and proximate result of the Johnson & Johnson Defendants' fraudulent conduct, Plaintiffs purchased and used the PRODUCTS in their perineal areas. As a direct and proximate result of such use, each Plaintiff developed ovarian cancer, and Plaintiffs were caused to incur medical bills, lost wages, and conscious pain and suffering.

- 134. Defendants had a duty to accurately and truthfully represent to the medical and healthcare community, Plaintiffs and the public that the PRODUCTS had been tested and found to be safe and effective for use in the perineal area. However, the representations made by Defendants, in fact, were false.
- 135. 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' high risk of unreasonable, dangerous, adverse side effects.
- 136. Defendants breached their duty in representing that the PRODUCTS were safe for use in the perineal areas of women.
- 137. At all relevant times, upon information and belief, the misrepresentations, omissions and concealments concerning the PRODUCTS made by the Defendants include, but are not limited to the following:
 - a. The Johnson & Johnson Defendants labeled and advertised the PRODUCTS in the following ways, among others: "For you, use every day to help feel soft, fresh, and comfortable;" "A sprinkle a day keeps the odor away;" "Your body perspires in more places than just under your arms;" "Use SHOWER to SHOWER to feel dry, fresh, and comfortable throughout the day; and "SHOWER to SHOWER can be used all over your body."
 - b. The Johnson & Johnson Defendants advertised the product SHOWER to SHOWER to be applied "all over," and in particular, urged women to use it to "Soothe Your

- c. Defendants, through the advertisements described above, among others, misrepresented to consumers, including the Plaintiffs, that the PRODUCTS were safe for use all over the body, including the female perineal area.
- d. Despite actual knowledge of the health risks of the PRODUCTS, the Defendants failed to disclose to the consumers and the Plaintiffs, through adequate warnings, representations, labeling, or otherwise, that the PRODUCTS were inherently dangerous and carcinogenic in nature, which poses serious health risks to consumers.
- e. Despite actual knowledge that the use of the PRODUCTS in the perineal area created a significantly increased risk of ovarian cancer, the Defendants failed to disclose to consumers and the Plaintiff, through adequate warnings, representations, labeling, or otherwise, that material fact.

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- f. Despite knowing about the carcinogenic nature of tale and its likelihood to increase the risk of ovarian cancer in women, the Johnson & Johnson Defendants falsely marketed, advertised, labeled and sold the PRODUCTS as safe for public consumption and usage, including for use by women to powder their perineal areas.
- 138. At all relevant times, Defendants failed to exercise reasonable care in ascertaining or sharing information regarding the safe use of PRODUCTS, failed to disclose facts indicating that the PRODUCTS were inherently dangerous and carcinogenic in nature, and otherwise failed to exercise reasonable care in communicating the information concerning the PRODUCTS to Plaintiff and/or concealed relevant facts that were known to them.
- 139. At all relevant times, Plaintiffs were not aware of the falsity of the foregoing misrepresentations, nor was she aware that material facts concerning tale and the PRODUCTS had been concealed or omitted. In reasonable reliance upon the Johnson & Johnson Defendants' misrepresentations and/or omissions, Plaintiffs were induced to and did purchase the PRODUCTS and did use the PRODUCTS on her perineal area. If the Defendants had disclosed true and accurate material facts concerning the risks of the use of the PRODUCTS, in particular

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 the risk of developing ovarian cancer from using the PRODUCTS in the female perincal area, Plaintiffs would not have purchased and/or received the PRODUCTS and/or used the PRODUCTS in that manner.

140. Plaintiffs' reliance upon the Defendants' misrepresentations and omissions was justified and reasonable because, among other reasons, those misrepresentations and omissions were made by individuals and entities who were in a position to know the material facts concerning the PRODUCTS and the association between the PRODUCTS and the incidence of ovarian cancer, while Plaintiff was not in a position to know these material facts, and because the Johnson & Johnson Defendants failed to warn or otherwise provide notice to the consuming public as to the risks of the PRODUCTS, thereby inducing Plaintiff to use the PRODUCTS in lieu of safer alternatives and in ways that created unreasonably dangerous risks to her health. At all relevant times, the Defendants' corporate officers, directors, and/or managing agents knew of and ratified the acts of the Johnson & Johnson Defendants, as alleged herein.

- 141. As a direct and proximate result of Defendants' conduct, Plaintiffs have been injured and sustained severe and permanent pain, suffering, disability, impairment, loss of enjoyment of life, loss of care and comfort, and economic damages.
- 142. WHEREFORE, Plaintiffs demand judgment against all Defendants for compensatory treble damages pursuant to California Civil Code Section 3345, and punitive damages in excess of the jurisdictional minimum of this Court, together with interest, costs of suit, attorneys' fees, and all such other relief, as this Court deems proper.

COUNT XII - WRONGFUL DEATH

(Against All Defendants)

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143. Plaintiffs hereby incorporate by reference all other paragraphs in this Complaint as if set forth fully herein.

144. As a direct and proximate result of the acts and/or omissions of Defendants as set forth herein, the Decedents named in this action used the PRODUCTS in their perincal areas. Subsequent to such use, Decedents developed ovarian cancer, suffered substantial pain and suffering, both physical and emotional in nature, and subsequently died.

145. Plaintiffs, on behalf of themselves and all of the next of kin or successors-in-interest of Decedents, are entitled to recover damages as Decedents would have if they were living, as a result of acts and/or omissions of Defendants.

146. Plaintiffs, on behalf of themselves and all of Decedents' next of kin or successors-ininterest are also entitled to recover punitive damages and damages for substantial pain and suffering caused to Decedents from the acts and/or omissions of Defendants as fully set forth herein, including without limitations, punitive damages.

147. As a direct and proximate result of Defendants' conduct, Plaintiffs and Decedents have been injured and sustained severe and permanent pain, suffering, disability, impairment, loss of enjoyment of life, loss of care and comfort, and economic damages.

148. WHEREFORE, Plaintiffs demand judgment against all Defendants for compensatory treble damages pursuant to California Civil Code Section 3345, and punitive damages in excess of the jurisdictional minimum of this Court, together with interest, costs of suit, attorneys' fees, and all such other relief, as this Court deems proper.

COUNT XIII - PUNITIVE DAMAGES

149. Plaintiffs hereby incorporate by reference all other paragraphs in this Complaint as if set forth fully herein.

150. California Code of Civil Procedure Section 3294 provides that "In an action for the breach of an obligation not arising from contract, where it is proven by clear and convincing evidence that the defendant has been guilty of oppression, fraud, or malice, the plaintiff, in addition to the actual damages, may recover damages for the sake of example and by way of punishing the defendant."

151. The Defendants have acted with oppression, fraud, and/or malice in the following ways in addition to the acts and/or omissions described throughout this Complaint:

a. Defendants knew of the unreasonably high risk of ovarian cancer posed by the PRODUCTS before manufacturing, marketing, distributing and/or selling the PRODUCTS, yet purposefully proceeded with such action;

b. Despite their knowledge of the high risk of ovarian cancer associated with the PRODUCTS. Defendants affirmatively minimized this risk through marketing and promotional efforts and product labeling:

c. Through the actions outlined above, Defendants expressed a reckless indifference to the safety of users of the PRODUCTS, including Plaintiffs. Defendants knew of the dangers and risks of the PRODUCTS, yet they concealed and/or omitted this information from labels and warnings contained on the PRODUCTS in furtherance of their conspiracy and concerted action. These actions were outrageous because of Defendants evil motive or a reckless indifference to the safety of users of the PRODUCTS.

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154. forth fully herein. counsel, and consortium.

As a direct and proximate result of the Defendants' acts of oppression, fraud and/or malice described throughout this Complaint, Plaintiffs have sustained damages as set forth above

WHEREFORE, Plaintiffs demand judgment against all Defendants for compensatory, treble damages pursuant to California Civil Code Section 3345, and punitive damages in excess of the jurisdictional minimum of this Court, together with interest, costs of suit, attorneys' fees, and all such other relief, as this Court deems proper.

COUNT XIV - LOSS OF CONSORTIUM

(Against All Defendants)

- Plaintiffs hereby incorporate by reference all other paragraphs in this Complaint as if set
- 155. Plaintiff's and Decedents' spouses were entitled to the comfort, care, affection companionship, services, society, advice, guidance, counsel, and consortium of their spouses.
- As a direct and proximate result of one or more of those wrongful acts or omissions of the Defendants described herein, Plaintiffs and Decedents' spouses have been and will be deprived of the comfort, care, affection, companionship, services, society, advice, guidance
- 157. WHEREFORE, Plaintiffs demand judgment against all Defendants for compensatory treble damages pursuant to California Civil Code Section 3345, and punitive damages, together with interest, costs of suit, attorneys' fees, and all such other relief as this Court deems proper.

TOLLING STATUTE OF LIMITATIONS

158. Plaintiffs hereby incorporate by reference all other paragraphs in this Complaint as if see forth fully herein,

 159. Plaintiffs have suffered an illness that has a latency period and does not arise until many years after exposure. Plaintiffs' illnesses did not distinctly manifest themselves until they were made aware that their ovarian cancer could be caused by their use of the Defendants' products. Consequently, the discovery rule applies to these cases, and the statute of limitations has been tolled until the day that Plaintiff's knew or had reason to know that their (or decedents') ovarian cancer was linked to their (or decedents') use of the Defendants' products.

160. Furthermore, the running of any statute of limitations has been equitably tolled by reason of Defendants' fraudulent concealment and conduct. Through their affirmative misrepresentations and omissions, Defendants actively concealed from Plaintiffs, decedents, and consumers the true risks associated with PRODUCTS.

161. As a result of Defendants' actions, Plaintiffs, decedents, and consumers were unaware, and could not reasonably know or have learned through reasonable diligence, that Plaintiffs and decedents had been exposed to the risks alleged herein and that those risks were the direct and proximate result of Defendants' acts and omissions.

162. Furthermore, Defendants are estopped from relying on any statute of limitations because of their concealment of the truth regarding the safety of PRODUCTS. Defendants were under a duty to disclose the true character, quality and nature of PRODUCTS because this was non-public information over which they continue to have exclusive control. Defendants knew that this information was not available to Plaintiffs, their medical providers and/or their health facilities, yet they failed to disclose the information to the public.

163. Defendants had the ability to and did spend enormous amounts of money in furtherance of their purposes of marketing and promoting a profitable product, notwithstanding the known or

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