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UNITED STATES D				
SOUTHERN DISTRIC	T OF CALIFORNIA			
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BENJAMIN A. VERKEST and LORI L. MCKEE-CALLANAN,	Case No. 14CV0106 JM JMA			
Plaintiffs,	G015DV 4.72VM			
VS.	COMPLAINT			
IANSSEN PHARMACEUTICALS INC				
JANSSEN PHARMACEUTICALS, INC. also known as ORTHO-NCNEIL-	JURY TRIAL DEMANDED			
JANSSEN PHARMACEUTICALS, INC.; JANSSEN PHARMACEUTICA,				
INC.; JANSSEN PHARMACEUTICA, INC.; JANSSEN LP; JOHNSON & JOHNSON, INC., JOHNSON &				
JOHNSON PHARMACEUTICAL				
RESEARCH AND DEVELOPMENT, LLC, and "JOHN DOE" 1-5 (said names				
being fictitious, as the true names are				
presently unknown), in their individual and official capacities,				
Defendants.				
PRELIMINARY	<u>STATEMENT</u>			
1. Plaintiff Benjamin A. Verkest	and his mother, Plaintiff Lori L. McKee			
Callanan ("Plaintiffs"), by and through their undersigned counsel, bring this action				
for strict liability, negligence, negligence	e per se, false advertising, fraudulent			
concealment, fraudulent misrepresentation	, failure to warn, breach of express and			

implied warranties, unfair business practices, negligent infliction of emotional

distress, intentional infliction of emotional distress, and reckless endangerment.

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This action is based upon Defendants' violations of the laws of the Unites States and of the State of California, including, but not limited to, 21 U.S.C. § 321, et seq. (the Federal Food, Drug, and Cosmetic Act), 31 U.S.C. §§ 3729-33; 3730(b)(1) (the False Claims Act), the Code of Federal Regulations (off-label promotion), 42 U.S.C. 1320a7b(b) (the federal anti-kickback statute), Section 13(b)(2)(A)(B) of the Securities and Exchange Act (15 U.S.C. § 78m(b)(2)(A)(B)), 21 U.S.C. §§ 331(a), 333(a)(1) and 352(f)(1)) (introduction of misbranded drugs into interstate commerce), California Business and Professions Code § 17200, et seq. and § 17500, and other applicable federal and California state requirements, in the manufacture, marketing and distribution of a defective and unreasonably dangerous medication, to wit, Risperdal, which is also known by the generic name risperidone (hereinafter "Risperdal" or "Risperdal/risperidone"). Plaintiffs seek compensatory, equitable, injunctive, punitive, and declaratory relief for the debilitating physical, psychological, pecuniary and related injuries for which Defendants are liable. Based upon personal knowledge and upon the investigation of their counsel, Plaintiffs respectfully allege the following:

- 2. This Court has subject matter jurisdiction pursuant to 28 U.S.C. §1332 because there is complete diversity of citizenship between Plaintiffs and each Defendant and because the amount in controversy exceeds \$75,000.00. This Court has personal jurisdiction over each Defendant pursuant to federal law and Cal. Code Civ. Proc. §410.10 due to Defendants' substantial, continuous and systematic presence and activity in California and due to Defendants' purposeful availment of the laws and privileges associated therewith.
- 3. Venue is properly laid in the United States District Court for the Southern District of California pursuant to 28 U.S.C. §1391(a) because the events giving rise to the claims alleged herein substantially occurred within the geographical boundaries of the District.

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This action arises in connection with Defendants' unlawful marketing 4 and promotion of the defective and unreasonably dangerous prescription medication Risperdal, which was taken by Plaintiff Verkest from 1997 to 2001. As a proximate result of Defendants' unlawful actions, Plaintiff Verkest suffered serious and debilitating physical, psychological, pecuniary and related injuries, including permanent disfigurement, significant and severe weight gain, diabetes mellitus, hyperglycemia, hyperprolactinemia, enuresis, damage to his sexual and endocrine functions, gynecomastia, impaired motor skills, dyssomnia, anxiety, embarrassment, difficulty concentrating, agitation, and impaired thinking, medical expenses, lost and/or diminished earning capacity and psychological trauma. Furthermore, as a proximate result of Defendants' unlawful actions, Plaintiff McKee-Callanan suffered various damages including, but not limited to, severe emotional distress, lost wages, inconvenience, medical expenses, and other damages. Plaintiffs seek, *inter alia*, compensatory, equitable, injunctive, punitive and declaratory relief for their injuries.

#### **BACKGROUND**

- 5. Plaintiff Benjamin A. Verkest is a 22 year old male, and a resident and citizen of the City of San Marcos, County of San Diego, and State of California. Plaintiff Lori L. McKee-Callanan is the biological mother of Plaintiff Verkest, and was at all times relevant hereto Plaintiff's Verkest's natural and legal parent and guardian.
- 6. At all times relevant to the acts alleged herein, Plaintiffs resided within the geographic confines of the Southern District of California.
- 7. Plaintiff Verkest resides with his mother Plaintiff McKee-Callanan, and additional family members.
- 8. In or about 1996, at about five years of age, Plaintiff Verkest began experiencing behavioral and psychological issues.

- 9. In or about November of 1997, at about six years of age, Plaintiff Verkest was prescribed and began to use Risperdal; Plaintiff continued to use Risperdal until approximately April of 2001, roughly one month before his tenth birthday.
- 10. During the time period that Plaintiff Verkest took Risperdal from 1997 to 2001, he experienced numerous serious side effects, including significant weight gain, enlarged nipples, and development of enlarged breasts.
- 11. Upon discontinuing Risperdal in or about April of 2001, Plaintiff Verkest experienced numerous deleterious effects, including increased aggression, acting out, and other emotional and behavioral problems.
- 12. Plaintiff Verkest has continued to experience negative effects resulting from his use of Risperdal from 2001 to the present, including gynecomastia, delayed onset of puberty, delayed and/or incomplete sexual development, impaired motor skills, dyssomnia, eneuresis, diabetes and/or diabetes-related symptoms, and other physical health issues and emotional disturbances.
- 13. Defendant Johnson & Johnson, Inc. ("J&J") is a New Jersey corporation with its principal place of business in New Jersey. Defendant J&J manufactures, markets, and sells a wide range of pharmaceutical, medical and related products. J&J is qualified to do business in California and does business in California.
- 14. Defendant Ortho-McNeil-Janssen Pharmaceuticals, Inc., also known as Janssen Pharmaceutica, Inc., and/or Janssen, LP, ("OMJPI") is a Pennsylvania corporation with its principal place of business in New Jersey.
- 15. Defendant Johnson & Johnson Pharmaceutical Research and Development, LLC ("J&JPRD") is a New Jersey limited liability company, whose sole member is Centocor Research & Development, Inc., a Pennsylvania

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27 28 corporation with its principal place of business in Pennsylvania. Medtronic Puerto Rico Operations Co. is a wholly-owned subsidiary of Defendant Medtronic, Inc., existing by virtue of the laws of the Cayman Islands, with its principal place of business at Road 149, km 56.3, Box 6001, Villalba, PR.

Defendants "John Doe" 1-5 are directors, officers, managers, 16. employees, agents, contractors, subsidiaries and/or closely related entities of the named and/or their subsidiaries who, at all times relevant to the allegations herein, acted within the scope of their authority and on behalf of the other Defendants.

#### **FACTUAL ALLEGATIONS**

- At all times relevant hereto, Defendants owned a patent on the 17. prescription drug Risperdal, which was approved by the Federal Food and Drug Administration ("FDA") in or around 1993. Defendants did during such times manufacture, create, design, test, label, sterilize, distribute, supply, prescribe, market, sell, advertise, purport to warn, purport to consult, and otherwise distribute in interstate commerce and in the State of California the product known as Risperdal.
- In or about 1997 and thereafter, Defendants made false and 18. misleading statements about the safety, cost and effectiveness of Risperdal and improperly influenced doctors and officials to promote and prescribe the medication.
- In or about 1999, Defendants received a warning letter from the FDA 19. regarding marketing Risperdal in a manner that was misleading, false and lacking in fair balance.
- 20. In or about 2001, Defendants were required by the FDA to change the labeling of Risperdal to include a statement that the safety and effectiveness of the drug in children had not been established.
  - 21. Documents released in connection with settlements, judgments and

plea agreements reached with the U.S. Department of Justice and various state attorneys general reflect that Defendants have concealed and/or minimized Risperdal's side effects and exaggerated Risperdal's effectiveness.

- 22. The New York Times has reported that between 1993 and 2008, more than 1,200 children have suffered serious complications in connection with taking Risperdal/risperidone, including 31 deaths.
- 23. On or about April 13, 2011, U.S. District Judge Gladys Kessler entered a Final Judgment against Defendant Johnson & Johnson in Civil Action No. 11-0686 in the U.S. District Court for the District of Columbia; the Final Judgment determined that Defendant Johnson & Johnson was "liable for disgorgement of \$38,227,826, representing profits gained as a result of the conduct alleged in the Complaint" brought by the U.S. Securities and Exchange Commission which alleged improper influence and failure to maintain proper records and accounting procedures.
- 24. That at all times relevant herein, Risperdal was widely and falsely advertised and promoted by JPI/JLP/J&J/JOHN DOE Defendants as a safe and effective treatment for Schizophrenia and Bipolar Disorder, and was falsely promoted by Defendants as a safe and effective treatment for non-FDA approved uses, such as for depressive symptoms, PTSD and MDD, and that JPI/JLP/J&J/JOHN DOE Defendants minimized and/or covered up the risk posed to patients taking Risperdal as prescribed and for those taking Risperdal for non-FDA approved uses, such as for depressive symptoms, Major Depression and PTSD.
- 25. That at all times relevant hereto, JPI/JLP/J&J/JOHN DOE Defendants knew that the product Risperdal was defective and that Risperdal was likely to cause hyperprolactenemia, gynecomastia, diabetes, excessive weight gain, gastrointestinal problems, urinary incontinence, fecal incontinence and other

medical problems.

- 26. At all times relevant hereto, JPI/JLP/J&J/JOHN DOE Defendants knew that Risperdal was no more effective and considerably less safe than other antipsychotic medications, yet engaged in an ongoing pattern of false and misleading conduct designed to increase Risperdal's perceived therapeutic and monetary value over cheaper, safer and more effective products.
- 27. JPI/JLP/J&J/JOHN DOE Defendants failed to disclose to physicians, patients, or Plaintiffs, and those similarly situated, that Risperdal was likely to cause hyperprolactenemia, gynecomastia, diabetes, excessive weight gain, gastrointestinal problems, urinary incontinence, fecal incontinence and other medical problems.
- 28. Defendants continued to promote Risperdal as safe and effective despite patient reports of adverse events, FDA warnings regarding Risperdal's dangers, and FDA requests to modify the warning labels.
- 29. As a direct result of ingesting Risperdal/Risperidone, Plaintiff Verkest has suffered severe physical and emotional injuries, including, but not limited to, hyperprolactenemia, gynecomastia, diabetes, excessive weight gain, gastrointestinal problems, urinary incontinence, fecal incontinence and other medical problems, fear, apprehension, despair, suicidality, social anxiety and other emotional problems.
- 30. Defendants failed to provide sufficient warnings and instructions that would have put Plaintiffs or the general public on notice of the dangers and adverse effects associated with Risperdal/Risperidone, including, but not limited to, hyperprolactenemia, gynecomastia, diabetes, excessive weight gain, gastrointestinal problems, urinary incontinence, fecal incontinence and other medical problems.

- 31. Risperdal/Risperidone were defective as marketed due to inaccurate warnings, instructions, and labeling in light of Defendants' knowledge that the product was likely to cause hyperprolactenemia, gynecomastia, diabetes, excessive weight gain, gastrointestinal problems, urinary incontinence, fecal incontinence and other medical problems.
- 32. Defendants manufactured and promoted Risperdal/Risperidone for sale within the State of California and elsewhere.
- 33. Defendants promoted Risperdal to physicians and consumers within the State of California and elsewhere.
- 34. Defendants knew or should have known that their false advertising and unlawful marketing activities in violation of the Fair Claims Act and other federal and state laws was likely to and did in fact cause physicians and consumers to rely on said advertising and marketing and to take Risperdal/risperidone without adequate knowledge of the risks associated therewith.
- 35. Defendants conducted an organized, coordinated, intentional and deliberate campaign to unlawfully market and promote off-label use of Risperdal/risperidone in spite of the risks associated therewith.
- 36. As a result of Defendants' unlawful actions, Risperdal became Defendants' best-selling drug.
- 37. As a result of Defendants' unlawful actions, physicians and consumers were deceived into using Risperdal/risperidone in lieu of first generation antipsychotic medications ("FGAs" or other medications in spite of the fact that Risperdal carried additional dangerous side effects, was not approved for pediatric use until 2006, and cost approximately 40-50 times as much as FGAs which were equally or more effective.
- 38. Defendants sought to create the image, impression and belief among consumers and physicians that the use of Risperdal/risperidone was safe for

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humans, including children, and that it had fewer side effects and adverse reactions than other medications; Defendants engaged in this unlawful behavior despite knowing that their representations were false and that there was no reasonable basis to believe them to be true.

39. Defendants purposefully concealed, obfuscated, downplayed and understated the health hazards and risks associated with Risperdal and actively promoted its off-label pediatric use in violation of federal and California state law.

# CLAIMS FOR RELIEF FIRST CAUSE OF ACTION

#### **NEGLIGENCE**

- 40. Plaintiffs hereby incorporate by reference all preceding paragraphs as if fully set forth herein.
  - 41. Defendants owed Plaintiffs a legal duty of care.
- 42. Defendants knew or should have known that there was a foreseeable risk that Plaintiff would suffer harmful side effects from Risperdal/risperidone and the resulting damages alleged herein.
  - 43. Defendants failed to act reasonably or with ordinary prudence.
- 44. It was reasonable for Plaintiffs to rely on Defendants' representations as to the safety and effectiveness of Risperdal/risperidone and Plaintiffs did so rely.
- 45. But for Defendants' breach of duty owed to Plaintiffs, and Plaintiffs' detrimental reliance thereon, Plaintiffs would not have suffered the harm alleged herein.
- 46. As a direct and proximate result of Defendants' wrongful conduct, Plaintiffs have suffered and will continue to suffer severe physical injuries and/or death, severe emotional distress, mental anguish, economic losses and other damages for which they are entitled to compensatory, equitable and other lawfully available relief in an amount to be proven at trial.

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#### SECOND CAUSE OF ACTION

#### **BREACH OF EXPRESS WARRANTY**

- 47. Plaintiffs hereby incorporate by reference all preceding paragraphs as if fully set forth herein.
- 48. At all times mentioned herein, Defendants expressly warranted to Plaintiffs by and through statements made by Defendants or their authorized agents or sales representatives, orally and in publications, package inserts and other written materials intended for physicians, medical patients and the general public, that the aforementioned products were safe, effective, fit and proper for their intended use.
- 49. In utilizing the aforementioned products, Plaintiffs relied on the skill, judgment, representations and foregoing express warranties of the Defendants, and each of them. Said warranties and representations were false in that the aforementioned products were not safe and were unfit for the uses for which they were intended.
- 50. As a result of the foregoing breach of express warranties by the Defendants, Plaintiffs suffered injuries and damages as alleged herein.

### THIRD CAUSE OF ACTION

#### **BREACH OF IMPLIED WARRANTY**

- 51. Plaintiffs hereby incorporate by reference all preceding paragraphs as if fully set forth herein.
- 52. At all times mentioned herein, Defendants expressly warranted to Plaintiffs by and through statements made by Defendants or their authorized agents or sales representatives, orally and in publications, package inserts and other

written materials intended for physicians, medical patients and the general public, that the aforementioned products were safe, effective, fit and proper for their intended use.

- 53. In utilizing the aforementioned products, Plaintiffs relied on the skill, judgment, representations and foregoing express warranties of the Defendants, and each of them. Said warranties and representations were false in that the aforementioned products were not safe and were unfit for the uses for which they were intended.
- 54. As a result of the foregoing breach of express warranties by the Defendants, Plaintiffs suffered injuries and damages as alleged herein.

### FOURTH CAUSE OF ACTION VIOLATION OF CAL. BUS. CODE § 17500, et seq.

- 55. Plaintiffs hereby incorporate by reference all preceding paragraphs as if fully set forth herein.
- 56. At all times mentioned herein, Defendants expressly warranted to Plaintiffs by and through statements made by Defendants or their authorized agents or sales representatives, orally and in publications, package inserts and other written materials intended for physicians, medical patients and the general public, that the aforementioned products were safe, effective, fit and proper for their intended use.
- 57. In utilizing the aforementioned products, Plaintiffs relied on the skill, judgment, representations and foregoing express warranties of the Defendants, and each of them. Said warranties and representations were false in that the aforementioned products were not safe and were unfit for the uses for which they were intended.

58. As a result of the foregoing breach of express warranties by the Defendants, Plaintiffs suffered injuries and damages as alleged herein.

## FIFTH CAUSE OF ACTION FRAUDULENT CONCEALMENT

- 59. Plaintiffs hereby incorporate by reference all preceding paragraphs as if fully set forth herein.
- 60. Defendants and Plaintiffs were in a fiduciary relationship, wherein Defendants manufactured, supplier and actively promoted a dangerous and untested prescription drug to young children;
- 61. Defendants intentionally concealed that they had not conducted proper tests and did not know of the risks and side effects of Risperdal, thereby intentionally failed to disclose important facts to Plaintiffs;
- 62. Defendants were in the unique position to know that they did not have generally accepted test results of the effects of Risperdal on young children
- 63. Plaintiffs did not know that Defendants lacked generally accepted testing results about the potential risks and side effects of young children taking Risperdal.
  - 64. Defendants intended to deceive Plaintiffs by concealing these facts;
- 65. Plaintiffs reasonably relied on Defendants' assertions, as passed on by their doctors;
  - Plaintiff was harmed by Defendants' fraudulent concealment;
- 67. Defendants' concealment was a substantial factor in causing Plaintiffs' harm.

#### **SIXTH CAUSE OF ACTION**

#### STRICT PRODUCTS LIABILITY-FAILURE TO WARN

- 68. Plaintiffs hereby incorporate by reference all preceding paragraphs as if fully set forth herein.
- 69. Plaintiffs claim that Risperdal lacked sufficient warnings of potential risks and side effects on young children.
  - 70. Defendants manufactured and distributed Risperdal;
- 71. Risperdal had potential risks and side effects for young boys that were known or knowable in the light of scientific and/or medical knowledge that was generally accepted in the scientific and/or medical community at the relevant times when Defendants were manufacturing and distributing Risperdal to physicians for off-label use with children;
- 72. The potential risks and side effects presented a substantial danger when Risperdal is used or misused in an intended or reasonably foreseeable way;
- 73. The potential risks and side effects are not the type of risks and side effects that ordinary consumers would recognize;
- 74. The potential risks and side effects were ignored by Defendants when advising doctors of the benefits of Risperdal in young boys;
- 75. Due to the strict requirements established by the FDA for approving anti-psychotic prescription drugs for any specific use, particularly in young children more susceptible to adverse effects, Defendants knew that Risperdal was not approved for use in young children, and yet they pushed this off-label use anyway this is the risk-amelioration intended by compliance with the FDA regulations for approval of these types of drugs in all persons, especially children.
  - 76. Defendants knew at all times that Risperdal was not approved by the

FDA for use by children.

- 77. Defendants failed to adequately warn of the potential risks and side effects;
  - 78. Plaintiffs were harmed;
- 79. The lack of sufficient instructions and warnings were substantial factors in causing Plaintiffs' harm.

## SEVENTH CAUSE OF ACTION NEGLIGENCE-FAILURE TO WARN

- 80. Plaintiffs hereby incorporate by reference all preceding paragraphs as if fully set forth herein.
- 81. Plaintiffs claim that Defendants were negligent by not using reasonable care to warn about Risperdal's dangerous condition or about facts that made Risperdal likely to be dangerous.
- 82. Defendants manufactured and distributed Risperdal from 1993 to present day.
- 83. Defendants knew or reasonably should have known that Risperdal was dangerous or was likely to be dangerous when used or misused in a reasonably foreseeable manner;
- 84. Defendants knew or reasonably should have known that users would not realize the danger;
- 85. Defendants failed to adequately warn of the danger or instruct on the safe use of Risperdal;
- 86. A reasonable manufacturer and distributor under the same or similar circumstance would have warned of the danger or instructed on the safe use of

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an anti-psychotic prescription drug manufacturer and distributor owes to children, giving rise to gross negligence or recklessness.

### NINTH CAUSE OF ACTION NEGLIGENT MISREPRESENTATION

- 97. Plaintiffs hereby incorporate by reference all preceding paragraphs as if fully set forth herein.
- 98. Defendants represented to Plaintiffs, doctors and the wider medical community that Risperdal was safe for young children.
- 99. Defendants' representations were not true, as Defendants had no approved use from the FDA for the use of Risperdal in children, and either did not know of the harm because they failed to adequately test the drug in children, or knew of the risks and side effects but marketed
- 100. Regardless of whether Defendants honestly believed that the representations were true, Defendants had no reasonable grounds for believing the representations were true when they made the statements;
  - Defendants intended that Plaintiffs rely on the representations;
  - 102. Plaintiffs reasonably relied on Defendants' representations;
- 103. Plaintiffs' reliance on Defendants' representations was a substantial factor in causing their harm.'

### TENTH CAUSE OF ACTION FALSE ADVERTISING

- 104. Plaintiffs hereby incorporate by reference all preceding paragraphs as if fully set forth herein.
- 105. Defendants violated Cal. Bus. and Prof. Code s. 17500 et seq. by publicly make false and misleading statements in promotion and marketing

Risperdal to California doctors and patients, including Plaintiffs.

- 106. Defendants knew or should have known through the exercise of reasonable care under the circumstances that the aforementioned statements were false and misleading because Defendants had no way of knowing the whether or not such statements were true without conducting proper studies.
- 107. Defendants directly or indirectly disseminated false and misleading information as a marketing scheme to increase sales in the market of antipsychotic prescriptions to children.
  - 108. Defendants' false advertising caused the proximate harm to Plaintiffs.

### ELEVENTH CAUSE OF ACTION FRAUDULENT MISREPRESENTATION

- 109. Plaintiffs hereby incorporate by reference all preceding paragraphs as if fully set forth herein.
- 110. Defendants represented to Plaintiffs or their doctors that Risperdal was safe for young children;
- 111. Defendants knew at the time they made such representations that they were false because they had not conducted proper tests and did not know of the risks and side effects of Risperdal, and therefore could not know what the risks and side effects were.
- 112. Defendants intended for Plaintiffs to rely on their representations in order for their scheme to sell off-label Risperdal to children, including Plaintiffs.
- 113. Plaintiffs reasonably relied on Defendants' representations because Defendants are a famous producer of health products and prescription medication.
  - 114. Plaintiffs were harmed by ingesting Risperdal.

115. Plaintiffs' reliance on Defendants' representations was a substantial factor in causing their harm.

### TWELFTH CAUSE OF ACTION NEGLIGENT INFLICTION OF EMOTIONAL DISTRESS

- 116. Plaintiffs hereby incorporate by reference all preceding paragraphs as if fully set forth herein.
- 117. Defendants were negligent in their promotion and marketing of offlabel use of Risperdal as safe for children because Defendants did not know and could not know of the risks or side effects such use would cause Plaintiff Verkest because Defendants had not conducted proper testing of the medication as per FDA rules.
- Plaintiff Verkest suffered serious emotional distress caused by the offlabel use of Risperdal both during the time he ingested it and continuing to present day due to inter alia gynecomastia and the bullying and emotional distress of a young boy growing up with breasts.
- 119. Plaintiff Verkest has been constantly bullied, cannot conduct himself like most boys or men due to the physical deformation caused by the off-label use of Risperdal this has caused him shame, humiliation, physical bullying, emotional bullying anxiety and more. No person should have to endure this.
- 120. Defendants had a duty to provide safe prescription drugs to Plaintiff Verkest or proper and adequate warnings of risks and side effects Defendants were fiduciaries in the delivery of properly tested medication to young boys like Plaintiff Verkest.
- 121. Defendants' negligence was a substantial factor in causing Plaintiff Verkest's serious emotional distress.

- 122. Defendants negligently cause serious injury to Plaintiff Verkest by fraudulently promoting off-label use of Risperdal as safe for young boys.
- 123. During the time that Plaintiff Verkest ingested Risperdal, his behavior became serious destructive and at times violent toward himself and others, including Plaintiff McKee-Callanan.
- During the time that Plaintiff Verkest ingested Risperdal, he was publicly humiliated by many persons in front of his mother, Plaintiff McKee-Callanan.
- 125. Plaintiff McKee-Callanan personally witnessed the torment Risperdal caused her son, the violence it caused him to act upon towards Plaintiff McKee-Callanan, the bullying and ridicule by others towards Plaintiff Verkest due to the effects of taking Risperdal.
- 126. Plaintiff McKee-Callanan was then aware of the effects that Risperdal was causing injury to Plaintiff Verkest.
- 127. Plaintiff McKee-Callanan suffered serious emotional distress, including without limitation horror, anguish, fright, anxiety, grief, humiliation, shame, etc. as proximate result thereof;
- 128. Plaintiff McKee-Callanan's serious emotions distress was beyond that which would be anticipated in a disinterested witness.
- 129. Defendants' conduct was a substantial factor in causing Plaintiff McKee-Callanan's serious emotions distress.

# THIRTEENTH CAUSE OF ACTION INTENTIONAL INFLICTION OF EMOTIONAL DISTRESS

130. Plaintiffs hereby incorporate by reference all preceding paragraphs as if fully set forth herein.

- 131. Defendants' conduct of manufacturing, distributing, marketing or promoting the off-label use of Risperdal in children as safe without conducting proper, FDA-approved trials to understand what the risks and side effects were is outrageous conduct;
- 132. Defendants acted with reckless disregard of the probability that Plaintiff would suffer emotional distress, knowing that Plaintiff was a young boy and the risks and side effects of testing conducted on adults would likely be equal to or greater than those in adults; and in the causing of Plaintiffs to ingest Risperdal without knowing the risks and side effects;
- 133. Plaintiffs suffered severe emotional distress during the time he ingested the off-label use of Risperdal, and continuing to present day to the gynecomastia that haunts him personally and elicits taunting and bullying from others no person should have to endure this.
- 134. Defendants' conduct was a substantial factor in causing Plaintiffs' severe emotional distress.

1	PRAYER FOR RELIEF					
2	WHEREFORE, Plaintiffs request judgment against Defendants as follows:					
3	1. Economic and non-economic damages in an amount exceeding \$75,000 as					
4	provided by law and supp	provided by law and supported by the evidence at trial;				
5	2. Compensatory and Punitive damages;					
6	3. Attorneys' fees and costs;					
7	4. Prejudgment interests and costs; and					
8	5. Such other and further relief, including equitable relief, as the Court may					
9	deem just and proper.	deem just and proper.				
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11	D . 1 . 1 . 1 . 2011					
12	Dated: January 15, 2014	THE LAW OFFICE OF CRISTOPHER G. SABOL				
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14		/s/ Cristopher G. Sabol				
15		Cristopher G. Sabol Attorney for Plaintiffs				
16		Attorney for Flaminis				
17						
18		LAW OFFICE OF CRISTOPHER G. SABOL				
19		CRISTOPHER G. SABOL (SBN 251317)				
20		Email: sabolesq@gmail.com				
21		7985 Santa Monica Blvd. Suite 109-80 West Hollywood, CA 90046				
22		Tel: 323-383-1155				
23		Attorneys for Plaintiffs				
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The JS 44 civil cover sheet and the information contained herein neither replace nor supplement the filing and service of pleadings or often paper as required by law, except as

provided by local rules of court purpose of initiating the civil do	t. This form, approved by the ocket sheet. (SEE INSTRUCT	ne Judicial Conference of TIONS ON NEXT PAGE OF	f the United States in September THIS FORM.)	1974, is required for the use of	the Clerk of Court for the	
I. (a) PLAINTIFFS Benjamin Verkest and Lo	ri McKee-Callanan		Pharmaceuticals,	euticals, Inc. a/k/a Ortho- Inc; Janssen Pharmaceu		
(b) County of Residence of First Listed Plaintiff San Diego, CA  (EXCEPT IN U.S. PLAINTIFF CASES)			County of Residence of First Listed Defendant Mercer, NJ  (IN U.S. PLAINTIFF CASES ONLY)  NOTE: IN LAND CONDEMNATION CASES, USE THE LOCATION OF THE TRACT OF LAND INVOLVED.			
(c) Attorneys (Firm Name, 2 Cristopher G. Sabol, Esq 7985 Santa Monica Blvd. West Hollywood, CA 900	. (SBN 251317) , Ste. 10980	•)	Attorneys (If Known) unknown			
II. BASIS OF JURISDI	CTION (Place an "X" in O	ne Box Only)	III. CITIZENSHIP OF P	RINCIPAL PARTIES	(Place an "X" in One Box for Plaintij	
☐ 1 U.S. Government Plaintiff	☐ 3 Federal Question (U.S. Government)	Not a Party)		PTF DEF  1		
☐ 2 U.S. Government Defendant	■ 4 Diversity (Indicate Citizenshi)	p of Parties in Item III)	Citizen of Another State	☐ 2 Incorporated and F of Business In A		
			Citizen or Subject of a Foreign Country	3 3 Foreign Nation	□ 6 □ 6	
IV. NATURE OF SUIT		ly) RTS	FORFEITURE/PENALTY	BANKRUPTCY	OTHER STATUTES	
□ 110 Insurance □ 120 Marine □ 130 Miller Act □ 140 Negotiable Instrument □ 150 Recovery of Overpayment & Enforcement of Judgment □ 151 Medicare Act □ 152 Recovery of Defaulted Student Loans (Excludes Veterans) □ 153 Recovery of Overpayment of Veteran's Benefits □ 160 Stockholders' Suits □ 190 Other Contract □ 195 Contract Product Liability □ 196 Franchise    REAL PROPERTY □ 210 Land Condemnation □ 220 Foreclosure □ 230 Rent Lease & Ejectment □ 240 Torts to Land □ 245 Tort Product Liability □ 290 All Other Real Property	PERSONAL INJURY  □ 310 Airplane □ 315 Airplane Product Liability □ 320 Assault, Libel &	PERSONAL INJURY    365 Personal Injury -   Product Liability     367 Health Care/     Pharmaceutical     Personal Injury     Product Liability     368 Asbestos Personal     Injury Product Liability     368 Asbestos Personal     Injury Product Liability     370 Other Fraud     370 Other Fraud     371 Truth in Lending     380 Other Personal     Property Damage     385 Property Damage     70 Product Liability     PRISONER PETITIONS     463 Alien Detainee     510 Motions to Vacate     510 Motions to Vacate     530 General     535 Death Penalty     Other:     540 Mandamus & Other     550 Civil Rights     555 Prison Condition     560 Civil Detainee -   Conditions of     Confinement	☐ 625 Drug Related Seizure of Property 21 USC 881 ☐ 690 Other  LABOR  TY ☐ 710 Fair Labor Standards Act ☐ 720 Labor/Management Relations ☐ 740 Railway Labor Act ☐ 751 Family and Medical Leave Act ☐ 790 Other Labor Litigation S ☐ 791 Employee Retirement Income Security Act  IMMIGRATION ☐ 462 Naturalization Application	□ 422 Appeal 28 USC 158 □ 423 Withdrawal 28 USC 157  PROPERTY RIGHTS □ 820 Copyrights □ 830 Patent □ 840 Trademark  SOCIAL SECURITY □ 861 HIA (1395ff) □ 862 Black Lung (923) □ 863 DIWC/DIWW (405(g)) □ 864 SSID Title XVI □ 865 RSI (405(g))  FEDERAL TAX SUITS □ 870 Taxes (U.S. Plaintiff or Defendant) □ 871 IRS—Third Party 26 USC 7609	□ 375 False Claims Act □ 400 State Reapportionment □ 410 Antitrust □ 430 Banks and Banking □ 450 Commerce □ 460 Deportation □ 470 Racketeer Influenced and □ Corrupt Organizations □ 480 Consumer Credit □ 490 Cable/Sat TV □ 850 Securities/Commodities/ □ Exchange □ 890 Other Statutory Actions □ 891 Agricultural Acts □ 893 Environmental Matters □ 895 Freedom of Information □ Act □ 896 Arbitration □ 899 Administrative Procedure □ Act/Review or Appeal of □ Agency Decision □ 950 Constitutionality of □ State Statutes	
	moved from	Appellate Court tute under which you are	4 Reinstated or Reopened 5 Transf Reopened 5 Transf Anoth (specify) Filing (Do not cite jurisdictional state	er District Litigation		
VI. CAUSE OF ACTION	Brief description of ca	use:				
VII. REQUESTED IN COMPLAINT:	CHECK IF THIS UNDER RULE 2.	IS A CLASS ACTION	DEMAND \$	CHECK YES only <b>JURY DEMAND:</b>	if demanded in complaint:	
VIII. RELATED CASI IF ANY	E(S) (See instructions):	<sub>JUDGE</sub> n/a		DOCKET NUMBER		
DATE 01/15/2014		signature of atto /s/Cristopher Sa				
FOR OFFICE USE ONLY	AOUNT	ADDI VINC IED	HIDCE	MAC HI	DOE.	