

**UNITED STATES DISTRICT COURT
DISTRICT OF MINNESOTA**

CLIFFORD STRAKA,)	
)	
Plaintiff,)	
)	
v.)	Civil Action No. _____
)	
JOHNSON & JOHNSON, ORTHO-McNEIL)	
PHARMACEUTICAL, INC., and JOHNSON)	
& JOHNSON PHARMACEUTICAL)	
RESEARCH AND DEVELOPMENT, LLC,)	
)	
Defendants.)	

**COMPLAINT AND
DEMAND FOR JURY TRIAL**

Plaintiff Clifford Straka, by and through his attorneys of record, hereby files this Complaint and Demand for Jury Trial against Defendants Johnson & Johnson, Ortho-McNeil Pharmaceutical, Inc. ("Ortho-McNeil"), and Johnson & Johnson Pharmaceutical Research & Development, LLC ("Johnson & Johnson Pharmaceutical" of "Johnson & Johnson PRD") (collectively "Defendants"), and states of information and belief as follows:

INTRODUCTION

1. This case involves the fluoroquinolone antibiotic, levofloxacin.
2. Levofloxacin was designed, formulated, promoted, sold and distributed by Defendants in the United States as Levaquin from 1997 through the present.
3. Levaquin was approved by the FDA for treatment of a variety of serious infections. However, Defendants market Levaquin as a first line therapy for common

bronchitis and sinusitis infections, and for which many other, safer, antibiotics are available.

4. As compared to other fluoroquinolone antibiotic drugs, Levaquin causes a higher incidence of tendon injuries, including tendon rupture, especially in persons over 60 years of age and/or who are on corticosteroid therapy, none of which was adequately disclosed to Plaintiff and his physicians.

5. Levaquin-induced tendon injury involves the degradation of the tendon tissue, leading to severe and permanent injuries.

6. Plaintiff in the above-captioned case suffered a severe and debilitating tendon injury after his use of the drug Levaquin.

7. This lawsuit asserts claims against Defendants for strict product liability for manufacturing and/or design defect; strict product liability for failure to warn; negligence; breach of express and implied warranties for the design, manufacture, production, testing, study, inspection, labeling, marketing, advertising, sales, promotion, and distribution of Levaquin; fraud; violation of consumer protection laws; and unjust enrichment.

JURISDICTION

8. This Court has jurisdiction over this action pursuant to 28 U.S.C. § 1332, because the amount in controversy exceeds Seventy-Five Thousand Dollars (\$75,000.00), exclusive of interest and costs, and because there is complete diversity of citizenship between the Plaintiff and all Defendants.

9. Venue is proper in this judicial district pursuant to 28 U.S.C. § 1391 because the Defendants researched, designed, licensed, manufactured, tested, marketed, distributed, and/or sold the prescription drug Levaquin within this judicial district and because Defendants are subject to personal jurisdiction within the State of Minnesota.

PARTIES

10. Plaintiff Clifford Straka is a citizen and resident of Edina, Minnesota.

11. Defendant Johnson & Johnson is a New Jersey corporation with its principal place of business in New Brunswick, New Jersey.

12. Defendant Ortho-McNeil is a Delaware corporation with its principal place of business in Raritan, New Jersey. Defendant Ortho-McNeil is a wholly owned subsidiary of Johnson & Johnson.

13. Defendant Johnson & Johnson Pharmaceutical Research & Development is a New Jersey corporation with its principal place of business in Raritan, New Jersey. Defendant Johnson & Johnson Pharmaceutical Research & Development is a wholly owned subsidiary of Johnson & Johnson and was formerly known as R.W. Johnson Pharmaceutical Research Institute.

14. At all times relevant herein, Defendants tested, studied, researched, designed, formulated, manufactured, inspected, labeled, packaged, promoted, advertised, marketed, distributed, and sold the prescription drug Levaquin in interstate commerce and throughout the State of Minnesota. At all times relevant herein, Defendants were registered to do business in the State of Minnesota.

GENERAL FACTUAL ALLEGATIONS

15. Levaquin, Defendants' brand name for the antibiotic levofloxacin, is a broad spectrum synthetic antibacterial agent approved for use in the treatment of a variety of upper respiratory infections, urinary tract infections, prostatitis, and other bacterial infections. It was first introduced into the U.S. market in 1997.

16. Levaquin is in a class of antibiotics known as fluoroquinolones. The original quinolone antibiotics were developed in the early 1960s and soon revealed themselves as highly effective against common gram-negative bacteria, but resistance developed rapidly. Twenty years later, in the early 1980s, fluorinated derivatives of the quinolones emerged, revealing a broader, more potent antibiotic, effective against many different types of infections. These so-called second generation fluoroquinolones included norfloxacin (Noroxin), ciprofloxacin (Cipro), ofloxacin (Floxin), and pefloxacin (never marketed in the U.S.).

17. Although considered highly effective at killing certain bacteria, fluoroquinolones have long been associated with serious side effects. Indeed, many fluoroquinolones have been removed from the market due to intolerable adverse events. For example, Omniflox (temafloxacin) was removed from the market in 1992 because of low blood sugar, kidney failure, and a certain rare form of anemia; Raxar and Zagam were removed because of QT-interval prolongation among other things; Trovan was removed from the market due to severe liver toxicity; and most recently, Tequin was removed from the market in 2006 amid reports of severe blood sugar reactions such as hyperglycemia and hypoglycemia.

18. In sum, though fluoroquinolones may share certain pharmacological properties, their safety profiles can differ immensely.

A. OFLOXACIN – THE FIRST GENERATION OF LEVAQUIN

19. To understand the pharmacological properties of Levaquin, one need look no further than to Levaquin's older brother, ofloxacin (Floxin), also manufactured and distributed by Defendants.

20. Both Floxin and Levaquin were created and developed by Daiichi, Japanese Company who holds the patent on both agents. Daiichi assigned the patents to Defendants and gave Defendants an exclusive license to manufacture and market both its fluoroquinolone compounds in the United States in return for royalty fees. Daiichi licenses levofloxacin to Aventis for manufacture and market in European counties. To date, Levaquin remains one of Daiichi's best selling pharmaceuticals.

21. Daiichi ensured that the post market surveillance of levofloxacin would be tracked world-wide by creating an international database to keep track of adverse events. This database ensured that Defendants could not ignore the post market experience of levofloxacin in other countries.

22. Ofloxacin was first introduced into the Japanese market in September 1985. Defendants introduced ofloxacin, under the brand name Floxin, in the United States six years later, in 1991.

23. Even before ofloxacin was marketed in Japan, Daiichi began researching products that could be the successor of ofloxacin. Daiichi wanted to develop a newer fluoroquinolone in order to be more competitive with Cipro and the other

fluoroquinolones by developing a drug with the same or better characteristics of ofloxacin that could be used both orally and by injection.

24. After many derivatives of ofloxacin were explored and synthesized, Daiichi isolated what is now known as levofloxacin. Levofloxacin is a purified version of one optically active form of ofloxacin, more specifically the L-isomer.

25. Accordingly, ofloxacin and Levaquin are pharmacologically very similar - in fact, so similar that Defendants alleged in their New Drug Application for Levaquin that the safety profile of Levaquin would be expected to mirror that of ofloxacin.

26. Unfortunately, while Levaquin did closely follow the safety profile of ofloxacin, Levaquin was worse with respect to certain adverse effects, including tendon toxicity.

B. EPIDIMIOLOGY OF FLUOROQUINOLONE TENDON TOXICITY – OFLOXACIN IS MORE TENOTOXIC THAN THE REST, AND THE ELDERLY AND USERS OF CORTICOSTEROIDS ARE AT A HEIGHTENED RISK

27. Tendonitis as a side effect of fluoroquinolones was first reported in 1983. The first case of Achilles tendon rupture was reported in 1991 in conjunction with pefloxacin – a fluoroquinolone that has never been approved in the U.S, in part due to its teno-toxicity. Potentially due to pefloxacin’s early use in France, by 1994, Dr. Pierfitte et al. identified over 100 French patients with fluoroquinolone tendon disorders (mostly pefloxacin), and was able to observe that tendon injury occurred more frequently in patients over 60 and especially in those who had received steroid therapy.

28. Although the Achilles tendon was affected the most, and bilaterally in many cases, Dr. Pierfitte reported that other tendons could be implicated as well.

Accordingly, the French regulatory body was one of the first to notify physicians and their patients about the risk of fluoroquinolones-induced tendon injury. Additionally, as a likely result of Dr. Pierfitte's published observations, pefloxacin became severely restricted in use by 1995.

29. Once pefloxacin became restricted, Defendants' first generation ofloxacin emerged as the most tenotoxic fluoroquinolone on the market.

30. One of the first published reports regarding the tendon toxicity of ofloxacin was published in 1995 in the British Journal of Clinical Pharmacology (Wilton, L.V., Pearce, G.L. Mann, RD, *A comparison of ciprofloxacin, norfloxacin, ofloxacin, azithromycin and cefixime examined by observational cohort studies*. Br J Clin Pharmacol 1996; 41:277-284).

31. In the Wilton report, an analysis of prescription event monitoring data in the United Kingdom (a country where pefloxacin was not approved for market) revealed that ofloxacin was more tenotoxic than the other fluoroquinolones examined.

32. The United Kingdom's Regulatory Authority issued a bulletin, published in July 1995, stating that it had received 21 reports of tendon damage associated with fluoroquinolone antibiotics. The Authority reported "elderly patients and those treated concurrently with corticosteroid are at particular risk."

33. Defendants submitted their New Drug Application regarding Levaquin to the FDA in 1995. Though they indicate that tendon disorders are associated with fluoroquinolone use, Defendants failed to report that ofloxacin was more tendon toxic

than other currently marketed fluoroquinolones and failed to report that the tendon toxicity was exacerbated in the elderly, and especially in those taking corticosteroids.

34. The first epidemiological study to evaluate the relative risk of fluoroquinolone-induced tendonitis was published in 1999 by pharmacoepidemiologists and researchers at the Department of Epidemiology and Biostatistics and Internal Medicine at Erasmus Medical Center in Rotterdam. Van der Linden PD, Van de Lie J, Nab HW, Knok A, Stricker B H Ch, *Achilles tendonitis associated with fluoroquinolones*, Br J Clin Pharmacol 1999; 48: 433-437.

35. Data analyzed in this retrospective cohort study from 41 general practices in the Netherlands from 1995 and 1996 revealed that that ofloxacin had the strongest association with Achilles tendonitis. The adjusted relative risk of tendonitis to fluoroquinolones was 3.7, while Achilles tendonitis associated with ofloxacin had a relative risk of 10.1. Upon information and belief, Defendants knew of this study and had an obligation to inform the FDA of this study by supplementing their New Drug Application.

36. A second epidemiological study published in 2002 by Van der Linden et al. analyzed data from the IMS Health database in the United Kingdom which contained general practice medical records on a source population of 1 to 2 million inhabitants. Van der Linden, PD, Sturdenboom MCJM, Herings, RMC, Leufkens HGM, Stricker BH Ch, *Fluoroquinolones and risk of Achilles tendon disorders: case control study*, BMJ 2002; 324:1306-1307.

37. In this nested case control study, the authors again found that ofloxacin was associated with an eleven fold increase in tendon disorders. More specifically, the authors found that the relative risk of Achilles tendon disorders following current use of fluoroquinolones was 1.9, but in patients over 60 years of age, the relative risk was 3.2. However, in the elderly, the relative risk was 11.5 for current use of ofloxacin, compared to 2.3 and 1.8 for ciprofloxacin and norfloxacin respectively. In patients of 60 years and older, concurrent use of corticosteroids and fluoroquinolones increased the risk to 6.2. Upon information and belief, Defendants knew of this study and had an obligation to inform the FDA of this study by supplementing their New Drug Application.

38. Soon thereafter, in 2003, Dr. Van der Linden published his final epidemiological study, a larger population-based case control study that analyzed cases of Achilles tendon rupture and fluoroquinolone use from 1988 to 1998. Stuningly, his report concludes that the relative risk of a tendon injury in patients over 60 years old taking ofloxacin was 27.7 compared to ciprofloxacin's 3.4. He also found that use of corticosteroids nearly doubled the risk for tendon injury for patients over 60 years old.

39. Even Daiichi, the inventor of ofloxacin and Levaquin, published a 1997 rat study that admitted that Levaquin and ofloxacin were the most toxic to tendons of all the fluoroquinolones marketed in the United States. The study was designed to not only better understand the pathophysiological mechanism of fluoroquinolone-induced tendon disorders, but also to compare the relative tendon toxicity of ten different fluoroquinolones.

40. Although the exact mechanism of how fluoroquinolones cause tendon injury is still being investigated, studies have suggested that fluoroquinolones can degrade tendon cells by causing apoptosis, or a programmable cell death, and therefore lose their integrity, and easily tear and/or rupture.

41. The outcome of Achilles tendon ruptures in persons over 60 - the population most affected by this adverse reaction, is not favorable. Treatment may include a corticosteroid to decrease inflammation - the very drug that, when combined with a fluoroquinolone, can dramatically increase the risk of a tendon rupture. In the event of a tendon rupture, the leg is often immobilized through a boot or other casting for anywhere between six weeks to six months, and physical therapy is ordered thereafter. Surgery is frequently not recommended in the elderly population due to poor recovery rates. However, even with immobilization for long periods of time and physical therapy, the Achilles tendons in the elderly rarely fully recover.

C. THE FIRST U.S. TENDON WARNING

42. According to the U.S. consumer watchdog organization, Public Citizen, by 1996 there were over 130 reports of tendon injury from around the world over a ten year period and 52 reports of tendon injury in the United States associated with fluoroquinolone use.

43. As there was no mention of any fluoroquinolone-induced tendon injury on the label, Public Citizen petitioned the FDA in 1996, based on the number of adverse event reports world wide, to require that manufacturers of fluoroquinolones revise their product label to alert physicians of this unusual adverse event.

44. The FDA responded by requiring the following warning on all fluoroquinolone labels:

“Ruptures of the shoulder, hand, and Achilles tendons that required surgical repair or resulted in prolonged disability have been reported with [the specific drug name]. [The specific drug name] should be discontinued if the patient experiences pain, inflammation, or rupture of a tendon. Patients should rest and refrain from exercise until the diagnosis of tendonitis or tendon rupture has been confidently excluded. Tendon rupture can occur at any time during or after therapy with [the specific drug name].”

45. By 1997, U.S. manufacturers of fluoroquinolones had modified their label. However, the label was buried in a long list of potential adverse reactions; it was not highlighted in any way, such as with bold lettering, or even a heading titled “tendon injury.” Moreover, no mention was made of the fact that age and corticosteroid use tripled the risk of tendon injury. No letter to physicians was disseminated, and Defendants did not highlight this unusual effect when promoting Levaquin to doctors.

D. LEVAQUIN’S EARLY POST-MARKET EXPERIENCE

46. Levaquin was first introduced in Japan in 1993 and later introduced in the United States in 1997.

47. As has previously been alleged and described, before Levaquin’s market launch in the United States, Defendants had knowledge that:

- a. Levaquin would be as toxic as ofloxacin;
- b. Ofloxacin was revealing itself as one of the most tenotoxic fluoroquinolones on the market; and
- c. The elderly, and especially those using corticosteroids, were at least three times as likely to suffer a tendon injury.

48. Despite this unique knowledge, Defendants chose to use the same label that the FDA required on all other fluoroquinolones. Accordingly, in 1997, most U.S. physicians did not understand fluoroquinolone tendon toxicity, and were completely ignorant of the elderly's exceptional vulnerability to this antibiotic, especially those dependent on corticosteroids.

49. A look at Defendants' sales materials could explain why: the very group that Levaquin was most toxic to was the very market Defendants were after. Defendants' target market for Levaquin was the elderly - especially those with upper respiratory infections who were likely to be chronic corticosteroid users.

50. More disturbing, Defendants' promotional campaign was themed on Levaquin's excellent safety profile and failed to disclose the risks of tendon injury.

51. Defendants capitalized on Levaquin's early introduction into Japan and other countries by using pre-U.S. prescription sales data to assert that Levaquin had been prescribed frequently with few adverse events.

52. For example, one such advertisement boasted that Levaquin had "An Outstanding Record of Safety" as "[o]ver 63,000,000 patients worldwide" had taken the drug and only diarrhea and nausea had shown up as adverse effects, albeit rarely.

53. Cleverly, the promotional literature only reported on adverse events in U.S. *clinical trials* where only a very small sampling of patients took their drug, and where many adverse events do not necessarily reveal themselves. So, Defendants claimed "proven performance" on the 63,000,000 million people that had used Levaquin outside

the United States, but chose not to disclose the adverse events that were being reported on this same population.

54. As Levaquin gained traction, its “Achilles heel” of heightened tenotoxicity revealed itself.

55. Levaquin enjoyed immediate popularity in the Italian market. Introduced to Italy in 1998, Levaquin became Italy’s best selling fluoroquinolone, surpassing Cipro, the major market leader, in just three years. Curiously, ofloxacin, Defendant’s previous fluoroquinolone, had the lowest market share, which was consistent with Daiichi’s plan to “cannibalize” ofloxacin in favor of Levaquin.

56. One of the first comparative studies that included post market experience with Levaquin was from Italy. The authors analyzed Italian adverse event data from 1999 to 2001 to help determine the relative toxicity of each marketed fluoroquinolone antibiotic.

57. The Italian study was published in 2003 and revealed 1) the most frequently reported serious reaction to fluoroquinolones were tendon disorders 2) levofloxacin was the fluoroquinolone with the highest tendonitis reporting rate, and 3) levofloxacin ranked first for tendonitis reports during the same period in the World Health Organization’s adverse event database, with 522 reports of levofloxacin-induced tendon disorders and ruptures.

58. Not surprisingly, in March 2002, the Italian Health Ministry issued a Dear Doctor letter to inform physicians of the risk of Levaquin tendon rupture.

59. France also reported a particularly large amount of tendon disorders soon after Levaquin was first marketed to that country in September 2000. By June 2001, in just nine months, 333 adverse reactions had been reported, with tendon disorder being the most frequently reported adverse event. Again, the adverse event data supported the epidemiological evidence finding that tendon injuries were more prominent in the elderly, especially when there had been co-administration of corticosteroids.

60. France's regulatory authority published a Dear Doctor letter to highlight this information in 2002.

61. Similarly, the Belgian regulatory authority received 161 reports of Levaquin-induced tendon injury, including 68 reports of tendon rupture, in the first two years of Levaquin's introduction to Belgium. Again, the average age of patients with levofloxacin-associated tendinopathy was 69 years old and about half were receiving concomitant corticosteroid treatment. As with other adverse event data, the tendon injuries were reported to occur soon after Levaquin was ingested. Belgium also noted, similar to Italy, that the number of tendon disorders associated with levofloxacin was much higher than that of the other quinolones. Not surprisingly, ofloxacin had the second highest reports of tendon injury.

62. Recognizing that the number of tendon effects from Levaquin were far more frequent than any of the older fluoroquinolones which had all been on the market over the past ten years, the Belgium regulatory authority also disseminated a Dear Doctor letter in 2002 highlighting their concerns about levofloxacin's toxicity and suggesting that levofloxacin is only justified for the treatment of community-acquired pneumonia in

patients who are allergic to beta-lactams. The agency stressed the elderly and people who used corticosteroids were particularly at risk and encouraged doctors that if levofloxacin treatment was necessary, to watch for tendon injury.

63. After nearly five years on the market in the United States, and following the post-marketing data out of Europe, Defendants finally decided to update their tendon warning.

E. LEVAQUIN'S SECOND TENDON WARNING

64. The pre-2002 Levaquin label bore the required tendon warning from its market launch in 1997. It was the last of the warnings listed, with no header or any other identification to alert a practitioner to this unusual side effect. The warning was behind gastrointestinal affects, hypersensitivity reactions, and even the rare event of anaphylactic shock.

65. In 2002, Defendants embedded the following in the existing tendon warning: "Post-marketing surveillance reports indicate that this risk may be increased in patients receiving concomitant corticosteroids, especially in the elderly."

66. Through an international database managed by Daiichi, Defendants had access to the post market surveillance data all over the world, and specifically France, Belgium, Italy, and the United Kingdom.

67. By 2002, the adverse event data in all those countries consistently and unequivocally revealed that the risk of tendon injury was nearly triple for people over 60 as compared to people under 60. Additionally, Defendants had knowledge of at least one

epidemiological study confirming the age effects of fluoroquinolone use. All data pointed to the fact that Levaquin was more tendon toxic than all other fluoroquinolones.

68. Despite a wealth of information, Defendants chose not to warn their target patient population - the elderly - with their 2002 warning. Instead, they muted their additional tendon warning by flipping the confounders. Rather than warn that the risk of tendon injury was increased (tripled) in the elderly, the warning stated that that the risk was possibly increased in those using corticosteroids. According to Defendants' warning, any elderly person not on corticosteroids therefore had no additional risk of a tendon injury, and the fact that the warning was so equivocal regarding corticosteroids diffused any possible effect of warning physicians of the effect of age on the frequency and severity and of this debilitating injury.

69. Nor did Defendants make any effort to highlight this new information to its prescribing doctors - Defendants did not send any Dear Doctor letters regarding the 2002 label change to any healthcare practitioners, as had been done in Italy, Belgium, and France.

70. Accordingly, despite the 2002 label change, Levaquin prescriptions only increased, and tendon injuries mounted.

F. DEFENDANTS THWART EFFORTS TO HIGHLIGHT LEVAQUIN'S INCREASED RISK OF TENDON INJURY

71. Alarmed by the early post market experience with Levaquin, France, Belgium, Italy, the United Kingdom and other European countries convened before the European Agency for the Evaluation of Medicinal Products (EMA) as early as September 2001 to discuss a heightened warning for levofloxacin.

72. The EMEA proposal was that levofloxacin would be singled out as the most tendon toxic of the fluoroquinolones with a warning that stated that levofloxacin (marketed under the brand name Tavanic) was associated with a greater frequency of tendinopathy and tendon rupture than other fluoroquinolones.

73. Aventis Pharmaceutical was the manufacturer and distributor of levofloxacin in Europe.

74. Under increasing pressure to agree to the proposed changes to the warning label, Aventis conducted two epidemiological studies in Europe regarding the relative tendon toxicity of levofloxacin. The first study used the United Kingdom's General Practitioners Research Database of medical records from 1997 through 2001, and the second used Germany's Mediplus database of medical records from 1998 to 2001.

75. Before releasing the results of the two epidemiology studies to the European regulatory authorities, and ostensibly because of the results of the studies, Aventis contracted with Defendants, specifically Johnson & Johnson Pharmaceutical Research & Development, to fund and co-author a study in the United States on tendon rupture and fluoroquinolones.

76. Advocating that the U.S. Study would be the largest epidemiological study to date and therefore provide the most definitive evidence of the relative risk of levofloxacin and tendon injury, and that the European studies to date were too small from which to base a label change, Aventis convinced the European regulatory authorities to forestall the proposed warning change until the preliminary data from the U.S. study was released.

77. In or around April 2002, Aventis submitted the results of their two European epidemiological studies to the United Kingdom's regulatory authority, the Medicines and Healthcare Products Regulatory Agency (MHRA).

78. The epidemiology studies conducted by Aventis in Europe concluded that levofloxacin was associated with a higher rate of tendon injury than all the other fluoroquinolones compared. Ofloxacin, the fluoroquinolone indicted in early epidemiological studies as the most teno-toxic, came in second.

79. An assessor at the MHRA concluded that the two epidemiological studies had findings "supporting a signal generated by spontaneous reporting with respect to an increased risk of tendinopathy with levofloxacin compared to other fluoroquinolones."

80. Moreover, the assessor remarked "the finding that ofloxacin (the racemate) is associated with an intermediate level of risk makes pharmacological sense, suggesting that the L-rather than the D-isomer of ofloxacin is likely to be responsible for tendon toxicity....given the consistency and plausibility of the findings, a real difference is the most likely explanation."

81. By the time Aventis released the results of their epidemiological studies, the preliminary results of the U.S. study was reportedly only six months away. Accordingly, the European regulatory authorities agreed to wait before forcing a label change.

82. The U.S. epidemiological study was funded and co-authored by Defendant Johnson & Johnson Pharmaceutical Research & Development.

83. Unlike the healthcare databases in Europe which contain computerized medical records, Johnson & Johnson PRD used data from the Ingenix Research database which consisted of U.S. health insurance claims data from 1997 to 2001. The study analyzed only Achilles tendon ruptures and sought to examine whether fluoroquinolone exposure was a risk factor for this injury. It did not assess the relative risk of Levaquin tendon toxicity, as had been requested by the United Kingdom.

84. Under the guise of data validation, Defendant Johnson & Johnson PRD created an algorithm that conveniently excluded nearly 70 percent of health claims for elderly persons who suffered Achilles tendon rupture.

85. The algorithm used CPT procedure codes that only related to surgical repair which thereby excluded all those Achilles tendon rupture cases where the patient was casted or booted, as is the case in the elderly population.

86. By manipulating the data, Defendant Johnson & Johnson PRD was able to exclude the very group that was prone to tendon rupture.

87. Not surprisingly, the results of the U.S. epidemiological study – the study upon which hinged regulatory action in Europe with ramifications to the U.S. market – revealed for the first time that there was no increased risk of Achilles tendon rupture associated with any fluoroquinolone use. Neither the confounders of age nor corticosteroid use altered these findings.

88. Indeed, when one includes the data that was excluded by the algorithm, the result becomes consistent with the approximately eight other epidemiological studies performed on the topic. See Seeger et al. *Achilles Tendon Rupture and its Associations*

with Fluoroquinolone Antibiotics and Other Potential Risk Factors in a Managed Care Population, *Pharmacoepidemiology and Drug Safety* 2006; 15: 784-792 (“There was a stronger association with fluoroquinolone antibiotic exposure among these “ruled-out” cases of ATR . . . than among the decision rule confirmed cases. This association was stronger with exposure close to the date of the rupture and was more pronounced among the elderly.”)

89. As a result of Defendants’ misrepresentations in the U.S. Study, the MHRA and the other European regulatory agencies chose not to revise the levofloxacin label as they had previously recommended.

G. DEFENDANTS DOWNPLAY THE RISK OF LEVAQUIN TO PHYSICIANS

90. Consistent with their plan to downplay Levaquin’s known risk of tendon injury, Defendants made no attempts to educate physicians in the United States about this unusual adverse event. Although Dear Doctors had been widely disseminated throughout Europe advising of Levaquin’s tendon toxicity and the vulnerability of this adverse event to the elderly, Defendants did not so advise the U.S. physicians.

91. Defendants plan was to hide behind the class warning and blame any tendon injuries reported on the general pharmacological properties of a fluoroquinolone antibiotic rather than on the L-isomer of the ofloxacin compound as the Aventis studies suggested.

92. Promotional material designed and distributed by Defendants, and more specifically, by Ortho-McNeil, consistently omits the risk of tendon injury on materials left with physicians.

93. Accordingly, physicians continued to prescribe Levaquin believing it to have the same safety profile as Cipro and unaware of the heightened affect of Levaquin on the elderly population.

H. AN EXPLOSION OF TENDON INJURIES RESULTS IN A THIRD LABEL CHANGE

94. A review of the events in the FDA Adverse Event database from 1997 through 2005, *for Levaquin alone*, showed 1,044 reports of tendon injuries, with 282 reports of tendon rupture. This six year figure for tendon affects associated with Levaquin far surpassed the ten year history of tendon affects from 1985 through 1995 associated with all pre-Levaquin fluoroquinolones.

95. After Cipro went generic in 2003, Levaquin became the number one prescribed fluoroquinolone in the United States. And when Zithromax, a highly popular macrolide antibiotic, went generic after its patent expired in 2005, Levaquin became the number one prescribed antibiotic in the world in 2006.

96. Corresponding with Levaquin's increased popularity, the number of adverse events reported to the FDA reported soared. 143 tendon related injuries were reported in 2006, and in just the first quarter 2007, 107 tendon related injuries were reported where Levaquin was the primary suspect.

97. The Levaquin phenomenon did not go unnoticed by the Illinois Attorney General. On May 18, 2005, the Attorney General submitted a petition to the FDA requesting a black box warning on fluoroquinolones. The Attorney General suggested that the black box was necessary to highlight the seriousness of tendon injuries and that the risk is increased in the elderly and in patients on corticosteroids.

98. The Attorney General also requested that the manufacturer issue a Dear Doctor letter to inform the health care providers about this significant hazard to health, as the tenotoxic affects of fluoroquinolones were not well known to practicing physicians.

99. In the Petition, the Attorney General's office reviewed the literature and stated that tendon injuries were not a rare complication of fluoroquinolone use. The Petition complained that the current tendon warning was "buried in lists of potential side effects which are both less frequent and less severe."

100. One year later, the same consumer watchdog organization that petitioned the FDA in 1996 for a tendon warning, petitioned the FDA again saying the first tendon warning did not go far enough. Citing the alarming increase in reports of tendon injury, Public Citizen joined the Illinois Attorney General's petition and urged that the FDA place a black box warning regarding the risk of tendonitis and tendon rupture.

101. At the request of the FDA, in April 2007, the Levaquin label changed for a third time with regard to tendon injuries. The April 2007 label was not a black box warning, but it did state that indeed the elderly are at an increased risk of tendon injury, and unequivocally stated that the risk of tendon injuries is increased with concomitant use of corticosteroids, contrary to the results of Defendant's Ingenix study.

102. Upon information and belief, Defendants negotiated with the FDA and insisted on a class warning to thereby minimize the heightened risk of tendon injury with Levaquin.

103. The April 2007 label change failed to alert physicians and prescribing health care providers that Levaquin is more toxic to the tendons than the other fluoroquinolones available in the U.S. market.

104. Defendants, upon information and belief, did not advise physicians of the 2007 label change and, therefore, it is not known whether physicians received this new information regarding the vulnerability of the elderly population to a Levaquin-induced injury.

I. THE NEW “BLACK BOX” WARNING

105. On July 8, 2008, the FDA notified Defendants and other fluoroquinolone antimicrobial manufacturers that a black box warning and Medication Guide must be added to the prescribing information for Levaquin and other fluoroquinolones to provide stronger warnings about the increased risk of suffering tendonitis and tendon rupture in patients using Levaquin and other fluoroquinolones.

106. Although the black box warning will indicate that the risk of tendinitis and tendon rupture is further increased in patients over 60; in kidney, heart and lung transplant recipients, and with the use of concomitant steroid therapy, it does not warn health care providers that Levaquin is much more tenotoxic than other fluoroquinolones and therefore physicians will interpret the relative risk of a Levaquin-induced tendon injury inappropriately.

107. Defendants continue to market Levaquin as a first line therapy for the common bronchitis and sinusitis infections, and for which many other, safer, antibiotics are available.

SPECIFIC FACTUAL ALLEGATIONS

108. Clifford Straka was 72 years of age when he was prescribed and began consuming Levaquin on March 27, 2006 to treat possible pneumonia. After using Levaquin for approximately nine days, Mr. Straka suffered bilateral Achilles tendon ruptures. Following the ruptures, Mr. Straka was placed in a walking boots and received physical therapy treatment. Prior to suffering rupturing both Achilles tendons, Mr. Straka was able to perform all activities of daily living independently. As a direct and proximate cause of his Levaquin-induced, bilateral Achilles tendon ruptures, Plaintiff Straka's ability to perform normal daily tasks has been compromised and his quality of life has been severely diminished.

**FIRST CAUSE OF ACTION
STRICT LIABILITY**

109. Plaintiff incorporates by reference all other paragraphs of this Complaint as if fully set forth herein and further alleges as follows:

110. At all relevant times hereto, Defendants were engaged in the development, testing, manufacturing, marketing and sales of Levaquin. Defendants designed, manufactured, marketed, and sold Levaquin to medical professionals and their patients, knowing it would be ingested for the treatment of infections.

111. Levaquin as designed, manufactured, marketed and sold by Defendants reached Plaintiff without substantial change in its condition and was used by Plaintiff in a reasonably foreseeable and intended manner.

112. Levaquin was "defective" and "unreasonably dangerous" when it entered the stream of commerce and was received by Plaintiff, because it was dangerous to an

extent beyond that which would be contemplated by the ordinary consumer. At no time did Plaintiff have reason to believe that Levaquin was in a condition unsuitable for proper and intended use among patients.

113. Levaquin was used in the manner for which it was intended, that is, for treatment of bacterial infections. This use resulted in injury to Plaintiff.

114. Plaintiff was not able to discover, nor could he have discovered through the exercise of reasonable care, the defective nature of Levaquin. Further, in no way could Plaintiff have known that Defendants had designed, developed, and manufactured Levaquin in such a way as to increase the risk of harm or injury to the recipients of Levaquin.

115. Levaquin is defective in design because of its propensity to cause tendon ruptures and other serious tendon injuries.

116. Levaquin is unreasonably dangerous because it was sold to Plaintiff without adequate warnings regarding, *inter alia*, the propensity of Levaquin to cause serious tendon injuries; the post-marketing experience with Levaquin; the increased risk of tendon injury in patients over the age of 60; the numbers of tendon-related adverse events reported; and the probability of suffering an acute tendon injury when ingesting corticosteroids concomitantly with Levaquin or post-Levaquin use.

117. Defendants failed to develop and make available alternative products that were designed in a safe or safer manner, even though such products were feasible and marketable at the time Defendants sold Levaquin to Plaintiff.

118. Defendants had knowledge and information confirming the defective and dangerous nature of Levaquin. Despite this knowledge and information, Defendants failed to adequately and sufficiently warn Plaintiff and his physicians that Levaquin causes serious tendon injuries including, without limitation, tendon rupture.

119. As a direct and proximate result of Defendants' wrongful conduct, including Levaquin's defective and dangerous design and inadequate warnings, Plaintiff has sustained and will continue to sustain severe and debilitating injuries, economic loss, and other damages including, but not limited to, cost of medical care, rehabilitation, lost income, permanent instability and loss of balance, immobility, and pain and suffering, for which he is entitled to compensatory and equitable damages and declaratory relief in an amount to be proven at trial.

SECOND CAUSE OF ACTION
NEGLIGENCE

120. Plaintiff incorporates by reference all other paragraphs of this Complaint as if fully set forth herein and further alleges as follows:

121. At all relevant times, Defendants had a duty to exercise reasonable care in the design, formulation, testing, manufacture, marketing, sale, and distribution of Levaquin, including a duty to ensure that Levaquin did not pose a significantly increased risk of bodily injury to its users.

122. Defendants had a duty to exercise reasonable care in the advertising and sale of Levaquin, including a duty to warn Plaintiff and other consumers, of the dangers associated with the consumption of Levaquin that were known or should have been known to Defendants at the time of the sale of Levaquin to the Plaintiff.

123. Defendants failed to exercise reasonable care in the design, testing, manufacture, marketing, sale and distribution of Levaquin because Defendants knew or should have known that Levaquin had a propensity to cause serious injury, including tendon rupture and other serious tendon injuries.

124. Defendants failed to exercise ordinary care in the labeling of Levaquin and failed to issue adequate pre-marketing or post-marketing warnings to prescribing doctors and the general public regarding the risk of serious injury, including, without limitation, tendon rupture.

125. Defendants knew or should have known that Plaintiff could foreseeably suffer injury as a result of Defendants' failure to exercise ordinary care as described above.

126. Defendants breached their duty of reasonable care to Plaintiff by failing to exercise due care under the circumstances.

127. As a direct and proximate result of Defendants' acts and omissions, including their failure to exercise ordinary care in the design, formulation, testing, manufacture, sale, and distribution of Levaquin, Plaintiff ingested Levaquin and suffered severe and debilitating injuries, economic loss, and other damages, including but not limited to, cost of medical care, rehabilitation, lost income, permanent instability and loss of balance, immobility, and pain and suffering, for which he is entitled to compensatory and equitable damages and declaratory relief in an amount to be proven at trial.

THIRD CAUSE OF ACTION
BREACH OF IMPLIED WARRANTIES

128. Plaintiff incorporates by reference all other paragraphs of this Complaint as if fully set forth herein and further alleges as follows:

129. Defendants designed, formulated, tested, manufactured, marketed, sold, and distributed Levaquin as has previously been alleged and described herein.

130. At the time Defendants marketed, sold and distributed Levaquin, Defendants knew of the use for which Levaquin was intended and impliedly warranted that Levaquin was merchantable, safe and fit for its intended purpose: namely that Plaintiff could ingest Levaquin without the risk of serious injury.

131. Plaintiff, foreseeable users of Levaquin, and Plaintiffs' physician(s), reasonably relied upon Defendants' judgment and implied warranties in purchasing and consuming Levaquin as intended.

132. Levaquin was defective, unmerchantable, and unfit for ordinary use when sold, and subjected Plaintiff to severe and permanent injuries.

133. Defendants breached their implied warranties because Levaquin was and continues to be neither of merchantable quality nor safe for its intended use in that Levaquin has the propensity to cause tendon rupture, other debilitating tendon injuries, and bodily harm.

134. As a direct and proximate result of Defendants' breach of the implied warranties of merchantability and fitness for its intended purpose, Plaintiff ingested Levaquin and suffered severe and debilitating injuries, economic loss, and other damages, including but not limited to, cost of medical care, rehabilitation, lost income, permanent

instability and loss of balance, immobility, and pain and suffering, for which they are entitled to compensatory and equitable damages in an amount to be proven at trial.

FOURTH CAUSE OF ACTION
BREACH OF EXPRESS WARRANTY

135. Plaintiff incorporates by reference all other paragraphs of this Complaint as if fully set forth herein and further alleges as follows:

136. Defendants through their marketing program, promotional activities, product labeling, package inserts, and other written and verbal assurances expressly warranted to physicians and consumers, including Plaintiff and/or his physicians, that Levaquin had been shown by scientific study to be safe for its intended use.

137. Plaintiff, and his physicians, reasonably relied upon Defendants' express warranties in purchasing consuming, and prescribing Levaquin.

138. Defendants breached their express warranties because Levaquin as manufactured and sold by Defendants does not conform to these express representations in that Levaquin has a propensity to cause tendon rupture, other serious tendon injuries, and bodily harm.

139. As a direct and proximate result of Defendants' breach of their express warranties, Plaintiff ingested Levaquin and suffered severe and debilitating injuries, economic loss, and other damages, including but not limited to, cost of medical care, rehabilitation, lost income, permanent instability and loss of balance, immobility, and pain and suffering, for which he is entitled to compensatory and equitable damages and declaratory relief in an amount to be proven at trial.

FIFTH CAUSE OF ACTION
FRAUD

140. Plaintiff incorporates by reference all other paragraphs of this Complaint as if fully set forth herein and further alleges as follows:

141. Defendants were under a duty and failed to discharge their duty to exercise reasonable care to disclose to Plaintiff and his physician, Dr. Katayoun Baniriah of the Mayo Clinic, the defective nature and risks that Levaquin can cause severe and permanent injuries, including, without limitation, tendon ruptures, of which they had special knowledge not available to Plaintiff or his doctors, and as to which they made affirmative representations in violation of all applicable laws, and concealed material facts relating to the defective nature and risks of Levaquin, which were peculiarly within its knowledge, knowing that Plaintiff and his doctors would rely on the presumption that no such facts exist.

142. Defendants knew that Levaquin can cause severe and permanent injuries, including, without limitation, tendon ruptures; indeed, Defendants knew that tendon injuries associated with Levaquin had occurred for years. Defendants had actual knowledge at the time of sale of Levaquin to the Plaintiff that Levaquin created a risk of serious bodily injury to its users, including, without limitation, tendon injuries, based, in part, upon test results, studies, adverse reaction reports, regulatory action in foreign countries, published reports, and their own clinical trials and post-marketing surveillance of Levaquin and its molecularly similar counterpart, ofloxacin.

143. At all times during the course of dealing between Defendants and Plaintiff, Defendants knowingly and recklessly omitted and concealed information peculiarly

within their knowledge to the Plaintiff, his doctors, the scientific community and to the general public - e.g., the dangers of Levaquin, including the special risk of tendon injury and tendon ruptures, particularly to the elderly - knowing that the scientific community, the general public, the Plaintiff and his doctors, would rely on the presumption that the dangers did not exist.

144. Defendants actively concealed from the Plaintiff, his doctors, the scientific community and the general public:

- i. that their own test results, published studies, and/or clinical trials showed a statistically high risk of serious tendon injuries associated with Levaquin including, without limitation, tendon ruptures; and/or
- ii. that Levaquin was not adequately tested for serious tendon injuries before or after its introduction on the market; and/or
- iii. that Levaquin was, in fact, unsafe as it posed a risk of injury which outweighed any purported benefits.

145. Defendants misrepresented that Levaquin was safe and effective for its intended uses by affirmative misrepresentation, and/or active concealment and omission of material facts regarding the safety and effectiveness of Levaquin, and by their course of conscious or intentional conduct succeeded in selling and marketing dangerous, defective, and ineffective antibiotics to be ingested by Plaintiff. Defendants intentionally omitted, concealed and/or suppressed this information from consumers, including Plaintiff and his doctors, in order to avoid losses in sales to consumers and market share to its major competitors.

146. Moreover, Defendants engaged in an aggressive marketing strategy, which included false representations regarding the safety profile and known adverse side effects of Levaquin to create the impression and to convey to Plaintiff and the general public that:

- i. Levaquin had a favorable safety profile and was fit for human consumption;
- ii. the benefits of taking Levaquin outweighed any associated risks; and
- iii. the use of Levaquin was safe and had fewer adverse health and side effects than were known or should have been known by Defendants at the time of these representations.

147. The omissions, misrepresentations and concealment described in the preceding paragraphs occurred, without limitation, in the Levaquin warning labels, advertisements and promotional materials, in the Johnson & Johnson funded or created scientific reports, and the failure to provide other special notification of the dangers of Levaquin to the Plaintiff or his physicians, for example, “Dear Doctor” letters. The Defendants’ statements omitted, concealed, and misrepresented the dangers of serious injury, including, but not limited to, tendon ruptures, particularly to the elderly, to Plaintiff and his prescribing doctors.

148. Defendants engaged in fraud by deliberately and affirmatively concealing and failing to disclose adverse reactions of Levaquin to Plaintiff, his doctors, the scientific community, and the general public, and by disseminating only positive and misleading scientific data, and by concealing scientific data that showed increased risk of

tendon-related injury, to Plaintiff, his doctors, the scientific community, and the general public.

149. Plaintiff Clifford Straka and his physician, Dr. Katayoun Baniriah of the Mayo Clinic, relied on the warning labels as they appeared in the prescribing information and patient package insert at the time they were prescribed and consumed Levaquin. The applicable warnings concealed and omitted material facts relating to the defective nature and risks of Levaquin. These dangers were peculiarly within the Defendants' knowledge, and were omitted and concealed knowing that Plaintiff and his doctors would rely on the presumption that no such facts exist.

150. Defendants knew or should have known that their representations and omissions regarding the safety of Levaquin were, in fact, false and/or misleading, and actively made such representations and omissions with the intent, design, and purpose that Plaintiff and others, including Plaintiff's prescribing physicians, rely on these representations leading to the prescription, purchase and consumption of Levaquin.

151. At all times herein, Plaintiff and his physicians, including Dr. Katayoun Baniriah, were unaware of the dangers of Levaquin with respect to tendon ruptures, including the special risk of tendon injury to the elderly, and were reasonably misled by the Defendants' omission of information about this danger.

152. At all times herein, Plaintiff and his physicians were unaware of the falsity underlying Defendants' statements and reasonably believed Defendants' false statements about the safety and efficacy of Levaquin to be true.

153. Plaintiff and his doctors could not have discovered Defendants' fraudulent and misleading conduct at an earlier date through the exercise of reasonable diligence because Defendants actively concealed their deceptive, misleading and unlawful activities.

154. Plaintiff and his physicians did, and could be expected to, reasonably and justifiably rely on Defendants' representations and omissions because Defendants held themselves out as having expertise and specialized knowledge in the pharmaceutical industry.

155. Plaintiff justifiably relied upon to his detriment and/or were induced by Defendants' false statements and active concealment over the safety of Levaquin, in part, because at no time did Plaintiff or his physicians have the knowledge or expertise necessary to independently evaluate the safety of Levaquin.

156. Defendants' misrepresentations, concealment, suppression and omissions were made willfully, wantonly, uniformly, deliberately, or recklessly, in order to induce Plaintiff to purchase Levaquin and Plaintiff and his physicians did reasonably and justifiably rely upon the material misrepresentations and omissions made by the Defendants about Levaquin when agreeing to purchase and/or ingest Levaquin.

157. As a direct and proximate result of Defendants' false representations and/or active concealment of material facts regarding the safety and efficacy of Levaquin, Plaintiff ingested Levaquin and suffered severe and debilitating injuries and economic loss, including but not limited to, cost of medical care, rehabilitation, lost income, permanent instability and loss of balance, immobility, and pain and suffering for which

he is entitled to compensatory and equitable damages and declaratory relief in an amount to be proven at trial.

SIXTH CAUSE OF ACTION
VIOLATION OF MINNESOTA'S UNFAIR AND DECEPTIVE TRADE
PRACTICES ACT (MINN. STAT. ANN. § 325D.13 and § 325D.44 ET SEQ.)

158. Plaintiff incorporates by reference all other paragraphs of this Complaint as if fully set forth herein and further alleges as follows:

159. Defendants have a statutory duty to refrain from unfair or deceptive acts or trade practices in the design, development, manufacture, promotion, and sale of Levaquin.

160. Had the Defendants not engaged in the deceptive conduct described above, Plaintiff would not have purchased and/or paid for Levaquin, and would not have incurred related medical costs.

161. Specifically, Plaintiff and his physicians at Mayo Clinic were misled by the deceptive conduct described herein.

162. Defendants' deceptive, unconscionable, or fraudulent representations and material omissions to patients, physicians and consumers, including Plaintiff, constituted unfair and deceptive acts and trade practices in violation of Minn. Stat. Ann. § 325D.13 and § 325D.44 *et seq.*

163. Defendants engaged in wrongful conduct while at the same time obtaining, under false pretenses, substantial sums of money from Plaintiff for Levaquin that they would not have paid had Defendants not engaged in unfair and deceptive conduct.

164. Defendants' actions, as complained of herein, constitute unfair competition or unfair, unconscionable, deceptive, or fraudulent acts or trade practices in violation of Minn. Stat. Ann. § 325D.13 and § 325D.44 *et seq.*

165. Plaintiff was injured by the cumulative and indivisible nature of Defendants' conduct. The cumulative effect of Defendants' conduct directed at patients, physicians and consumers was to create a demand for and sell Levaquin. Each aspect of Defendants' conduct combined to artificially create sales of Levaquin.

166. The medical community relied upon Defendants' misrepresentations and omissions in determining which antibiotic to utilize.

167. By reason of the unlawful acts engaged in by Defendants, Plaintiff has suffered ascertainable loss and damages.

168. As a direct and proximate result of Defendants' wrongful conduct, Plaintiff was damaged by paying in whole or in part for Levaquin.

169. As a direct and proximate result of Defendants' violations of Minn. Stat. Ann. § 325D.13 and § 325D.44 *et seq.*, Plaintiff has sustained economic losses and other damages for which he is entitled to statutory and compensatory damages and declaratory relief in an amount to be proven at trial.

SEVENTH CAUSE OF ACTION
MINNESOTA'S SENIOR CITIZEN AND HANDICAPPED PERSON
CONSUMER FRAUD ACT (MINN. STAT. ANN. § 325F.71)

170. Plaintiff incorporates by reference all other paragraphs of this Complaint as if fully set forth herein and further alleges as follows:

171. Pursuant to Minn. Stat. § 325F.71(2), this Count applies to those Plaintiffs who are Senior Citizens and/or Handicapped persons.

172. Minn. Stat. §325F.71, subdiv. 2 incorporates Minn. Stat. 325D.43-48 regarding deceptive trade practices, § 325F.67 regarding false advertising and § 325F.68-70 regarding consumer fraud and provides special remedies if violations of those statutes are directed against senior citizens or handicapped people, including priority of restitution § 325F.71, subdiv 3, and the recovery of “damages, including costs of investigation and reasonable attorney’s fees” and to “receive other equitable relief as determined by the court.” Id. § 325F.71, subdiv. 4.

173. The affirmative misrepresentations and the pattern of omissions by Defendants described above, violated Minn. Stat. § 325F.44, subdiv. 1, (5) because, through affirmative misrepresentations and the pattern of omissions, Defendants represented that Levaquin had “characteristics, ingredients, uses [and/or] benefits...that they do not have” - a per se violation of § 325F.71.

174. The affirmative misrepresentations and the pattern of omissions by Defendants described above, violated § 325.44, subdiv. 1, (7) because, through those affirmative misrepresentations and the pattern of omissions, Defendants represented that Levaquin was of a “particular standard, quality, or grade”, when it was, in fact, of a much lower standard, quality or grade – a per se violation of § 325F.71.

175. The affirmative misrepresentations and the pattern of omissions by Defendants in their Annual Reports, advertising literature, press releases, and other public

statements, constitutes false advertising as prohibited by § 325F.67 - a per se violation of § 325F.71.

176. The conduct, affirmative misrepresentations, and the pattern of omissions by Defendants described above constitutes a “fraud, false pretenses, false promise, misrepresentation, misleading statement or deceptive practice, with the intent that others rely thereon in connection with the sale of ...[Levaquin]”, in violation of Minn. Stat. § 325F.69, subdiv. 1 – a per se violation of § 325F.71.

177. Pursuant to Minn.. Stat. § 325F.71, subdiv. 4, Plaintiff is entitled to recover all damages arising out of Defendants’ violation of Minn. Stat. § 325F.44, subdiv 1, (5) and/or (7), §325F.67, and/or Minn. Stat. § 325F.69, subdiv. 1.

178. As a direct and proximate result of Defendants’ wrongful conduct, Plaintiff has sustained and will continue to sustain severe physical injuries, economic losses, and other damages, including but not limited to, cost of medical care, rehabilitation, lost income, permanent instability and loss of balance, immobility, and pain and suffering for which he is entitled to statutory and compensatory damages and declaratory relief in an amount to be proven at trial.

179. In addition, Plaintiff is entitled to recover “costs of investigation and reasonable attorney’s fees”, pursuant to Minn. Stat. §325F.71, subdiv. 4 and the Court is urged to give priority to the remedy of restitution pursuant to Minn. Stat. §325F.71, subdiv. 3.

EIGHTH CAUSE OF ACTION
VIOLATION OF MINNESOTA'S CONSUMER FRAUD ACT
(MINN.STAT. § 325F.69)

180. Plaintiff incorporates by reference all other paragraphs of this Complaint as if fully set forth herein and further alleges as follows:

181. Defendants acted, used, and/or employed fraud, false pretense, false promise, misrepresentation, misleading statements and/or deceptive practices, concerning the safety, use, efficacy, and testing of Levaquin with the intent that others, including Plaintiff, rely upon those false and deceptive acts in determining whether to use Levaquin.

182. In its marketing, direct-to-consumer advertising, promotion, sale, and distribution of Levaquin, Defendants knowingly, unfairly, and deceptively promised and represented that Levaquin is a safe and effective antibiotic while failing to disclose the known properties, ingredients, characteristics, qualities and risks associated with Levaquin when the Defendants had actual knowledge or should have known of the serious adverse health effects associated with Levaquin, including but not limited to, tendon ruptures.

183. Defendants made such misrepresentations and omissions of material fact with the intent, design, and purpose that consumers, including Plaintiff, rely on such representations in choosing to purchase Levaquin.

184. As a direct and proximate result of Defendants' fraudulent sale and marketing, Plaintiff ingested Levaquin, and suffered severe and debilitating injuries and economic loss, including but not limited to, cost of medical care, rehabilitation, lost

income, permanent instability and loss of balance, immobility, and pain and suffering for which he is entitled to statutory and compensatory damages and declaratory relief in an amount to be proven at trial.

NINTH CAUSE OF ACTION
VIOLATION OF MINNESOTA'S FALSE ADVERTISING ACT
(MINN. STAT. § 325F.67)

185. Plaintiff incorporates by reference all other paragraphs of this Complaint as if fully set forth herein and further alleges as follows:

186. Defendants knowingly misrepresented Levaquin as a safe and effective antibiotic and knowingly made false statements and omissions of material fact concerning the properties, ingredients, characteristics, qualities, benefits, uses, efficacy, safety, and/or testing of Levaquin to the Plaintiff and the general public.

187. In its labeling, marketing, direct-to-consumer advertising, promotion, sale, and distribution of Levaquin, Defendants made untrue, deceptive, and/or misleading material assertions, representations, and/or statements downplaying risks associated with Levaquin and exaggerating the drug's safety to Plaintiff and the general public when Defendants had actual knowledge of the serious, adverse health effects associated with Levaquin including, but not limited to, tendon ruptures.

188. Defendants intended to increase the sale and consumption of Levaquin by falsely marketing Levaquin as safe and effective, and by concealing facts regarding the dangerous properties of Levaquin, to thereby induce Plaintiff's physicians to prescribe Levaquin and to ultimately cause Plaintiff to purchase and consume Levaquin.

189. In purchasing and consuming Levaquin, Plaintiff reasonably relied upon Defendants' false and misleading assertions and omissions of material fact that Levaquin was safe and effective for the treatment their illness.

190. Defendants' actions as described herein constitute unlawful, unfair, and deceptive trade practices within the meaning of Minn. Stat § 325F.67.

191. As a direct and proximate result of Defendants' false statements as herein alleged, Plaintiff ingested Levaquin and suffered severe and debilitating injuries and economic loss, including but not limited to, cost of medical care, rehabilitation, lost income, permanent instability and loss of balance, immobility, and pain and suffering for which he is entitled to statutory and compensatory damages and declaratory relief in an amount to be proven at trial.

TENTH CAUSE OF ACTION
UNJUST ENRICHMENT

192. Plaintiff incorporates by reference all other paragraphs of this Complaint as if fully set forth herein and further alleges as follows:

193. As the intended and expected result of their conscious wrongdoing, Defendants have profited and benefited from the purchase and implementation of Levaquin by Plaintiff.

194. Defendants have voluntarily accepted and retained those profits and benefits, derived from Plaintiff, with full knowledge and awareness that, as a result of Defendants' fraud and other conscious and intentional wrongdoing, Plaintiff was not receiving a product of the quality, nature, or fitness that had been represented by Defendants, or that Plaintiff, as a reasonable consumer, expected to receive.

195. By virtue of the conscious wrongdoing alleged above, Defendants have been unjustly enriched at the expense of Plaintiff, who is entitled in equity, and hereby seek, the disgorgement and restitution of Defendants' wrongful profits, revenues and benefits, to the extent and in the amount deemed appropriate by the Court; and such other relief as the Court deems just and proper to remedy Defendants' unjust enrichment.

WHEREFORE, Plaintiff prays for relief against Defendants, jointly and severally, as follows:

1. Compensatory damages according to proof, in excess of the amount required for federal diversity jurisdiction, and in an amount to fully compensate Plaintiff for all of his injuries and damages, both past and present;
2. Special damages according to proof, in excess of the amount required for federal diversity jurisdiction and in an amount to fully compensate Plaintiff for all of his injuries and damages, both past and present, including but not limited to, past and future medical expenses, costs for past and future rehabilitation and/or home health care, permanent disability, including permanent instability and loss of balance, and pain and suffering.
3. Double or triple damages as allowed by law;
4. Disgorgement of profits;
5. A full refund of cost of all Levaquin prescriptions;
6. Attorneys' fees, expenses, and costs of this action;

7. Pre-judgment and post-judgment interest in the maximum amount allowed by law; and
8. Such further relief as this Court deems necessary, just, and proper.

JURY DEMAND

Plaintiff demands a trial by jury of all claims asserted in this Complaint.

Dated: October 15, 2008

Respectfully submitted,

/s/ Ronald Goldser, Esq.
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