

**FILED**  
U. S. DISTRICT COURT  
EASTERN DISTRICT ARKANSAS

JUN 12 2019

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

JAMES W. MCCORMACK, CLERK  
By: [Signature]  
DEP CLERK

ROBERT BEST,

Plaintiff,

v.

CASE NO: 1:19-cv-46-JM

CONSENSUS ORTHOPEDICS, INC.  
and JOHN DOES 1-10,

**JURY DEMANDED PER F.R.C.P. 38**

Defendants.

**CIVIL ACTION COMPLAINT**

COMES NOW, Plaintiff Robert Best, by and through his counsel of record, and files this complaint against Defendants Consensus Orthopedics Inc. and John Does 1-10, and alleges as follows:

**PARTIES, JURISDICTION, AND VENUE**

1. At all times relevant hereto, Plaintiff Robert Best is an individual and resident of Cleburne County, Arkansas.

2. Defendant Consensus Orthopedics Inc. ("Consensus") is and at all times herein mentioned was a California corporation, or other business entity, organized and existing under the laws of the State of California with a principal place of business located at 1115 Winfield Way, Suite 100, El Dorado Hills, California 95762. Defendant Consensus may be served with process by personally serving Paul Rugg as registered agent for service of process at the above address or by serving the Arkansas Secretary of State pursuant to Arkansas Code Annotated § 16-58-120(2) (2019).

**This case assigned to District Judge Moody  
and to Magistrate Judge Deere**

3. John Does 1-10 are other persons or entities yet to be identified, who may be liable for the damages alleged herein for any reason, including, but not limited to, their involvement in the design, manufacture, sale, supply, distribution, marketing, inspection or maintenance of the subject knee implantation device.

4. This Court has personal jurisdiction over Defendants pursuant to Arkansas' Long Arm Statute § 16-58-120, as Defendants have conducted substantial or systemic business in Arkansas by designing, manufacturing, producing, making, marketing, distributing or selling the below-described product in Arkansas, including the one implanted in Plaintiff.

5. This Court has subject matter jurisdiction pursuant to 28 U.S.C. § 1332, because the parties are diverse and the amount in controversy exceeds \$75,000.

6. Venue is proper in this Judicial District pursuant to 28 U.S.C. § 1391, because a substantial part of the events or omissions giving rise to the claims occurred in this judicial district.

#### **ALLEGATIONS COMMON TO ALL COUNTS**

7. Defendants designed, fabricated, produced, compounded, processed, assembled, marketed, distributed and/or sold the Consensus Total Knee System Replacement prosthesis.

8. On March 7, 2014, Plaintiff underwent a right total knee replacement with surgical placement of a Consensus Knee System artificial knee joint.

9. On September 19, 2017, x-rays of Plaintiff's right knee showed that the polyethylene insert component of the Consensus implant had failed and become dislodged from the tibial tray.

10. On February 14, 2018, Plaintiff underwent a total revision right total knee arthroplasty.

11. Consensus Total Knee System Replacements and its components were defectively designed in that the polyethylene tibial insert would dislodge from the tibial tray causing dangerous instability, injury, and pain.

12. Additionally, the Consensus Total Knee System Replacements and its components were defectively manufactured in such a manner as to cause the tibial insert to bend, move, slip, and/or break.

13. As a result of the device's failure, Plaintiff has endured and incurred physical pain and suffering (both past, present, and future), emotional pain and suffering (both past, present, and future), permanent impairment and scarring, medical bills and expenses (both past, present, and future), loss of enjoyment of life, pre-judgment and post-judgment interest, statutory and discretionary costs, and any and all such other and further legal and equitable relief to which he is entitled.

## **CAUSES OF ACTION**

### **COUNT I – STRICT PRODUCT LIABILITY**

14. Plaintiff hereby incorporates all other paragraphs of this complaint as though fully set forth herein.

15. Plaintiff was damaged by the defective and unreasonably dangerous Consensus Total Knee System Replacement prosthesis, including having to undergo a revisions surgery less than 4 years after post-implant.

16. Defendants designed, fabricated, produced, compounded, processed, assembled, marketed, distributed and/or sold the Consensus Total Knee System Replacement prosthesis implanted in Plaintiff's right knee, by which he has been damaged.

17. The Consensus Total Knee System Replacements and related components were in a defective condition that rendered it unreasonably dangerous in that the polyethylene tibial insert,

tibial replacement components, and other components, would bend, move, slip, and/or break, and thus were defective and unreasonably dangerous to consumers, including Plaintiff.

18. The Consensus Total Knee System Replacements and its components were defective, unreasonably dangerous and unsafe for reasonably foreseeable use and consumption due to their design, manufacture and/or Defendants' failure to warn of dangers beyond that which would be contemplated by the ordinary and reasonable user, consumer, patient, and medical professional.

19. After recovery from the respective surgeries, and after regular use for their intended purpose, the Consensus Total Knee System Replacement and, subsequently, the Consensus revision knee replacement components failed, causing pain and forcing Plaintiff to undergo additional surgeries.

20. The defective nature of the devices was a direct and proximate cause of injury and damages to Plaintiff, including physical pain and suffering (both past, present, and future), emotional pain and suffering (both past, present, and future), permanent impairment and scarring, medical bills and expenses (both past, present, and future), loss of enjoyment of life, pre-judgment and post-judgment interest, statutory and discretionary costs, and any and all such other and further legal and equitable relief to which he is entitled.

#### **COUNT II – NEGLIGENCE**

21. Plaintiff hereby incorporates all other paragraphs of this complaint as though fully set forth herein.

22. Defendants owed a duty of reasonable care to the general public, including Plaintiff, when it designed, manufactured, assembled, inspected, tested, marketed, placed into the stream of commerce, and sold the Consensus Total Knee System Replacements to protect users from

unreasonable risk of harm when using the device for its intended purpose in a reasonably foreseeable manner.

23. Defendants breached this duty by designing, manufacturing, assembling, inspecting, testing, marketing, distributing, and selling the Consensus Total Knee System Replacements in a defective and unreasonably safe condition including, but not limited to, its foreseeably appreciated risk of harm of the tibial insert's propensity to dislodge from the tibial tray. A reasonably careful medical device manufacturer would not have acted in this manner.

24. Defendants knew or should have known that the subject products were defective and could fail and that use of the products involved an unreasonable, foreseeable danger, for which Defendants were required to warn.

25. Defendants' knowledge included, but was not limited to, knowledge that the polyethylene tibial insert, tibial replacement components, and other components, would bend, move, slip, and/or break, and thus were defective and dangerous to consumers, including Plaintiff.

26. Defendants had a duty to take steps to prevent any additional implantations of the defective devices and/or to warn patients who already had the devices implanted of the possibility of failure.

27. Defendants had a duty to warn Plaintiff of the dangers of the devices as the dangers were such that they were not generally known or not reasonably expected by a purchaser or user to find in said devices.

28. Defendants failed to provide adequate warning.

29. As a result of Defendants' failure to warn, the devices continued to be implanted in additional patients, including Plaintiff, and their sudden failure without warning caused Plaintiff to suffer injuries and losses.

30. Defendants' negligent design, manufacture, sale, and failure to warn were the direct and proximate causes of Plaintiff's injuries and damages, including physical pain and suffering (both past, present, and future), emotional pain and suffering (both past, present, and future), permanent impairment and scarring, medical bills and expenses (both past, present, and future), loss of enjoyment of life, pre-judgment and post-judgment interest, statutory and discretionary costs, and any and all such other and further legal and equitable relief to which he is entitled.

### **COUNT III – GROSS NEGLIGENCE**

31. Plaintiff hereby incorporates all other paragraphs of this complaint as though fully set forth herein.

32. Defendants knew or should have known, in light of the surrounding circumstances and past litigation, that the Consensus Total Knee System Replacements were defective and unreasonably dangerous and knew or should have known that their products and conduct would naturally and probably result in injury or damage to others, including that suffered and incurred by the Plaintiff. Defendants acted willfully, wantonly, and/or recklessly in disregard of the consequences; and therefore, in addition to his other damages as alleged herein, Plaintiff seeks exemplary and punitive damages against Defendants in an amount to be determined at trial.

### **COUNT IV – BREACH OF WARRANTY**

33. Prior to the time that Plaintiff used the products for their intended purpose, Defendants expressly and/or impliedly warranted to Plaintiff that the products were of merchantable quality, reasonably fit, and safe for their ordinary use and intended purpose.

34. At the time of contracting for sale and the retail sale of the subject Consensus Total Knee System Replacement prostheses and revision components, Defendants knew or had reason

to know the particular purpose for which the goods were required and that Plaintiff was relying on Defendants' skill and judgment to select and furnish suitable goods.

35. In a reasonable and foreseeable manner, Plaintiff relied on Defendants' express and implied representations and warranties in consenting to knee implant surgery using Defendants' devices.

36. Defendants' breached their express and implied representations and warranties regarding the safety and merchantability and fitness for particular purpose of their devices.

37. The subject devices were not safe, not fit for their intended use, nor of merchantable quality as warranted by Defendants.

38. Defendants knew or had reason to know that the devices were not safe, not fit for their intended use, nor of merchantable quality as warranted by Defendants.

39. On information and belief, Defendants' knowledge included, but was not limited to, knowledge that the polyethylene tibial insert, tibial replacement components, and other components, would bend, move, slip, and/or break, and thus were defective and dangerous to consumers, including Plaintiff.

40. As a direct and proximate result of Defendants' breaches of warranty, Plaintiff has suffered injuries, losses and damages including physical pain and suffering (both past, present, and future), emotional pain and suffering (both past, present, and future), permanent impairment and scarring, medical bills and expenses (both past, present, and future), loss of enjoyment of life, pre-judgment and post-judgment interest, statutory and discretionary costs, and any and all such other and further legal and equitable relief to which he is entitled.

**PRAYER FOR RELIEF**

WHEREFORE, ALL PREMISES CONSIDERED, Plaintiff requests the Court grant judgment against Defendants Consensus Orthopedics, Inc. and John Does 1-10, jointly and severally, for COMPENSATORY DAMAGES and PUNITIVE DAMAGES as alleged herein, in an amount considered fair and reasonable by a jury, and for all such further relief, both general and specific, to which the Court deems proper, just, and equitable under the circumstances. Plaintiff also seeks general relief.

**JURY DEMAND**

Plaintiff requests a jury trial of all matters appropriately tried to a jury.

DATED this 11 day of June 2019.

Respectfully submitted,

/s/ Richard Underwood

RICHARD UNDERWOOD, ARB 2006137

**FARRIS BOBANGO & BRANAN, PLC**

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*Counsel for Plaintiff Robert Best*



CIVIL COVER SHEET 1:19-cv-46-JM

The JS 44 civil cover sheet and the information contained herein neither replace nor supplement the filing and service of pleadings or other papers as required by law, except as provided by local rules of court. This form, approved by the Judicial Conference of the United States in September 1974, is required for the use of the Clerk of Court for the purpose of initiating the civil docket sheet. (SEE INSTRUCTIONS ON NEXT PAGE OF THIS FORM.)

PLAINTIFFS

(b) County of Residence of First Listed Plaintiff Cleburne (EXCEPT IN U.S. PLAINTIFF CASES)

(c) Attorneys (Firm Name, Address, and Telephone Number) Richard Underwood, 999 S Shady Grove Road, Ste 500, Memphis, TN 38120, 901.259.7100

DEFENDANTS

CONSENSUS ORTHOPEDICS, INC., and JOHN DOES 1-10

County of Residence of First Listed Defendant (IN U.S. PLAINTIFF CASES ONLY)

NOTE: IN LAND CONDEMNATION CASES, USE THE LOCATION OF THE TRACT OF LAND INVOLVED.

Attorneys (If Known)

II. BASIS OF JURISDICTION (Place an "X" in One Box Only)

- 1 U.S. Government Plaintiff, 2 U.S. Government Defendant, 3 Federal Question (U.S. Government Not a Party), 4 Diversity (Indicate Citizenship of Parties in Item III)

III. CITIZENSHIP OF PRINCIPAL PARTIES (Place an "X" in One Box for Plaintiff and One Box for Defendant)

- Citizen of This State, Citizen of Another State, Citizen or Subject of a Foreign Country, PTF DEF, Incorporated or Principal Place of Business In This State, Incorporated and Principal Place of Business In Another State, Foreign Nation

IV. NATURE OF SUIT (Place an "X" in One Box Only)

Click here for: Nature of Suit Code Descriptions.

Table with 5 columns: CONTRACT, REAL PROPERTY, TORTS, CIVIL RIGHTS, PRISONER PETITIONS, FORFEITURE/PENALTY, LABOR, IMMIGRATION, BANKRUPTCY, SOCIAL SECURITY, FEDERAL TAX SUITS, OTHER STATUTES. Includes various legal codes and descriptions.

V. ORIGIN (Place an "X" in One Box Only)

- 1 Original Proceeding, 2 Removed from State Court, 3 Remanded from Appellate Court, 4 Reinstated or Reopened, 5 Transferred from Another District (specify), 6 Multidistrict Litigation - Transfer, 8 Multidistrict Litigation - Direct File

VI. CAUSE OF ACTION

Cite the U.S. Civil Statute under which you are filing (Do not cite jurisdictional statutes unless diversity): 28 USC 1332

Brief description of cause: Products Liability- medical device

VII. REQUESTED IN COMPLAINT:

CHECK IF THIS IS A CLASS ACTION UNDER RULE 23, F.R.Cv.P. DEMAND \$ CHECK YES only if demanded in complaint: JURY DEMAND: X Yes O No

VIII. RELATED CASE(S) IF ANY

(See instructions):

JUDGE

DOCKET NUMBER

DATE

SIGNATURE OF ATTORNEY OF RECORD

FOR OFFICE USE ONLY

RECEIPT # AMOUNT APPLYING IFP JUDGE MAG. JUDGE