

UNITED STATES DISTRICT COURT
NORTHERN DISTRICT OF OHIO
WESTERN DIVISION

JESSICA FARSON)	
)	
<i>Plaintiff</i>)	
)	
v.)	Case No.
)	
COOPERSURGICAL, INC.,)	
THE COOPER COMPANIES, INC.,)	
FEMCARE, LTD. – UK SUBSIDIARY OF)	
UTAH MEDICAL PRODUCTS, INC., and)	
UTAH MEDICAL PRODUCTS, INC.)	
)	
<i>Defendants</i>)	

PLAINTIFF’S COMPLAINT AND JURY DEMAND

NOW COMES Plaintiff, Jessica Farson (hereinafter “Plaintiff” and/or “Ms. Farson”) by and through her counsel, Griffin Purnell LLC and The Henry Law Firm and for her cause of action against Defendants CooperSurgical, Inc., The Cooper Companies, Inc., Femcare, Ltd. – UK subsidiary of Utah Medical Products, Inc., and Utah Medical Products, Inc. (collectively hereinafter “Defendants”), all jointly and severally, as the companies and/or successors in interest to the companies that designed, developed, manufactured, tested, labeled, packaged, distributed, marketed and/or sold the Filshie Clip medical device that was surgically used in Plaintiff and others throughout the United States and the world. Accordingly, Plaintiff alleges and states to the Court as follows:

I. INTRODUCTION

1. Plaintiff brings this civil action to recover damages within the jurisdictional limits of this Court including all (1) General Damages; (2) Special Damages; and (3) Punitive Damages as well as all other damages allowable under Ohio law as a result of the use, design, manufacture,

surveillance, sale, marketing, advertising, promotion, labeling, packaging, and distribution of Filshie Clips.

2. Plaintiff brings claims fully set forth below asserting: (1) design defect; (2) manufacturing defect; (3) strict liability; (4) negligence; (5) gross negligence; and (6) punitive damages.

3. This claim arises from Ms. Farson's Filshie Clip tubal ligation procedure which, because of Defendants' actions and omissions, resulted in a series of damages.

II. PARTIES

4. Plaintiff, Jessica Farson lives in Fremont, Ohio, and is subject to the jurisdiction of this Court, and is deemed to be a resident of the State of Ohio for purposes of venue and jurisdiction.

5. Defendant, The Cooper Companies, Inc. ("Cooper Companies") is a Delaware corporation with its principal place of business located at 6101 Bollinger Canyon Road, in San Ramon, California. For diversity of citizenship purposes, Defendant Cooper Companies, Inc. is a citizen of both Delaware and California. Cooper Companies, Inc. may be served with process by serving its registered agent at 251 Little Falls Drive, Wilmington, DE 19808.

6. Defendant CooperSurgical, Inc. ("CooperSurgical") is a Delaware corporation with its principal place of business located at 95 Corporate Drive in Trumbull, Connecticut. CooperSurgical may be served with process by serving its registered agent at CooperSurgical, Inc., 95 Corporate Drive, Trumbull, CT 06611.

7. Defendant Femcare, Ltd. is a UK subsidiary of Utah Medical Products, Inc. with its principal place of business located at 32 Premier Way, Romsey, Hampshire SO51 9DQ, United Kingdom. Femcare, Ltd. – UK Subsidiary of Utah Medical Products, Inc. may be served with

process by serving its registered agent Karen Elizabeth Glasbey, FemcareUK, 32 Premier Way, Romsey, Hampshire, United Kingdom SO519DQ.

8. Defendant Utah Medical Products, Inc. is the parent company of Femcare, Ltd. with its principal place of business located at 7043 South 300 West, Midvale, Utah 84047-1048 and may be served with process by serving its registered agent Ben Shirley at 7043 South 300 West, Midvale, UT 84047.

9. CooperSurgical is a subsidiary of Defendant Cooper Companies, Inc. Defendant CooperSurgical is a citizen of both Delaware and Connecticut for diversity of citizenship purposes. Cooper Companies, Inc. and CooperSurgical, Inc. are referred to collectively hereinafter as “CooperSurgical.”

10. Femcare, Ltd. is a UK subsidiary of Utah Medical Products, Inc., and a citizen of England for diversity of citizenship purposes. Utah Medical Products, Inc. is a citizen of Utah for diversity of citizenship purposes.

11. All acts and omissions of the Defendants as described herein were done by its agents, servants, employees and/or owners, acting in the course and scope of its respective agencies, services, employments and/or ownership.

III. JURISDICTION & VENUE

12. This Court has subject matter original jurisdiction through diversity of citizenship pursuant to 28 U.S.C. §1332(a) because the Plaintiff is a citizen of Ohio, the named Defendants are citizens of different states and the amount in controversy exceeds the sum of value of \$75,000.00, exclusive of interest and costs.

13. This Court has specific jurisdiction over these Defendants because they purposefully availed themselves of the privilege of conducting business in the state of Ohio and

established minimum contacts sufficient to confer jurisdiction over these Defendants, and the assumption of jurisdiction over Defendants will not offend traditional notions of fair play and substantial justice and is consistent with constitutional requirements of due process.

14. CooperSurgical, Femcare, Ltd., and Utah Medical Products sell their products and intend that they be used by medical professionals treating patients in Ohio.

15. At all times relevant hereto and alleged herein, the Defendants conducted and continue to regularly conduct substantial business within the state of Ohio which included and continues to include, the research, safety surveillance, manufacture, sale, distribution and/or marketing of Filshie Clips which are distributed through the stream of interstate and intrastate commerce in the state of Ohio, and within the Northern District of Ohio.

16. Venue is proper in this District pursuant to 28 U.S.C. § 1391(b) and 18 U.S.C. §1965 (a) because a substantial part of the events or omissions giving rise to the claim occurred in this District and each Defendant transacts business affairs and conducts activity that gave rise to the claim of relief in this District.

IV. FACTUAL BACKGROUND

a. *Plaintiff Brings this Action Because Filshie Clips Injured her after migration.*

17. Plaintiff in this action seeks compensation for injuries she sustained in connection from the use of Filshie Clips, a medical device used in tubal ligations.

18. This action is brought by Plaintiff. Ms. Farson was implanted with a female birth control device known as a Filshie Clip. In short, this device is intended to cause bilateral occlusion (blockage) of the fallopian tubes by applying a clip onto the fallopian tubes which then anchors and elicits tissue growth, theoretically causing a closure of the tubes. However, in reality, the clips migrate from the tubes wreaking havoc on the female body.

b. What is a Filshie Clip and How is it Supposed to Work?

19. Filshie Clips are part of the “Filshie Clip system” for laparoscopic tubal ligation which involves applying a titanium clip with silicone rubber lining around each of the fallopian tubes.

20. The Filshie Clip works by exerting continued pressure on the fallopian tube, causing avascularization for the 3 to 5 mm area it encompasses. The silicone continues this pressure even after necrosis starts and the fallopian tube decreases in size. Fibrosis then occurs, and the clip is peritonealized if all goes as planned.

21. Defendants’ disposable delivery system consists of an applicator which allows insertion into the women’s body to allow the clip to be snapped onto the fallopian tube.

22. A women’s choice of birth control is a deeply personal decision, particularly when choosing a long-acting form of birth control like a tubal ligation which should permanently alter a women’s body.

c. Background on Filshie Clips and the FDA Process.

23. Femcare, the manufacturer of the Filshie Clip, obtained Conditional Premarket Approval (PMA) by the Food and Drug Administration (FDA). The Defendants’ failure to conform with the FDA requirements prescribed in the PMA and violations of relevant state and federal law form the basis of this lawsuit.

24. Class III medical devices are those that either “present a potential unreasonable risk of illness or injury or are for a use in supporting or sustaining human life or for a use which is of substantial importance in preventing impairment of human health.” 21 U.S.C. § 360(c)(1)(c)

25. Because Filshie Clips are classified as a Class III medical device the FDA evaluated Filshie Clips’ safety and effectiveness prior to granting the product Conditional PMA in 1996.

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