

Ernesteen JONES, Plaintiff-Appellant,  
v.  
**NOVARTIS PHARMACEUTICALS  
COMPANY**, A corporation, Defendant,  
Novartis Pharmaceuticals  
Corporation, Defendant-Appellee.

United States Court of Appeals, Eleventh Circuit.  
April 30, 2018  
--- Fed.Appx. ----  
2018 WL 2015488

Appeal from the United States District Court for the  
Northern District of Alabama, D.C. Docket No. 2:13-  
cv-00624-VEH

#### Attorneys and Law Firms

**Tammy Smith**, Taylor & Taylor, Birmingham, AL, for  
Plaintiff-Appellant

**Frederick G. Helmsing, Jr., Edward S. Sledge, III,**  
McDowell Knight Roedder & Sledge, LLC, Mobile,  
AL, **Robert E. Johnston, Stephen A. Klein, Andrew  
L. Reissaus, Tamara F. Barago**, Hollingsworth LLP,  
Washington, D.C., for Defendant-Appellee

Before **WILSON, JORDAN**, and **HIGGINBOTHAM**, \*  
Circuit Judges.

\* Honorable Patrick E. Higginbotham, United States  
Circuit Judge for the Fifth Circuit, sitting by  
designation.

#### Opinion

PER CURIAM:

\*1 Ernesteen Jones appeals the district court’s exclusion  
of testimony (either in whole or in part) offered by  
her four experts, as well as the district court’s grant of  
summary judgment in favor of Novartis Pharmaceuticals  
Corporation. After careful review of the record, the  
parties’ briefs, and with the benefit of oral argument, we  
find no reversible error and, accordingly, affirm.

Ms. Jones offered the testimony of Dr. William Banks  
Hinshaw, who opined that general causation was  
established between the medication **Reclast** and atypical

femur fractures. “General causation refers to the ‘general  
issue of whether a substance has the potential to cause the  
plaintiff’s injury.’” *Chapman v. Procter & Gamble Distrib.,  
LLC*, 766 F.3d 1296, 1306 (11th Cir. 2014) (quoting *Guinn  
v. AstraZeneca Pharms. LP*, 602 F.3d 1245, 1248 n.1 (11th  
Cir. 2010)).

The district court determined that Dr. Hinshaw, although  
qualified, employed unreliable methodologies in reaching  
that conclusion and excluded his testimony in full.  
*See Fed. R. Evid. 702(c)* (requiring admissible expert  
testimony to be “the product of reliable principles and  
methods”); *Daubert v. Merrell Dow Pharms., Inc.*, 509  
U.S. 579, 592–93, 113 S.Ct. 2786, 125 L.Ed.2d 469  
(1993) (discussing factors in evaluating reliability of a  
methodology). “[W]e must affirm [this conclusion] unless  
we at least determine that the district court has made  
a clear error of judgment, or has applied an incorrect  
legal standard.” *McClain v. Metabolife Intern., Inc.*,  
401 F.3d 1233, 1238 (11th Cir. 2005) (quotation marks  
omitted). Nothing in our review of the record, including  
Dr. Hinshaw’s deposition testimony, expert reports, and  
supporting exhibits, leads us to believe that the district  
court committed a “clear error of judgment,” *id.*, or  
that its decision was “manifestly erroneous,” *Rink v.  
Cheminova, Inc.*, 400 F.3d 1286, 1291 (11th Cir. 2005), so  
we affirm the district court’s exclusion of Dr. Hinshaw’s  
general causation opinions.

We agree with the district court that this case falls  
within *McClain’s* second category and that, therefore,  
Ms. Jones was required to offer admissible testimony  
on general causation. *See McClain*, 401 F.3d at 1239.  
She conceded as much at oral argument, stating that  
she “need[s] Dr. Hinshaw.” Because the district court  
did not abuse its discretion in excluding Dr. Hinshaw’s  
general causation opinions, summary judgment in favor  
of Novartis was appropriate. *See Chapman*, 766 F.3d  
at 1316 (noting that the plaintiff was “required to have  
*Daubert*-qualified, general and specific-causation-expert  
testimony that would be admissible at trial to avoid  
summary judgment”) (emphasis in original).

Having concluded that summary judgment was  
appropriate due to the exclusion of Dr. Hinshaw, we need  
not analyze whether the district court erred in excluding  
or limiting the testimony of Ms. Jones’ remaining three  
experts.

**AFFIRMED.**

**All Citations**

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