

IN THE UNITED STATES DISTRICT COURT
FOR THE MIDDLE DISTRICT OF NORTH CAROLINA

JOHN AND JANE DOE 2,)	
INDIVIDUALLY, and as GUARDIANS)	
AD LITEM, OF MINOR CHILD DOE 2,)	
)	
Plaintiffs,)	
v.)	1:03CV00669
)	
ORTHO-CLINICAL)	
DIAGNOSTICS, INC.,)	
)	
Defendant.)	

MEMORANDUM OPINION

BEATY, District Judge.

Plaintiffs John and Jane Doe 2 (“Plaintiffs”) have initiated this lawsuit based upon their contention that the thimerosal in Defendant Ortho-Clinical Diagnostics, Inc.’s (“Ortho-Clinical” or “Defendant”) biologic product RhoGAM caused their child’s autism. This matter is presently before the Court on three motions: Defendant’s Motion to Exclude All Testimony that Thimerosal-Containing RhoGAM Causes Autism [Document #63], Defendant’s Motion to Exclude Plaintiffs’ Expert Suzanne Parisian, M.D. [Document #65], and relatedly, Defendant’s Motion for Summary Judgment [Document #94]. The Court heard testimony concerning Defendant’s motions to exclude witnesses for three days at the end of May 2006. Based upon the testimony at that hearing, the Court will grant Defendant’s Motion to Exclude All Testimony that Thimerosal-Containing RhoGAM Causes Autism. More specifically, the focus of the

Court's present Memorandum Opinion is the testimony of Plaintiffs' expert witness, Dr. Mark Geier. Dr. Geier was the only expert offered in this case by Plaintiffs who is designated to testify as to both general *and* specific causation. For the reasons given by the Court herein, Dr. Geier's testimony is specifically being excluded pursuant to Defendant's Motion to Exclude. As such, without Dr. Geier's testimony, Plaintiffs are unable to meet their burden to demonstrate that the thimerosal in Defendant's RhoGAM product caused Plaintiff Minor Child Doe 2's autism, a result that leads directly to the failure of all of Plaintiffs' claims. Accordingly, for the reasons detailed below, the Court will also grant Defendant's Motion for Summary Judgment.

I. FACTUAL BACKGROUND¹

Plaintiffs allege that Minor Child Doe 2 ("Minor Child Doe") has suffered severe neurodevelopmental disorders and permanent injuries from exposure to toxic levels of mercury. Plaintiffs claim that this mercury exposure resulted from one single shot of RhoGAM that Jane Doe received while 28-weeks pregnant and another shot of RhoGAM that Jane Doe received shortly after Minor Child Doe's birth.² Plaintiffs argue that this limited amount of thimerosal,

¹ The Court in this Memorandum Opinion will refer to some scholarly works and other documents that were attached to Defendant's Brief in Support of the Motion to Exclude All Testimony that Thimerosal-Containing RhoGAM Causes Autism. For ease in citing to these articles, the Court will refer to the Appendix filed by Defendants and located in the record as Documents ##67-70. The format for such references will be "Def. Apx. Tab #, name of article, p.#." In other instances, the Court will refer to articles discussed by either Plaintiffs' or Defendant's experts during the hearing, and will cite those articles as appropriate.

² RhoGAM is used to suppress the immune response of Rh negative women to Rh positive red blood cells. This treatment is used whenever it is suspected that fetal red blood cells have entered the circulation of an Rh negative mother, unless either the fetus or the father is

which contains a mercury derivative, in both of those shots given to his mother caused Minor Child Doe to develop autism approximately sixteen months after his birth. Based upon these allegations, Plaintiffs' several claims against Defendant consist of the following: (1) negligence; (2) negligent failure to warn; (3) breach of express warranty; (4) breach of implied warranty; (5) negligent misrepresentation; (6) intentional misrepresentation and fraud; (7) unfair and deceptive trade practices; (8) inadequate design; (9) negligent infliction of emotional distress; (10) gross negligence; (11) loss of consortium; and (12) punitive damages. It is significant to the Court's ultimate disposition of this matter that the viability of all of the listed claims hinge on Plaintiffs' ability to prove that the thimerosal in RhoGAM caused Minor Child Doe's autism.

In order to prove their claims, Plaintiffs designated three experts on the question of whether thimerosal could cause autism: Dr. Mark Geier ("Geier"), Boyd Haley, Ph.D. ("Haley") and George Lucier, Ph.D. ("Lucier"). As a separate matter, Plaintiffs have designated a fourth expert, Dr. Suzanne Parisian, to testify about the FDA regulatory process as it relates to claims regarding negligent failure to warn and inadequate design. The Court conducted a Daubert hearing in this matter that included the testimony of Dr. Geier, as well as testimony from several of Defendant's experts who asserted that Dr. Geier's methodology that supports his proffered opinion on the causal connection, both general and specific causation, as it relates to the autism

shown to be Rh negative. Without such treatment, the immune response of the mother could cause Hemolytic Disease of the Newborn, which in turn may lead to perinatal injury or death.

of Minor Child Doe, was flawed, as well as his conclusions in that regard.³

II. THE DAUBERT STANDARD

The nature and necessity of a Daubert hearing is derived from the case of Daubert v. Merrell Dow Pharms., 509 U.S. 579, 113 S. Ct. 2786 (1993). Under Daubert, this Court must rule on the admissibility of expert scientific testimony. See id. at 598, 113 S. Ct. at 2799. Daubert requires a two-part analysis: first, this Court must determine whether an expert's testimony reflects "scientific knowledge," whether the findings are "derived by the scientific

³ At the hearing, Plaintiffs also made available expert reports from their other two experts, Dr. Haley and Dr. Lucier, as to the general causation question. The Court has appropriately taken this information into account, but nevertheless finds that Dr. Haley's report does not state an expert opinion that thimerosal causes autism, rather just that he has a *theory*, (see Def. Apx. Tab 18, Dep. of Boyd Haley, at 190), about how such a thing could happen. At best, he expressed "strong belief" that the cause of "neurodevelopmental disorders in infants" is exposure to an organic-mercury compound such as thimerosal. (See Def. Apx. Tab. 14, Haley Expert Report, p. 5.) Additionally, Plaintiffs proffered the report of Dr. Lucier, who is an expert in *methylmercury* and not *ethylmercury*, which is the substance in RhoGAM. Dr. Lucier does not offer an opinion that methylmercury causes autism, but rather that it may cause "developmental disorders." Significantly, the Court notes that neither Dr. Haley nor Dr. Lucier asserts that he is an expert on autism nor are they offered as such. In any event, the Court finds that neither of the proffered reports of Dr. Haley nor Dr. Lucier are sufficiently reliable under Daubert on the general causation issue because neither is relevant to the "task at hand." It would be an unacceptable scientific leap to suggest that they serve as proof, by a preponderance of the evidence, of Plaintiff's claim that the thimerosal in RhoGAM can cause autism. See Dunn v. Sandoz Pharms. Corp., 275 F. Supp. 2d 672, 684 (M.D.N.C. 2003) ("While hypothesis is essential in the scientific community because it leads to advances in science, speculation in the courtroom cannot aid the fact finder in making a determination of whether liability exists. Ultimately, speculation is unreliable evidence and is inadmissible."). Accordingly, the majority of the Court's opinion is focused only on the testimony of Dr. Geier, Plaintiffs' primary general causation expert and single specific causation expert. Nevertheless, it is the Court's finding that the discussion and analysis herein regarding the relevant scientific literature applies equally to the proffered reports of Drs. Haley and Lucier, which also purport to survey the relevant literature in coming to their respective conclusions.

method,” and whether the work product is “good science.” Id. at 590 & 593, 113 S. Ct. at 2795 & 2797. Second, this Court must determine whether the expert’s testimony is “relevant to the task at hand.” Id. at 597, 113 S. Ct. at 2799. This gatekeeping function is important because, “due to the difficulty of evaluating their testimony, expert witnesses have the potential to be both powerful and quite misleading.” Westberry v. Gislaved Gummi AB, 178 F.3d 257, 261 (4th Cir. 1999) (quoting Daubert, 509 U.S. at 595, 113 S. Ct. 2786) (internal quotation marks omitted).

In Daubert and related cases, the U.S. Supreme Court has elucidated a number of factors for District Courts to consider when determining whether to admit expert testimony under Federal Rule of Evidence 702.⁴ For example, the U.S. Supreme Court stated in Daubert that courts may consider whether the theory or technique employed by the expert is generally accepted in the scientific community; whether it has been subjected to peer review and publication; whether it can be and has been tested; whether the known or potential rate of error is acceptable; and the existence and maintenance of standards and controls. Id. at 593-95, 113 S. Ct. at 2796-98. These factors are not exclusive nor dispositive. Since Daubert, the U.S. Supreme Court and lower courts have also identified additional factors that may be considered, such as whether an expert has unjustifiably extrapolated from an accepted premise to an unfounded

⁴ Rule 702 of the Federal Rules of Evidence provides that “if scientific, technical, or other specialized knowledge will assist the trier of fact to understand the evidence or to determine a fact in issue, a witness qualified as an expert by knowledge, skill, experience, training, or education, may testify thereto in the form of an opinion or otherwise, if (1) the testimony is based upon sufficient facts or data, (2) the testimony is the product of reliable principles and methods, and (3) the witness has applied the principles and methods reliably to the facts of the case.”

conclusion, see GE v. Joiner, 522 U.S. 136, 146, 118 S. Ct. 512, 519 (1997), whether an expert has adequately accounted for obvious alternative explanations, see Claar v. Burlington N. R.R., 29 F.3d 499, 502 (9th Cir. 1994), or whether an expert is proposing to testify about matters “growing naturally and directly out of research they have conducted independent of the litigation, or whether they have developed their opinions expressly for purposes of testifying.” Daubert v. Merrell Dow Pharms., 43 F.3d 1311, 1317 (9th Cir. 1995) (hereinafter, Daubert II).

In this case, much, but not all, of Dr. Geier’s proposed testimony is not based upon his own research, but instead upon a review of the relevant literature. Where proffered expert testimony is not based on independent research, but instead on such a literature review, the party proffering such testimony must “come forward with other objective, verifiable evidence that the testimony is based on ‘scientifically valid principles.’ One means of showing this is by proof that the research and analysis supporting the proffered conclusions have been subjected to normal scientific scrutiny through peer review and publication.” Daubert II, 43 F.3d at 1318. Thus, the research Dr. Geier relied upon must itself be able to meet the Daubert test. The fact that a journal is peer-reviewed is a significant consideration. Id.

While Daubert itself focused on an expert’s methodology, the Court notes that later decisions have gone beyond methodology in certain instances. While in Daubert the U.S. Supreme Court stated that a Daubert analysis must “focus . . . solely on principles and methodology, not on the conclusions that they generate,” the Court later recognized that “conclusions and methodology are not entirely distinct from one another.” GE, 522 U.S. at 146,

118 S. Ct. at 519. Put another way – “Trained experts commonly extrapolate from existing data. But nothing in either Daubert or the Federal Rules of Evidence requires a district court to admit opinion evidence which is connected to existing data only by the *ipse dixit* of the expert. A court may conclude that there is simply too great an analytical gap between the data and the opinion proffered.” Id. Finally, a bold statement of the experts’ qualifications, conclusions, and assurances of reliability are not enough to satisfy the Daubert standard. See Daubert II, 43 F.3d at 1319.

In addition to a consideration of the Daubert analysis, the Court notes that it must also distinguish in this case between Dr. Geier’s proffered testimony as to both “general causation” and “specific causation.” See, e.g., Dunn v. Sandoz Pharms. Corp., 275 F. Supp. 2d 672, 676 (M.D.N.C. 2003) (citing Reference Manual on Scientific Evidence 444 (2d ed. 2000)). On the one hand, “[g]eneral causation ‘is established by demonstrating . . . that exposure to a substance can cause a particular disease.’” Id. “Specific, ‘or individual causation, however[,] is established by demonstrating that a given exposure is the cause’ of a particular individual’s disease.” Id. Where a “plaintiff is not able to establish general causation, it is unnecessary to consider whether the plaintiff can establish specific causation.” Id.; see also Raynor v. Merrell Pharms., 104 F.3d 1371, 1376 (D.C. Cir. 1997).

With these legal standards in mind, it was the Court’s undertaking to determine whether Plaintiffs’ evidence satisfied their burden of proof to show that their experts used proper scientific methodology in reaching their ultimate conclusion that Minor Child Doe’s autism was

caused by the thimerosal in Defendant's product RhoGAM. At the close of Plaintiffs' presentation at the Daubert hearing, Plaintiffs argued that their evidence would support such a conclusion. In response to Plaintiffs' position, Defendant challenged Plaintiffs' proffer by way of a cross examination of Plaintiffs' expert Dr. Geier and by offering its own experts to demonstrate that Plaintiffs' experts used unsound methodology or otherwise failed to follow sound protocol. Having closely considered the evidence and arguments both by Plaintiffs and Defendant, the Court has made a number of findings with respect to the testimony by Plaintiffs' primary expert Dr. Geier. These findings form the basis of the Court's ultimate conclusion that Plaintiffs have not met their burden under the Daubert analysis.

III. ANALYSIS OF DAUBERT AS APPLIED TO DR. MARK GEIER

As initial background information with respect to the qualifications of Dr. Geier, the Court notes that he is the president of his own company, The Genetic Centers of America. He is a medical doctor who specializes in obstetrical genetics with a Ph.D. as well in genetics. He is board certified in medical genetics and forensic medicine. However, it is significant to the Court that he is not board certified in pediatrics or in pediatric neurology, nor is he certified as an epidemiologist or biostatistician. Dr. Geier did serve as a researcher at the National Institutes of Health for 10 years and worked as a professor at John Hopkins University. While he has published more than 50 peer-reviewed medical papers, none of these prior publications were on the specific issue at hand, that is, whether RhoGAM with thimerosal causes autism. The Court has taken into account, as well, the fact that Dr. Geier has testified as an expert witness in about

one hundred cases before the National Vaccine Injury Compensation Program of the United States Court of Federal Claims. It is noteworthy that in more than ten of these cases, particularly in some of the more recent cases, Dr. Geier's opinion testimony has either been excluded or accorded little or no weight based upon a determination that he was testifying beyond his expertise.⁵ In this case, subject to the Court's Daubert analysis, Dr. Geier's testimony is being offered by Plaintiffs for presentation at trial to support Dr. Geier's ultimate conclusion that the

⁵ See, e.g., Piscopo v. Sec'y of HHS, 66 Fed. Cl. 49, 55 (May 26, 2005) (approving Dr. Geier's exclusion under Daubert because his training is in genetics and obstetrics, which is "largely irrelevant to the expertise needed to establish a causal relationship between the Hepatitis B vaccine and the petitioner's autoimmune disorder"); Weiss v. Sec'y of HHS, No. 03-190V, 2003 WL 22853059 (Fed. Cl. Oct. 9, 2003) (finding Dr. Geier is a "professional witness in areas for which he has no training, expertise, and experience"); Thompson v. Sec'y of HHS, No. 99-436V, 2003 WL 21439672 (Fed. Cl. May 23, 2003) (finding Dr. Geier was not qualified because his causation theory was filled with "speculation that is directly contrary to the conclusions reached in well-respected and numerous epidemiologic and medical studies ranging over two decades"); Bruesewitz v. Sec'y of HHS, No. 95-0266V, 2002 WL 31965744 (Fed. Cl. Dec. 20, 2002) (finding Dr. Geier unqualified to diagnose neurological diseases); Raj v. Sec'y of HHS, No. 96-294V, 2001 WL 963984 (Fed. Cl. July 31, 2001) (finding Dr. Geier "wholly unqualified to testify concerning the two major issues in this case [encephalopathy and infantile spasms] . . . because he is neither board certified nor has formal training in pediatrics and pediatric neurology"); Haim v. Sec'y of HHS, No. 90-1031V, 1993 WL 346392 (Fed. Cl. Aug. 27, 1993) (finding that Dr. Geier's testimony did not reach "the level of evidentiary reliability that Daubert requires because it is not based upon scientific validity, valid methodology, peer review or testing, and more than minimal support within the scientific community"); Marascalco v. Sec'y of HHS, No. 90-1571V, 1993 WL 277095 (Fed. Cl. July 9, 1993) (finding Dr. Geier's testimony "intellectually dishonest" and that his affidavit was "nothing more than an egregious example of blatant, result-oriented testimony").

Plaintiffs point to a number of cases in which Dr. Geier was apparently able to give his opinion as to causation in vaccine cases. However, the Court finds that the majority of cases cited in Plaintiffs' Response concern testimony by Dr. Geier on vaccine issues prior to 1995. Plaintiffs, however, do not assert that any of these cases or any previous testimony by Dr. Geier concern or even address the issue of whether thimerosal in RhoGAM causes autism.

thimerosal in RhoGAM caused Minor Child Doe's autism. However, appropriately under Daubert, the Court's focus, principally, is on the acceptability of the methodology by which Dr. Geier reached his ultimate conclusion. The specific methodology at issue here is Dr. Geier's method of using (1) a review of the relevant literature and his own studies as related to the incidence of autism, so as to support a general causation theory, and furthermore, (2) his use of a differential diagnosis of Minor Child Doe, so as to establish specific causation with respect to Minor Child Doe's own condition. The Court will address each of these areas in turn.

A. Dr. Geier's Review of Literature Concerning Whether RhoGAM Causes Autism

In examining Dr. Geier's methodology, the Court notes that, in fact, a literature review can be an appropriate part of a method of determining general causation. See, e.g. Benedi v. McNeil-P.P.C., Inc., 66 F.3d 1378, 1384 (4th Cir. 1995) (allowing an expert opinion based upon patient history, examination, lab and pathology data, and a study of the peer-reviewed literature). However, a literature review must still be performed appropriately. As revealed by his testimony at the Daubert hearing, Dr. Geier, however, relied upon a number of disparate and unconnected studies, including the findings of Dr. Haley and Dr. Lucier, to reach a piecemeal conclusion with respect to general causation that the small amount of thimerosal received in this case by the mother of Minor Child Doe during the course of her pregnancy and shortly after the child's birth, could cause autism. Dr. Geier's methodology consisted of attempting to connect various individual studies that had developed the existence of certain findings such as the

