

**IN THE COURT OF COMMON PLEAS
OF PHILADELPHIA COUNTY, PENNSYLVANIA**

**FIRST JUDICIAL DISTRICT OF PENNSYLVANIA
TRIAL DIVISION - CIVIL**

IN RE	:	
	:	
YAZ®, YASMIN®, OCELLA®	:	SEPTEMBER TERM, 2009
LITIGATION	:	NO. 01307
	:	

DISCOVERY MASTER ORDER NO. 3


AND NOW, to wit, this 21st day of October, 2010, following a conference with counsel on October 18, 2010, the Court-appointed Discovery Master, Harris T. Bock, Esquire, hereby **ORDERS** as follows:

1. **Production of SAS Data from GIAD Database:** The Bayer Defendants shall produce SAS data from the studies of Yaz, Yasmin, and Ocella in the GIAD database by the close of business on November 17, 2010.
2. **Production of SAS Data from TOSCA Database:** The Bayer Defendants will provide a report to the Discovery Master on the status of producing SAS data for YAZ, Yasmin, and Ocella studies from the TOSCA database by the close of business on November 8, 2010.
3. **Production from STEEPROCK Database:** The Bayer Defendants will produce an export of the Yaz and the Yasmin data from the STEEPROCK database by the close of business on October 31, 2010.
4. **Redacted Documents:** By agreement of liaison counsel, Case Management Order No. 4, Section I(B)(2)(d) dated December 11, 2009, is hereby amended so

that it is consistent with the Orders dated September 22, 2010 and October 15, 2010 entered by the Honorable David Herndon who is presiding over the “MDL” litigation involving Yaz, Yasmin and Ocella. Judge Herndon’s September 22, 2010 and October 15, 2010 Orders (the “MDL Orders”) are attached hereto and their provisions and requirements as amended or modified, are adopted in this case. Consistent with the requirements and provisions of the MDL Orders, the Bayer Defendants shall produce to the Pennsylvania Plaintiffs the same and identical un-redacted documents that the Bayer Defendants produce in the MDL Litigation pursuant to the MDL Orders attached hereto, and any amendments or modifications to the MDL Orders.

5. **Next Discovery Master Conference:** The next Discovery Master Conference shall be held on Monday, November 15, 2010 at 10:00 a.m. at the offices of The Dispute Resolution Institute. As soon as practicable, counsel shall submit to the Discovery Master the agenda for the next Discovery Master Conference, along with any supporting materials for the Discovery Master to review prior to the conference.

BY THE DISCOVERY MASTER:



HARRIS T. BOCK, ESQ.

UNITED STATES DISTRICT COURT
SOUTHERN DISTRICT OF ILLINOIS

IN RE YASMIN AND YAZ)	
(DROSPIRENONE) MARKETING, SALES)	3:09-md-02100-DRH-PMF
PRACTICES AND PRODUCTS LIABILITY)	
LITIGATION)	MDL No. 2100

This Document Relates to:

ALL CASES

ORDER

INTRODUCTION AND BACKGROUND

On February 18, 2010, this Court entered Case Management Order Number 10 (“CMO 10”) (3:09-cv-2100 Doc. 672), governing the redaction of documents and claims of privilege. This Case Management Order, was the result of extensive arm’s length negotiations among highly experienced and informed counsel on both sides. The terms contained in CMO 10 were voluntarily agreed to and drafted by the parties. Pursuant to CMO 10, Defendants are permitted to “redact from produced documents, materials or other things, or portions thereof, the following items:”

Those portions of documents that contain information relating to Bayer’s non-Drospirenone-containing medicines or products. With respect to Drospirenone-containing medicines other than YAZ/Yasmin/Ocella, Defendants may redact those

portions of any material that relates to any business strategy, marketing, or sales or that otherwise does not contain safety, adverse event, efficacy, or scientific study information[.]

(3:09-cv-2100 Doc. 672 ¶ A(5)). Since the entry of CMO 10, the Bayer Defendants have produced approximately 1.3 million documents consisting of approximately 33 million pages. Over the course of that production, the Bayer Defendants have been redacting documents as permitted under CMO 10. Plaintiffs contend that of the 1.3 million documents produced, approximately 14% have been redacted based on the provisions agreed to by the parties in ¶ A(5) (3:09-cv-2100 Doc. 1273 p. 2 n. 1).

Plaintiffs now bring this motion to modify CMO 10, pursuant to Rule 22.6 of the Manual for Complex Litigation. Specifically, Plaintiffs seek to amend CMO 10, paragraph 5(A) so that “(1) Defendants are prohibited from redacting business strategy, marketing, or sales information regarding Defendants’ other Drospirenone (“DRSP”)-containing medicines other than YAZ/Yasmin/Ocella; (2) Defendants are prohibited from redacting business strategy, marketing and sales information pertaining to Bayer’s hormonal contraceptives; (3) Defendants are required to un-redact any and all documents that have redacted thus far on these grounds.” (3:09-cv-2100 Doc. 1272).

Fortunately, the parties have worked together on issues throughout this litigation and consistently meet and confer thoroughly before bringing a dispute to the Court. The parties’ efforts to resolve the present dispute are no

exception; both parties met and conferred on these issues in an effort to resolve the dispute. Although the Court encourages and appreciates such efforts, the parties also have a responsibility to bring disputes to the Court's attention in a timely manner to avoid unnecessarily compounding the problem. Regrettably, in this instance, 33 million pages were produced before the matter was brought to the Court's attention and as a result the issue of burden, argued extensively by the Defendants, has grown.

ANALYSIS

A. Scope of Discovery

The federal discovery rules are liberal in order to assist in the preparation for trial and settlement of litigated disputes. *See Bond v. Utreras*, 585 F.3d 1061, 1075 (7th Cir.2009). Pursuant to Federal Rule of Civil Procedure 26(b)(1), “[p]arties may obtain discovery regarding any nonprivileged matter that is relevant to any party's claim or defense... Relevant information need not be admissible at the trial if the discovery appears reasonably calculated to lead to the discovery of admissible evidence.” Fed. R. Civ. P. 26(b)(1).

B. Modification of Case Management Orders

The Manual for Complex Litigation, published by the Federal Judicial Center, is a primary reference text in this litigation. Pursuant to Rule 22.6 the Court has the authority to update, and modify case management orders as the litigation unfolds. The Court also notes, however, that case management orders

are designed to facilitate the complex litigation process and are a vital tool in managing the discovery process. Thus, the Court does not take requests to modify case management orders lightly, especially when a case management order is voluntarily entered into after extensive negotiations between the parties.

C. Modification of Case Management Order Number 10 ¶ 5(A)

At oral argument, Plaintiffs theorized that Drospirenone, an active ingredient in YAZ and Yasmin, is the component that caused Plaintiffs' alleged injuries. Plaintiffs contend that their primary concern with ¶ 5(A) is that it allows Defendants to redact sales and marketing information relating to other Drospirenone containing products. Plaintiffs contend that the sales, marketing, scientific, and regulatory information pertaining to YAZ and Yasmin and to other Drospirenone containing products are so intertwined, the redaction protocols in ¶ 5 (A) are preventing Plaintiffs from obtaining discovery of relevant information and from fully understanding the documents that have been produced.

Defendants argue, credibly, that to go back over the documents that have been produced thus far, 90% of what Plaintiffs say they are requesting, will be burdensome. There are three categories of redactions to this point; relevance, privilege and confidentiality. Of the relevance redactions, some are for the subject matter drugs at issue in this motion and others are for completely unrelated drugs. Therefore, the Defendants, submit they cannot simply go back and remove all relevance redactions. Plaintiffs counter that the percentage of non-subject

matter relevance redactions are a small percentage of the total of relevance redactions. However, they can only speculate about that comparison based on context of matters discussed in the material.

Burden on the Defendants is a factor the Court should weigh and balance against the Plaintiffs need to access relevant information in the possession of the Defendants or information which is likely to lead to the discovery of relevant or admissible evidence. *See* Fed. R. Civ. P. 26 (b)(2)(iii) (discovery shall be limited by the Court “if it determines that ... the burden or expense of the proposed discovery outweighs its likely benefit, taking into account the needs of the case, the amount in controversy, the parties' resources, the importance of the issues at stake in the litigation, and the importance of the proposed discovery in resolving the issues.” However, it is a factor that cannot outweigh a litigant’s right to discover relevant evidence. Just as the Court, in this litigation, has dismissed a number of cases when plaintiffs have failed to provide the Defendants with relevant evidence to which they are entitled in Plaintiff Fact Sheets.

On the other side of this issue, the Plaintiffs argue, credibly, that the redactions in the past, and presumably to come, make the documents produced nonsensical for the most part and impossible for Plaintiffs “to get traction” on the gathering of meaningful information as they build a case of admissible evidence in preparation for trial. From the exhibits Counsel demonstrated in open court, it

was clear that the redaction complained of made the production virtually useless in too many instances.

Although the Court is reluctant to modify an agreed case management order, especially after such a large number of documents have been produced, the Court agrees that information pertaining to Defendants' other Drospirenone containing medicines and information pertaining to Drospirenone containing medicines in general could lead to the discovery of admissible evidence. Moreover, as noted, the Court feels that the redaction complained of makes the production virtually unusable. Accordingly, the Court concludes that the redaction provisions in ¶ 5(A) should be modified in the following way:

(1) Defendants are prohibited from redacting business strategy, marketing, or sales information regarding Defendants' other Drospirenone ("DRSP")-containing medicines other than YAZ/Yasmin/Ocella; and

(2) Defendants are required to un-redact any and all documents that have redacted thus far on these grounds.

The Court, however, will not grant Plaintiffs' request that the Defendants be prohibited from redacting business strategy, marketing and sales information pertaining to Bayer's "hormonal contraceptives."

This request is overly broad. The Court feels that this decision strikes an appropriate balance between the burden that will be placed on the Defendants and the Plaintiffs' right to discovery of relevant information.

IT IS SO ORDERED.

Dated September 22, 2010

/s/ David Herndon

**Chief Judge
United States District Court**

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**U.S. District Court
Southern District of Illinois**

Notice of Electronic Filing

The following transaction was entered on 10/15/2010 at 4:03 PM CDT and filed on 10/15/2010

Case Name: In Re: Yasmin and Yaz (Drospirenone) Marketing, Sales Practices and Products Liability Litigation
Case Number: 3:09-md-02100-DRH -PMF
Filer:
Document Number: 1336(No document attached)

Docket Text:

MINUTE ORDER RE: Deadline for complying with the production of documents pursuant to the Order amending portions of CMO 10 [1308] and for complying with CMO 22 [1303]. On September 22, 2010, the Court entered an Order (3:09-cv-2100 Doc.1308) modifying the redaction provisions in Case Management Order Number 10 (CMO 10) (3:09-cv-2100 Doc. 672). As a result, the Defendants are currently in the process of generating documents that are responsive to CMO 10 and compliant with the Courts recent modification to the redaction provisions in CMO 10. On September 14, 2010, the Court issued Case Management Order Number 22 (CMO 22) with regard to the Clintrace database and making that database available to the Plaintiffs (3:09-cv-2100 Doc.1303). Defendants report that they are diligently working to meet their obligations pursuant to the above described orders. The Court concludes that it would be beneficial to impose a deadline for compliance with the above described orders. Accordingly, the Court hereby ORDERS the Defendants to produce the un-redacted documents described in the Courts September 22, 2010 Order [1308] which amended CMO 10 and to comply with their obligations pursuant to CMO 22 [1303] by Friday October 22, 2010.. Signed by Chief Judge David R. Herndon on 10/15/2010. (dsw)THIS TEXT ENTRY IS AN ORDER OF THE COURT. NO FURTHER DOCUMENTATION WILL BE MAILED.

3:09-md-02100-DRH -PMF Notice has been electronically mailed to:

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