

UNITED STATES DISTRICT COURT
SOUTHERN DISTRICT OF INDIANA
INDIANAPOLIS DIVISION

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|--------------------------------|--------------------------|
| ELI LILLY AND COMPANY, |) |
| |) Cause No. |
| Plaintiff, |) 1:10-CV-01376-TWP-DKL |
| |) Indianapolis, Indiana |
| vs. |) August 22, 2013 |
| |) 9:04 a.m. |
| TEVA PARENTERAL MEDICINES, |) |
| INC., APP PHARMACEUTICALS, |) |
| LLC, PLIVA HRVATSKA D.O.O., |) |
| TEVA PHARMACEUTICALS USA, |) |
| INC., BARR LABORATORIES, INC., |) |
| |) |
| Defendants. |) |
| |) |

V O L U M E IV

Before the Honorable
TANYA WALTON PRATT

OFFICIAL REPORTER'S TRANSCRIPT OF
BENCH TRIAL

| | |
|-----------------|---------------------------------|
| Court Reporter: | David W. Moxley, RMR, CRR, CMRS |
| | United States District Court |
| | 46 East Ohio Street, Room 340 |
| | Indianapolis, Indiana 46204 |

PROCEEDINGS TAKEN BY MACHINE SHORTHAND
TRANSCRIPT CREATED BY COMPUTER-AIDED TRANSCRIPTION

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1 approving the drug, was saying no. So, it was a tough
2 situation, yeah. We had to act quickly.

3 Q. And during the call, was there a discussion about what the
4 response should be?

5 A. Yes, there was.

6 Q. Can we take a look at Exhibit 2262, please? And let's
7 pull up 1-1. Thank you.

8 Is this the letter that Lilly sent back to the FDA the
9 next day, on December 22nd, Doctor?

10 A. Yes, it is.

11 Q. And in the beginning, does the letter just recast what the
12 response was -- what the FDA's fax said?

13 A. Yes, in the fax of December 21st, late afternoon.

14 Q. And that's -- the FDA said that the information in the
15 annual report about the toxicities in the trial does not
16 appear to support the addition of vitamins?

17 A. That's correct.

18 Q. What is the annual report referring to?

19 A. The annual report is now still referring to that
20 September 10th cutoff; and actually, in a sense, it's probably
21 comprehensible at this time if the FDA reviewer is still
22 referring to this document, because that document didn't
23 include the deaths -- the sudden deaths we saw post that
24 annual report.

25 Q. Did Lilly in this letter describe those -- that additional

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1 information?

2 A. Yeah, we had to right away.

3 Q. Okay. Let's look at 1-4, still on the first page.

4 Can you describe what Lilly is telling the FDA here?

5 A. Yeah. Lilly is telling the FDA that we have actually
6 additional information on the safety profile that we have seen
7 in addition to what we had as of September 10th.

8 Q. Okay. And then 1-3, please.

9 And then, in the bottom of that page, and onto the
10 next page, can you explain what Lilly is stating here?

11 A. Yeah. Here, Lilly is actually updating the FDA that
12 within the time that we're going back and forth on the
13 reaction, especially from that report, we have seen actually
14 patients dying from drug-related deaths; and that was
15 extremely concerning to us.

16 Q. Can we take a look at 2-3? This is on the second page of
17 the letter. There's a paragraph.

18 And the paragraph says, "Lilly has consulted a number
19 of oncology experts regarding patients' safety"; and then it
20 says, "These consultants were in unanimous agreement that
21 intervention was necessary to promote patients' safety in the
22 pemetrexed trials."

23 Does that accurately reflect what the experts told you
24 on that conference call you had?

25 A. Yes.

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1 Q. The next sentence, Doctor, says, "They all further
2 suggested that supplementation with folic acid would offer the
3 best chance of reducing serious toxicity to the broadest
4 patient population."

5 Does that accurately reflect what you were told or
6 what Lilly was told by the experts on that call?

7 A. Yes.

8 Q. Then the next sentence says, "These experts felt that
9 supplementation with low levels of folic acid would not
10 adversely affect efficacy of pemetrexed."

11 Does that statement accurately reflect what the
12 experts were saying on the call?

13 A. No, actually not.

14 Q. What were the experts saying about efficacy on the call,
15 Doctor?

16 A. What I understood on the call was that the risk/benefit
17 had shifted towards intervening to protect the patients from
18 the toxicities, and this was warranted by these drug-related
19 deaths now being observed, and that we had --

20 MR. WIESEN: Your Honor, now we've gone into hearsay
21 that's contradicting documents they submitted to the FDA.

22 MR. GENDERSON: Your Honor, this is not hearsay now.
23 It's not for the truth. It's for what was stated. And we're
24 going to explain that the person who wrote this letter wasn't
25 on the call. All of this happened over a -- literally an

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