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US District Court - New Jersey

**Jazz Pharmaceuticals
v.
Roxane Laboratories**



Video Deposition of:
Glenn Van Buskirk, Ph.D.
October 28, 2015

[REDACTED]

7 G L E N N A . V A N B U S K I R K,
8 Ph.D., Nonclinical Drug Development Consulting
9 Services, LLC, having been duly sworn, was
10 examined and testified as follows:

[REDACTED]

[REDACTED]

23 Q. You didn't attend any advisory committee
24 meetings for -- for GHB or Xyrem in front of the
25 FDA, did you?

1 A. No, I did not.

2 Q. And you were not aware of any 2001
3 advisory committee meetings before the FDA,
4 correct?

5 A. No. Until you mentioned advisory
6 meetings, I didn't even know there were such
7 things. Oh, I know they exist; I didn't know they
8 existed for this compound. So, no, I was not
9 aware of it.

10 Q. Something I forgot to get in. Your
11 educational background, you have a Bachelor of
12 Science in pharmacy, correct?

13 A. I do.

14 Q. And a master's in pharmaceutical
15 science?

16 A. Correct.

17 Q. And a Ph.D. in pharmaceutical science,
18 correct?

19 A. That is also correct.

20 Q. And you -- you are also a licensed
21 pharmacist?

22 A. Yes, I am.

[REDACTED]

[REDACTED]

[REDACTED]

[REDACTED]

[REDACTED]

[REDACTED]

[REDACTED]

[REDACTED]

[REDACTED]

[REDACTED]

[REDACTED]

9 Q. Okay. Do you monitor the Federal
10 Register for notices of advisory committee
11 meetings for unapproved drugs?

12 A. No, that's not something I do.

13 Q. Do individuals with degrees in pharmacy
14 generally monitor the Federal Register for notices
15 on unapproved drugs?

16 A. Not unless they're in regulatory
17 affairs. And from my perspective as a scientist,
18 we monitor it for the general regulations around
19 such things as good manufacturing practices and
20 specifications and things that affect our daily
21 lives. Other people monitor the regulatory
22 component.