

UNITED STATES NUCLEAR REGULATORY COMMISSION WASHINGTON, D. C. 20555

October 23, 1990

Dr. Donna Bush Chief, Drug Testing Section Division of Applied Research National Institute on Drug Abuse 5600 Fishers Lane (Room 9-A-54-) Rockville, Maryland 20857

Dear Dr. Bush:

Forwarded for your action in accordance with Section 2.8(e) of Appendix A to 10 CFR Part 26, are 4 reports of investigations from Portland General Electric Company.

- Letter, dtd 2/28/90, reports a false negative on a blind performance test reported by American BioTest Laboratories, Santa Clara, California. The error was caused by the failure of a laboratory technician to properly calibrate the immunoassay equipment. The root cause was determined to be failure by the laboratory to establish an adequate procedure.
- Letter, dtd 5/25/90, reports a false negative on a blind performance test reported by Laboratory Pathology, Seattle, Washington. Sample degradation was ruled out, and the report concludes that the probable cause was that certain physiological parameters unique to that specimen caused interference with immunoassay screening instruments, but did not affect the GC/MS technology.
- Letter, dtd 6/21/90, reports a false negative on a blind performance test reported by Laboratory Pathology. The cause of the error was reported to be incomplete hydrolysis during preparation of the specimen for GC/MS.
- 4. Portland General Electric Company letter, dtd 9/17/90, concerning a false negative report of a blind performance test specimen by Laboratory Pathology. The investigation disclosed that the technician transposed the last 2 digits of the sample number and another aliquot was selected for confirmation testing. Also, the laboratory failed to determine why the GC/MS response was zero after the initial presumptive positive screen.

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Dr. Donne Bush

Should you have any questions, please call me at 492-0944.

Sincerely,

Loren L. Bush, Chief Program Development and Review Section Office of Nuclear Reactor Regulation

Enclosures: As stated

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