

# WISCONSIN DEPARTMENT OF REGULATION & LICENSING



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IN THE MATTER OF \_\_\_\_\_ :  
DISCIPLINARY PROCEEDINGS AGAINST \_\_\_\_\_ :  
\_\_\_\_\_ : ORDER DENYING STAY  
LISA GARDNER, R.N., \_\_\_\_\_ :  
RESPONDENT. \_\_\_\_\_ :

4. Respondent shall in addition refrain from the consumption of over-the-counter medications or other substances which may mask consumption of controlled substances or of alcohol, or which may create false positive screening results, or which may interfere with respondent's treatment and rehabilitation. Respondent shall report all medications and drugs, over-the-counter or prescription, taken by respondent to the Supervising

Health Care Provider within 24 hours of ingestion or administration, and shall identify the person or persons who prescribed, dispensed, administered or ordered said medications or drugs. Within 24 hours of a request by the Supervising Health Care Provider or the Board or its designee, Respondent shall provide releases which comply with state and federal laws authorizing release of all health care records by the person who prescribed, dispensed, administered or ordered this medication for respondent. These releases shall also authorize the Supervising Health Care Provider, the Board or its designee to discuss the Respondent's health care with the person who prescribed, dispensed, administered or ordered this medication. The terms of this paragraph shall not be deemed to modify or negate Respondent's obligations as set forth in this Order.

5. The Department Monitor is the individual designated by the Board as its agent to coordinate compliance with the terms of the Order, including receiving and coordinating all reports and petitions, and requesting additional monitoring and surveillance. The Department Monitor may be reached as follows:

Department Monitor  
Department of Regulation Division of Enforcement  
P.O. Box 8935  
Madison, WI 53708-8935  
FAX (608) 266-2264  
TEL. (608) 267-7139

6. Respondent shall provide and keep on file with the Supervising Health Care Provider, all treatment facilities and personnel, laboratories and collections sites current releases which comply with state and federal laws authorizing release of all urine, blood and hair specimen screen results and medical and treatment records and reports to, and permitting the Supervising Health Care Provider and all treating physicians and therapists to disclose and discuss the progress of respondent's treatment and rehabilitation with the Board or any member thereof, or with any employee of the Department of Regulation and Licensing acting under the authority of the Board. Copies of these releases shall be filed simultaneously with the Department Monitor.

7. Respondent shall supply on at least a weekly basis, random monitored urine, blood or hair specimens as the Supervising Health Care Provider shall direct. The Supervising Health Care Provider (or designee) shall request the specimens from Respondent and these requests shall be random with respect to the hour of the day and the day of the week. In addition, the Board or its designee may at any time request a random monitored urine, blood or hair specimen from Respondent by directing the Department Monitor in the Department of Regulation and Licensing, Division of Enforcement to contact Respondent and request Respondent provide a specimen. To prevent the respondent's ability to predict that no further screens will be required for a given period (because the minimum frequency for that period has been met), the program of monitoring shall require respondent to provide in each quarter at least two (2) random screenings in excess of the minimums specified in this Order.

8. Respondent shall keep the Supervising Health Care Provider informed of Respondent's location and shall be available for contact by the Supervising Health Care Provider at all times.

9. All requested urine, blood or hair specimens shall be provided by Respondent within five (5) hours of the request for the specimen. All urine specimen collections shall be a split sample accomplished by dividing urine from a single void into two specimen bottles. The total volume of the split sample shall be at least 45 ml. of urine. All split sample urine specimens, blood specimens and hair specimens shall be collected, monitored and chain of custody maintained in conformity with the collection, monitoring and chain of custody procedures set forth in 49 CFR Part 40. Urine specimen collections shall be by direct observation if:

a. The Respondent must provide an additional specimen because Respondent's initial specimen was outside of the normal temperature range (32.5 - 37.7 C/90.5 - 99.8 F) and respondent refuses to have an oral body temperature measurement or respondent does provide an oral body temperature measurement and

the reading varies by more than 1 C/1.8 F from the temperature of the urine specimen;

b. Respondent's last provided specimen was determined by the laboratory to have a specific gravity of less than 1.003 and creatinine concentration below 0.2 g/l;

c. The collection site person observes Respondent acting in such a manner to provide reason to believe that Respondent may have attempted or may attempt to substitute or adulterate the specimen. The collection site person, if he or she believes that the initial urine specimen may have been adulterated or a substitution made, shall direct Respondent to provide an additional observed urine specimen;

d. The last provided specimen resulted in a positive or suspected positive test result for the presence of controlled substances; or

e. The Board (or any member of the Board), the Department Monitor, or Respondent's Supervising Health Care Provider directs that the urine specimen collection be by direct observation.

If either of the above conditions (a) or (c) requires collection of an additional observed urine specimen, the collection of the subsequent specimen shall be accomplished within the required five (5) hours of the request for the initial specimen; the collection of the initial specimen shall not satisfy the requirement that the urine specimen be collected within five (5) hours of the request for the initial specimen.

10. The drug and alcohol treatment program in which Respondent is enrolled shall at all times utilize a United States Department of Health and Human Services certified laboratory for the analysis of all specimens collected from Respondent.

11. The drug and alcohol treatment program in which Respondent is enrolled shall utilize only those urine, blood and hair specimen collection sites for collection of Respondent's urine, blood or hair specimens as comply with the United States Department of Transportation collection and chain of custody procedures set forth in 49 CFR Part 40.

12. The Supervising Health Care Provider, treatment facility, laboratory and collection site shall maintain a complete and fully documented chain of custody for each urine, blood or hair specimen collected from Respondent.

13. Every urine specimen collected from Respondent shall be analyzed at the time of collection for tampering by measurement of the temperature of the specimen and the oral temperature of Respondent. Every urine specimen collected from Respondent shall be further analyzed at the laboratory for tampering by measuring the creatinine concentration and the specific gravity of the specimen. The laboratory may at its discretion or at the direction of a Supervising Health Care Provider or the Board or any member thereof conduct additional tests to evaluate the urine specimen for tampering including, but not limited to, pH, color and odor.

14. Every urine, blood or hair specimen collected from Respondent shall be analyzed for alcohol, amphetamine, cocaine, opiates, phencyclidine, marijuana, methadone, propoxyphene, methaqualone, barbiturates, benzodiazepines or the metabolites thereof. The Board or its designated agent may at any time direct that screens for additional substances and their metabolites be conducted by scientific methods and instruments appropriate to detect the presence of these substances. The laboratory shall conduct confirmatory tests of positive or suspected positive test results by appropriate scientific methods and instruments including, but not limited to, gas chromatography and mass spectrometry.

15. All urine, blood or hair specimens remaining after testing shall be maintained in a manner necessary to preserve the integrity of the specimens for at least seven (7) days; and all positive or suspected positive urine, blood or hair specimens remaining after testing shall be so maintained for a period of at least one (1) year. The Supervising Health Care Provider or the Board or any member thereof may direct that the urine, blood or hair specimens be maintained for a longer period of time.

16. For the purpose of further actions affecting Respondent's license under the Order, it shall be presumed that all confirmed positive reports are valid. Respondent shall have the burden of proof to establish that the positive report was erroneous and that the respondent's specimen sample did not contain alcohol or controlled substances or their metabolites.

17. If any urine, blood or hair specimen is positive or suspected positive for any controlled substances or alcohol, Respondent shall promptly submit to additional tests or examinations as the Supervising Health Care Provider shall determine to be appropriate to clarify or confirm the positive or suspected positive urine, blood or hair specimen test results.

18. The Supervising Health Care Provider shall report immediately to the Department Monitor in the Department of Regulation and Licensing, Division of Enforcement by FAX or telephonic communication: any failure of Respondent to provide a urine, blood or hair specimen within five (5) hours from the time it was requested; or of any inability to locate Respondent to request a specimen. The laboratory shall immediately report all urine specimens suspected to have been tampered with and all urine, blood or hair specimens which are positive or suspected positive for controlled substances or alcohol to the Department Monitor, and to the Supervising Health Care Provider.

19. The laboratory shall within 48 hours of completion of each drug or alcohol analysis mail the report from **all** specimens requested of Respondent under this Order to the Department Monitor (regardless of whether the laboratory analysis of the specimen was positive or negative for controlled substances, their metabolites or alcohol). Each report shall state the date and time the specimen was requested; the date and time the specimen was collected; the results of the tests performed to detect tampering; and the results of the laboratory analysis for the presence of controlled substances and alcohol.

20. The Supervising Health Care Provider shall submit formal written reports to the Department Monitor in the Department of Regulation and Licensing, Division of Enforcement, P.O. Box 8935, Madison, Wisconsin 53708-8935 on a quarterly basis, as directed by the Department Monitor. These reports shall assess Respondent's progress in the drug and alcohol treatment program and summarize the results of the urine, blood or hair specimen analyses. The Supervising Health Care Provider shall report immediately to the Department Monitor [Division of Enforcement, P.O. Box 8935, Madison, Wisconsin 53708-8935, FAX (608)266-2264, telephone no. (608)267-7139] any violation or suspected violation of the Board's Final Decision and Order.

21. Respondent is responsible for compliance with all of the terms and conditions of the Final Decision and Order. It is the responsibility of Respondent to promptly notify the Department Monitor, of any suspected violations of any of the terms and conditions of the Order, including any failures of the Supervising Health Care Provider, treatment facility, laboratory or collection sites to conform to the terms and conditions of this Order.

22. If the Board determines that the Supervising Health Care Provider, treatment facility, laboratory or collection sites have failed to satisfy the terms and conditions of the Final Decision and Order, the Board may, at its sole discretion, direct that Respondent continue treatment and rehabilitation under the direction of another Supervising Health Care Provider, treatment facility, laboratory or collection site which will conform to the terms and conditions of this Final Decision and Order.

23. Respondent may petition the Board for modification of the terms of the limited license. Any such petition shall be accompanied by a written recommendation from respondent's Supervising Health Care Provider expressly supporting the specific modifications sought. Denial of the petition in whole or in part shall not be considered a

denial of a license within the meaning of Sec. 227.01(3)(a), Stats. and Respondent shall not have a right to any further hearings or proceedings on any denial in whole or in part of the petition for modification of the limited license.

After two years of continuous active professional practice under the Order and without relapse, and upon recommendation of the Supervising Health Care Provider, respondent may petition the Board for a termination of all limitations on the license, and restoration of an unlimited license. Such restoration shall be in the sole discretion of the Board, and denial of the petition in whole or in part shall not be considered a denial of a license within the meaning of Sec. 227.01(3)(a), Stats. and Respondent shall not have a right to any further hearings or proceedings on any denial in whole or in part of the petition for termination of the limitations and restoration of unlimited licensure.

24. Respondent shall be responsible for all costs and expenses incurred in conjunction with the monitoring, screening, supervision and any other expenses associated with compliance with the terms of this Order.

25. Respondent shall refrain from access to or the administration of controlled substances in her work setting until such time as access or administration is approved by the Board.

26. Respondent shall practice only under the general supervision of a licensed professional nurse or other licensed health care professional approved by the Board or in a work setting pre-approved by the Board or its designated agent. Until further order of the Board Respondent shall not work as a nurse in a home health care setting.

27. Respondent shall arrange for her employer to provide formal written reports to the Department Monitor in the Department of Regulation and Licensing, Division of Enforcement, P.O. Box 8935, Madison, Wisconsin 53708-8935 on a quarterly basis, as directed by the Department Monitor. These reports shall assess Respondent's work performance.

28. Respondent shall report to the Board any change of employment status, residence, address or telephone number within five (5) days of the date of a change.

29. Respondent shall furnish a copy of the Order to all present employers immediately upon issuance of this Order, and to any prospective employer when respondent applies for employment as a health care provider.

On August 8, 2003, the board considered Ms. Gardner's request for a stay of the suspension of her license. The board also considered irregularities in the screening procedure and the panel utilized. Based upon all information on record herein, the board orders as follows:

#### ORDER

NOW, THEREFORE, IT IS ORDERED that the request for a stay of the suspension of the license of Lisa Gardner, RN is hereby denied. Ms. Gardner may renew her request at the board's September 5, 2003, meeting, and she shall appear before the board on that date in support of her request.

Date this 14<sup>th</sup> day of August, 2003

WISCONSIN BOARD OF NURSING

Linda M. Sanner, RN

Chair