

## Validity Testing Information Part I

Laboratory Name:

NWT Drug Testing

Address:

1141 E. 3900 S.

Salt Lake City, UT 84124

Responsible Person:

Dr. David J. Kuntz

(Printed Name)

I certify that the answers and information provided are true and correct as of this date. Any false, fictitious, or fraudulent answers or information provided may violate Federal Law and could subject me to prosecution, monetary penalties, or both (Sec 18 U.S.C. 1001; 31 U.S.C. 3801-812).

 ALT-RP  
Signature, Responsible Person

10-04-00  
Date

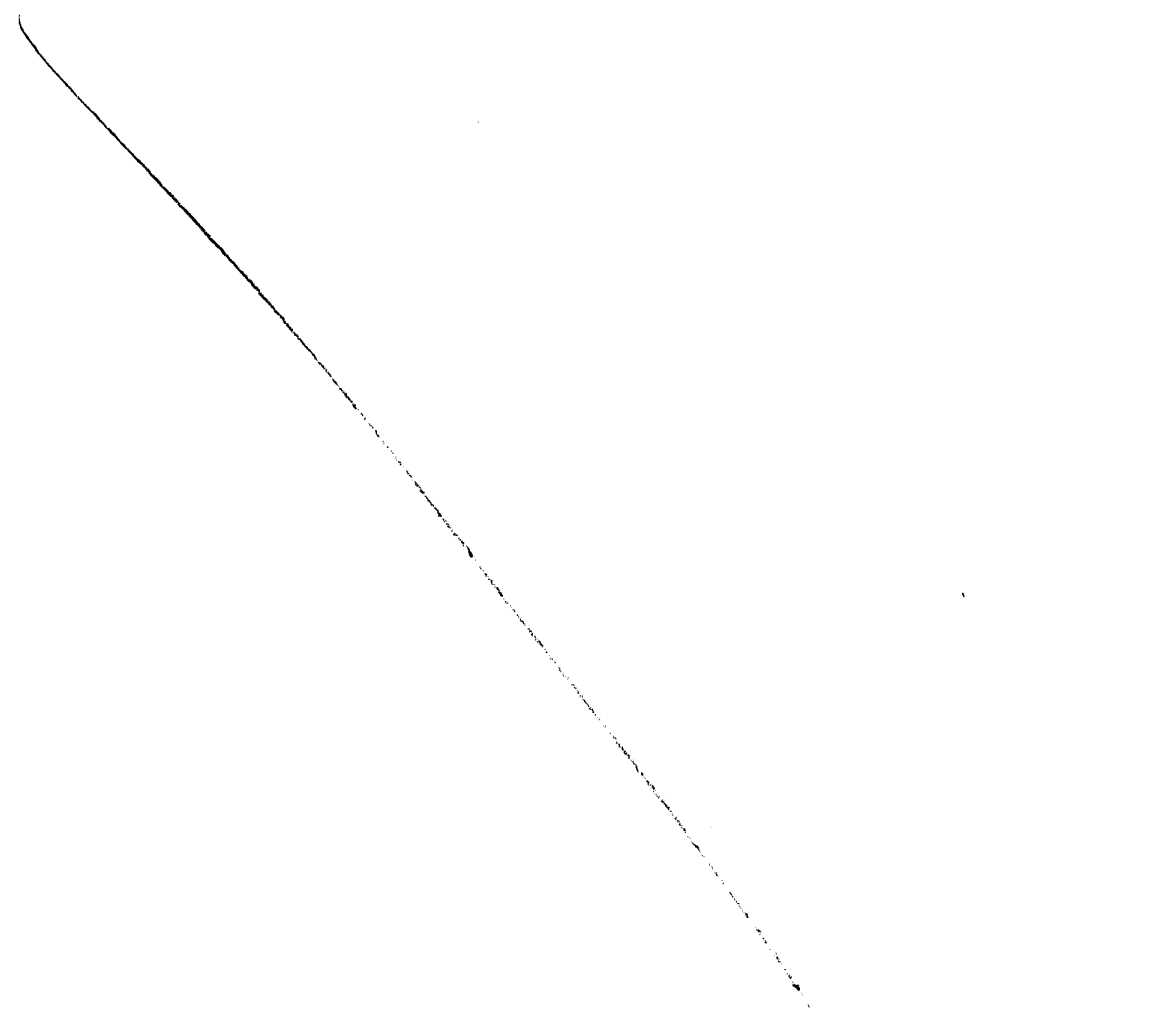
Laurie Tobler ALT-RP  
Printed Name, Responsible Person

# Validity Testing Audit NWT, Inc. Oct 17, 2000

Specimen ID Number	Laboratory Accession Number	Date of Receipt	Date Reported	Reported Result	Quantitative Test Result
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Specimen ID Number	Laboratory Accession Number	Date of Receipt	Date Reported	Reported Result	Quantitative Test Result
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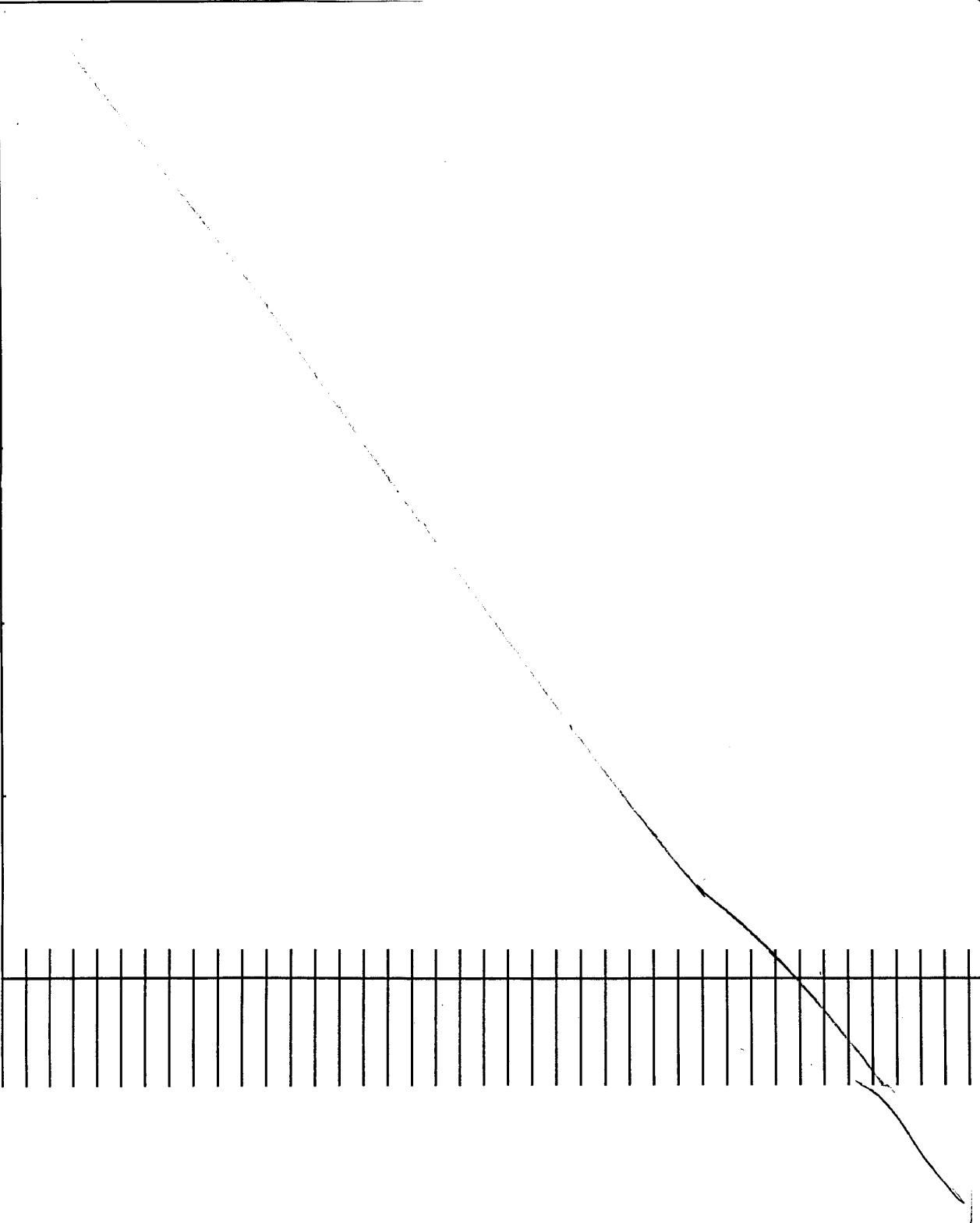
Specimen ID Number	Laboratory Accession Number	Date of Receipt	Date Reported	Reported Result	Quantitative Test Result
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Specimen ID Number	Laboratory Accession Number	Date of Receipt	Date Reported	Reported Result	Quantitative Test Result
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RESEARCH TRIANGLE INSTITUTE



National Laboratory Certification Program

December 27, 2000

0074  
Dr. David J. Kuntz  
NWT Drug Testing  
1141 East 3900 South  
Salt Lake City, UT 84124

Dear Dr. Kuntz:

We have reviewed the material provided in your correspondence of December 7, 2000, submitted in response to issues raised during the November 1-3, 2000, specimen validity testing inspection of your laboratory as outlined in our correspondence of November 17, 2000. The information submitted by the laboratory appears to demonstrate that appropriate corrective actions have been completed to address the issues raised.

Based upon our review of the material submitted, it appears that the laboratory's specimen validity testing procedures are in compliance with program guidance. All corrective actions must be implemented within 30 days of the receipt of this correspondence. Failure to comply may result in the laboratory's suspension to perform specimen validity testing on federally regulated specimens. Corrective actions will be reviewed during the next inspection.

If you have any questions or if we can be of further assistance, please call me at (919) 541-6176 or Dr. Michael R. Baylor at (919) 541-7043.

Sincerely,

  
Susan Crumpton  
NLCP Technical Analyst

cc: Project Files/SVT074





RESEARCH TRIANGLE INSTITUTE

National Laboratory Certification Program

November 17, 2000

0074  
Dr. David J. Kuntz  
NWT Drug Testing  
1141 East 3900 South  
Salt Lake City, UT 84124

Dear Dr. Kuntz:

The enclosed critique was developed from the inspection report associated with the November 1-3, 2000, specimen validity testing inspection of your laboratory under the National Laboratory Certification Program (NLCP). The laboratory's procedures were not in full compliance with program guidance issued in Program Document 035 (September 28, 1998) and Program Document 037 (July 28, 1999). The laboratory must submit information to address the following issues raised:



Dr. Kuntz  
Page 2 of 2  
11/17/00

The laboratory must submit, within 30 calendar days of receipt of this letter, documentation to demonstrate that corrective actions have been implemented to address the issues raised. In responding to these issues, please organize the material in your document in accordance with the sections and item numbers as listed in this correspondence. The laboratory must also review the enclosed critique and take all necessary corrective actions. All corrective actions must be implemented within 30 days receipt of this correspondence and will be reviewed at the next inspection.

If you have any questions or if we can be of further assistance, please call me at (919) 541-6176 or Dr. Michael Baylor at (919) 541-7043.

Sincerely,

  
Susan Crumpton  
NLCP Technical Analyst

Enclosure

cc: Project Files/svt074

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## NATIONAL LABORATORY CERTIFICATION PROGRAM

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### Document Review and Critique

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Laboratory I.D. Number: 0074  
Document No. Final

Laboratory: NWT Drug Testing

Location: Salt Lake City, UT

Document Reviewed:  Specimen Validity Testing Inspection Report

Date: 1 November 2000

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A review of the National Laboratory Certification Program (NLCP) consensus inspection report has been completed. Issues identified during the inspection are described on the following pages. Evidence that appropriate remedial action has been taken is required for continued certification.

The following comments were noted, and appear in the same order as the corresponding questions in the Laboratory Inspection Report:

Section E. Standard Operating Procedures - Procedures Manual

Section F. Chain-of-Custody, Accessioning, and Security

Section G. Quality Control and Quality Assurance

Section I. Specimen Validity Tests

Section K. Records Audit

Section L. Certification and Reporting

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WORKPLACE  
DRUG TESTING

A Division of  
NWT Inc.

1141 EAST 3900 SOUTH

SALT LAKE CITY

UTAH 84124 USA

801-293-2300

800-322-3361

FAX 801-293-2387

December 7, 2000

Susan Crumpton  
National Laboratory Certification Program  
Research Triangle Institute  
PO Box 12194  
3040 Cornwallis Road  
Research Triangle Park, NC 27709

received  
SDC 12/12/00

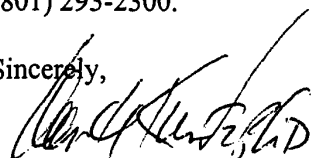
Reference: Specimen Validity Testing Inspection; November 1-3 2000  
Corrective Action Response

Dear Ms. Crumpton:

In response to your letter describing deficiencies which require corrective action,  
we have addressed the following issues:

If you have any questions concerning the submitted data, please contact me at (801) 293-2300.

Sincerely,



David J. Kuntz, Ph.D.  
Responsible Person

RESEARCH TRIANGLE INSTITUTE



National Laboratory Certification Program

December 27, 2000

0074  
Dr. David J. Kuntz  
NWT Drug Testing  
1141 East 3900 South  
Salt Lake City, UT 84124

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If you have any questions or if we can be of further assistance, please call me at (919) 541-6176 or Dr. Michael R. Baylor at (919) 541-7043.

Sincerely,

  
Susan Crumpton  
NLCP Technical Analyst

cc: Project Files/SVT074