



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Alcohol, Drug Abuse and
Mental Health Administration
Rockville MD 20857

Robert L. Stout, Ph.D.
Clinical Reference Lab
11850 West 85th Street
Lenexa, Kansas 66214

NOV 2 1989

Dear Dr. Stout:

I am pleased to inform you that Clinical Reference Lab, Lenexa Kansas, has successfully met all of the requirements for laboratory certification as specified in the Department of Health and Human Services' (HHS) Mandatory Guidelines for Federal Workplace Drug Testing Programs (53 FR 11970).

Clinical Reference Lab will be placed on the list of laboratories certified as eligible to bid on contracts to perform drug testing for Federal Drug-Free Workplace Programs. The list of laboratories certified by the National Institute on Drug Abuse (NIDA) on behalf of the Department will be sent to all Federal Agencies. Bi-monthly updates to this list will be published in the Federal Register, and made available to the general public upon request.

To maintain certification from HHS, Clinical Reference Lab must continue to meet all the requirements of the Federal Guidelines as specified in Subpart C--Certification of Laboratories Engaged in Urine Drug Testing for Federal Agencies. Maintenance of certification requires participation in an every-other-month performance testing program plus periodic, on-site inspections (see Sections 3.2(b), 3.17, 3.19 and 3.20).

If you have any questions concerning NIDA's National Laboratory Certification Program, please contact the Division of Applied Research (formerly the Office of Workplace Initiatives), at (301) 443-6780.

The HHS laboratory standards for urine drug testing certification were designed to assure Federal Agencies and their employees that the laboratories and the scientific and methodological procedures used are of the highest quality. Your laboratory is to be congratulated for meeting all the requirements of the department's program.

Sincerely,

Charles R. Schuster

Charles R. Schuster, Ph.D.
Director
National Institute on Drug Abuse

ATTACHMENT C

ABOUT CRL

Certifications, Licenses & Accreditations

Certification / Certifying Body

American Board of Forensic Toxicology (ABFT)

www.abft.org

California - Department of Public Health

www.cdph.ca.gov

Centers for Disease Control and Prevention (CDC)

National Heart, Lung and Blood Institute Lipid Standardization Program

www.cdc.gov

Clinical Laboratory Improvement Amendments (CLIA)

www.phppo.cdc.gov/clia/

College of American Pathologists (CAP)

www.cap.org

Drug Enforcement Agency (DEA)

www.usdoj.gov/dea

Florida - Health Care Administration (AHCA)

ahca.myflorida.com

Forensic Toxicologist Certification Board (FTCB)

home.usit.net

Hawaii - Department of Health

hawaii.gov/health

Department of Health & Human Services (HHS)

Substance Abuse and Mental Health Services Administration (SAMHSA)

www.samhsa.gov

www.hhs.gov

Iowa - Department of Health (IDPH)

www.idph.state.ia.us

Kansas Department of Health & Environment (KDHE)

Laboratory Improvement Program

www.kdheks.gov

Kansas State Board of Pharmacy

www.accesskansas.org/pharmacy

Maine Center for Disease Control and Prevention

Office of the Department of Health and Human Services

www.maine.gov/dhhs/boh

Maryland Department of Health and Mental Hygiene

Office of Health Care Quality

www.dhmdh.state.md.us

Document

(Toxicology)

(General Testing Lab)

(Clinical Trials)

(Clinical Trials # 17D1021310)

(Gen Lab # 17D0667123)

(Toxicology # 17D0667123)

(Clinical Trials # 7185553)

(Gen Lab # 3021101)

(Toxicology # 3021103)

(Toxicology - Analytical Lab)

(Toxicology - Researcher)

(Toxicology)

(General Testing Lab)

(Toxicology)

(Toxicology)

(Toxicology)

Approved Laboratories List

(Clinical Trials)

(Toxicology)

(General Testing lab)

(Toxicology)

(Toxicology)

(General Testing Lab)

(Toxicology)

<u>National Glycohemoglobin Standardization Program (NGSP)</u>	<u>(Clinical Trials)</u>
<u>Level I Laboratory</u>	
<u>www.ngsp.org</u>	
<u>Nevada Department of Health and Human Services</u>	<u>(Toxicology)</u>
<u>dhhs.nv.gov/</u>	
<u>New York State Department of Health</u>	<u>(C.R.L. - Permit)</u>
<u>www.health.state.ny.us</u>	<u>(General Testing Lab - C.Q.)</u>
	<u>(Toxicology - C.Q.)</u>
<u>Oklahoma State Department of Health (OSDH)</u>	<u>(Toxicology)</u>
<u>www.ok.gov/health</u>	
<u>OSHA Blood Lead Certified Laboratory List</u>	<u>Blood Lead Laboratory List</u>
<u>www.osha.gov/SLTC/bloodlead/</u>	
<u>Pennsylvania - Department of Health</u>	<u>(General Testing Lab)</u>
<u>www.dsf.health.state.pa.us</u>	
<u>Texas Department of Public Safety</u>	<u>(Toxicology)</u>
<u>www.txdps.state.tx.us</u>	
<u>University of Washington - Cholesterol Reference Laboratory</u>	<u>(General Testing Lab)</u>
<u>Vermont - Department of Health</u>	<u>(General Testing Lab)</u>
<u>healthvermont.gov</u>	<u>(Toxicology)</u>
<u>West Virginia Department of Health and Human Resources</u>	<u>(General Testing Lab)</u>
<u>(WVDHHR)</u>	
<u>www.wvdhhr.org</u>	

Forensic Toxicologist Certification Board

hereby declares that the professional education, experience, and examination of

David Kuntz

has been found satisfactory and the requirements of the Board have been met. The Board, therefore, grants this certificate of qualification in

Forensic Drug Testing Toxicologist

at the rank of

Granted the 1st day of December 1999

Certificate Number 001

For the Board, I am,

Secretary

1999, Dallas, Texas



National Laboratory Certification Program

For more information, please visit our website at www.nlcplab.com or call us at (919) 541-7043.

c0007LU_070531_KuntzD_RP

May 31, 2007

0007
Mr. John Irving
Clinical Reference Lab
8433 Quivira Road
Lenexa, KS 66215-2802

Dear Mr. Irving:

The National Laboratory Certification Program (NLCP) staff at RTI has reviewed the acceptability of Dr. David Kuntz as a Responsible Person (RP) for Clinical Reference Lab in Lenexa, KS. After review of submitted documentation and the inspectors' comments from the laboratory's 33rd maintenance inspection on April 26-28, 2006, Dr. Kuntz has been found acceptable to serve as a Responsible Person for the laboratory.

The NLCP must be apprised of and approve all changes in the RP status of the laboratory. Guidance for effecting changes in the RP for a laboratory may be found in NLCP Program Document 12 (PD 012). If there are any questions concerning this matter, please contact me at (919) 541-7043.

Sincerely,

A handwritten signature in black ink, appearing to read "Michael R. Baylor".

Michael R. Baylor, Ph.D.
Co-Director, NLCP

ATTACHMENT E



National Laboratory Certification Program

3040 Cornwallis Road ■ PO Box 12194 ■ Research Triangle Park, NC 27709-2194 ■ USA
Telephone 919-541-7242 ■ Fax 919-541-7042 ■ www.rti.org

c0007LU_060504_KolbowD__ALTok

May 4, 2006

0007
Mr. Randal Clouette
Clinical Reference Lab
8433 Quivira Road
Lenexa, KS 66215-2802

Dear Mr. Clouette:

The National Laboratory Certification Program (NLCP) staff at RTI has reviewed the acceptability of Dr. Daniel Kolbow as an alternate (temporary) Responsible Person (altRP) for Clinical Reference Lab in Lenexa, KS. After review of submitted documentation and the inspectors' comments from the laboratory's 31st maintenance inspection on April 27-29, 2006, Dr. Kolbow has been found acceptable to serve as an alternate Responsible Person for the laboratory.

The NLCP must be apprised of and approve all changes in the RP status of the laboratory. Guidance outlining the conditions under which Dr. Kolbow may assume responsibility for the laboratory's operations under the Mandatory Guidelines is found in NLCP Program Document number twelve (PD 12).

If there are any questions concerning this matter, please contact me at (919) 541-7043.

Sincerely,

A handwritten signature in black ink, appearing to read "Michael R. Baylor".

Michael R. Baylor, Ph.D.
Co-Director, NLCP

ATTACHMENT E