RN Expertise, Inc 214 Hickman Dr. Ste# 102 Sanford, FL 32771 (866) 321-8611 toll free 407-321-8611 phone 407-321-6168 fax

RN Expertise, Inc.



To:	John Estep	From: Christine Steele	
Fax:	304-558-3970	Pages:	
Phone	N	Dates 4 28 29	
Re:		cci	
_			(1)
	CRFQ 0803 FOT2400	000085 Drug and Alcohol Testing Services due 4-29-24 @	Đ

1:30pm

• COMMENTS:

RECEIVED

2024 APR 29 AM 9: 12

WV PURCHASING DIVISION



RN Expertise, Inc.

"For Convenient and Quality Service"

April 27, 2024

WV Department of Administration Purchasing Division Attn: John Estep 2019 Washington Street East Charleston WV 25305

RE: CRFQ 0803 DOT2400000085

Dear Mr. Estep:

RN Expertise, Inc. is pleased to participate in the above referenced RFQ. RN Expertise, Inc. certifies that it has read the RFQ in its entirety and is able to meet all of the service requirements listed in the RFQ. All questions and answers have been reviewed. RN Expertise, Inc. has over 31 years experience in the drug testing industry and is very qualified to perform the drug and alcohol testing services required by the State of West Virginia.

RN Expertise, Inc. serves as a national third party administrator for drug testing programs for both private and governmental agencies. We provide DOT and non-DOT drug and alcohol testing services for over 1,200 clients and arrange off-site and on-site drug screen and breath alcohol testing services to meet the specific needs of each individual customer. RN Expertise, Inc. provides accurate and reliable services in a cost effective manner. RN Expertise, Inc. originated in 1993 and began performing paramedical exams and on-site drug testing. We began providing DOT drug testing services in 1995 when the Omnibus Transportation Testing Act of 1991 was implemented.

The President of RN Expertise, Inc. attended training offered by the individual Dr. Donna Smith who was one of the authors of CFR 49 Part 40 guidelines. Ms. Steele attended this training in 1994 and began implementing TPA services for Department of Transportation workplaces in the United States. We were one of the original TPA's who started providing these services when the program began. Ms. Steele served on the Board of the Drug and Alcohol Testing Industry Association during that time. She is a Certified Breath Alcohol Instructor and is certified by Intoximeter. Ms. Steele is also a Certified Department of Transportation Collector Trainer. She has trained hundreds of collectors and breath alcohol technicians over the years. Ms. Steele has attended DOT trainings throughout the years on any updates and changes in Federal guidelines. She also subscribed to the RED BOOK and receives all updates in Federal and state drug testing

guidelines through ODAPC. The staff at RN Expertise is updated on any changes in guidelines.

RN Expertise, Inc. is a Women Business Enterprise and is 100% owned by Ms. Christine Steele, RN. RN Expertise, Inc. is a Subchapter S Corporation and is a small business. Our Federal Id Tax number is 59-3172603. The principal place of business is:

RN Expertise, Inc.
214 Hickman Dr Ste 102
Sanford, FL 32771
(407) 321-8611
Fax (407) 321-6166
Contact: Christine Steele, President
Rnexinc@aol.com

The president of RN Expertise, Inc. prepared this proposal and is available to answer any questions that may arise during evaluation. It is RN Expertise's goal to provide the most affordable, expedient and quality services possible. We strive for excellence. We have an excellent reputation in the industry and take pride in customer service. Our staff is always available to assist our clients. RN Expertise, Inc. provides 24 hour turn around on negative results and 48-72 hour turn around on positive results.

The administrative staff of RN Expertise has over 22 years experience with the company. They are well educated on federal drug testing guidelines. They provide data entry, statistical reporting, billing, MRO assistance, customer service, random generation, and communicate with Ms. Steele on any customer service issues. The administrative staff communicates with clients and serves as a liaison between the laboratory and the clients to assist with any questions that occur regarding the drug testing process. RN Expertise, Inc. utilizes the Medical Review Officer services of Dr. Emily Vives. She has over 16 years experience as a Medical Review Officer and is certified by AAMRO.

RN Expertise, Inc. provides all services required by Federal, State, and non-regulated programs. We are very experienced in originating new programs and with arranging any form of drug alcohol testing services. We have national access to collection sites and have a 31 year relationship history with the major drug testing laboratories. RN Expertise also specializes in on-site testing and has a network of on-site testing partners located throughout the US.

As mentioned, we provide services for many different varieties of drug free workplaces. Some of our clients are: The Greater Orlando Aviation Authority, State of West Virginia DHHR, Trillium Driver Solutions, Arkansas DHS, The Kentucky State Police, The Kentucky Transportation Cabinet, and The Raleigh School District, to name a few. We provide services for Department of Transportation workplaces, State of Florida Drug Free workplaces and numerous non-Dot workplaces throughout the nation. It is our goal to provide an excellent turnkey drug and alcohol testing program for West Virginia DOT.

We utilize the laboratory services of Abbott formerly Alere, a DHHS SAMHSA certified laboratory and CRL. Abbott(formerly Alere) and CRL have many years of experience in forensic toxicology. All laboratory services for all drug test types and panels will be performed by Alere and CRL. We are contracted with Alere and CRL with a price agreement for services and the results are sent from the laboratory to our Medical Review Officers at RN Expertise, Inc.

RN Expertise appreciates the potential opportunity to provide services to the State of WV. If any questions arise during the evaluation period, I may be contacted at (407) 321-8611. It would be my pleasure to answer any questions and to assist you with the administration of your drug and alcohol testing requirements.

Sincerely,

Christine Steele

President RN Expertise, Inc. 214 Hickman Drive Ste 102

Sanford, FL 32771 (407) 321-8611



Department of Administration **Purchasing Division** 2019 Washington Street East Post Office Box 50130 Charleston, WV 25305-0130

State of West Virginia **Centralized Request for Quote** Medical

1394254

Doc Description: DRUG AND ALCOHOL TESTING 8324C000076

Reason for Modification:

ADDENDUM NO 1

Vendor Questions and

Responses

Bid Opening Moves to 04/29/2024

Proc Type:

Central Master Agreement

Date Issued

Solicitation Closes

Solicitation No

Version

2024-04-22

2024-04-29 13:30

CRFQ

0803 DOT2400000085

BID RECEIVING LOCATION

BID CLERK

DEPARTMENT OF ADMINISTRATION

PURCHASING DIVISION

2019 WASHINGTON ST E

CHARLESTON

WV 25305

US

VENDOR : 3

Vendor Customer Code: V5 00000 12883

Vendor Name: RN Expertise Inc.
Address: 214 Mickinsh Pr St. 102

5m sel

Street:

City:

State: F2

4.0

zlp: 32777

Principal Contact: MRISTING Stale

Vendor Contact Phone: 45 3M 8111

Extension:

FOR INFORMATION CONTACT THE BUYER

John W Estep 304-558-2566

john.w.estep@wv.gov

Vendor Signature

All offers subject to all terms and conditions contained in this solicitation

Date Printed: Apr 22, 2024

· Page: 1

FORM ID: WV-PRC-CRFQ-002 2020/05

ADDITIONAL INFORMATION

ADDENDUM NO 1

Addendum No_1 issued to publish and distribute the attached Information to the Vendor Community

REQUEST FOR QUOTATION:

The West Virginia Purchasing Division is soliciting bids on behalf of the West Virginia Department of Transportation (WVDOT) to establish an open-end contract for drug and alcohol testing services involving WVDOT agency employees, and individuals proposed to become WVDOT agency employees, in all 55 countles of the State of West Virginia. Per the Bid Requirements, Specifications, Terms and Conditions attached to this solicitation.

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2	On-Site Urine Collection - After Hours	10.00000	EA	21150	2160
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3	On-Site Alcohol (Breath) Test	250.00000	EA	10	2500.
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9	Scheduled Clinic Visit - Urine Collections	900.00000	EA	39.33	35,397

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11	Reasonable Suspicion Test-Urine Collection- Business Hours	5.00000	EA	340	172.50

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Line	Comm Ln Desc	Qty	Unit Issue	Unit Price	Total Price
12	Reasonable Suspicion Test-Urine Collection- After Hours	5.00000	EA	34.50	172.5t

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13	Reasonable Suspicion Test-Alcohol Breath Test-Business Hours	5.00000	EA	10	50.00

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14	Reasonable Suspicion Test-Alcohol Breath Test-After Hours	5.00000	EA	10	800

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SCHEDUIE:OBJEVENTIS	Section Line 1 - All All All All All All All All All A
Line Event Date 1 Tech Questions due by 10:00am 2024-04-10	

Exhibit A

Pricing Pages

All per costs are to be based upon an all-Inclusive collection as seen throughout the entire process of analysis, culminating with the certification of results and proper reporting of such results of the Human Resources Division or the appropriate Agency Program Manager. Alcohol and drug screening requires separate pricing. Regular hours testing (Monday - Friday, 6:30 am - 5:00 pm) and after hours testing (Saturday, Sunday, and weekdays 5:01 pm - 6:29 am) requires separate pricing. "On-Site" refers to urine or breath collections which are conducted at a DOH, Parkways, or State Rail work location.

The West Virginia Department of Transportation requests your bid on the following services and estimated quantities:

	Usage Sample Totals	Unit of Measure	. Cost Of Each Test/Item	Total For Each Line
1. On-Site Urine Collection	1400	pertest	34.5	48,300.00
On-Site Urine Colleciton- After Hours	10	per test	34.5	345.00
2. On-Site Alcohol (Breath) Test	250	per test	10	2,500.00
On-Site Alcohol (Breath) Test-After Hours	10	per test	10	100.00
3. Professional Services				
Collector Testimony	1	per day	0	0.00
Deposition	1	per day	0	0.00
Expert Witness Testimony	1	per day	0	0.00
Laboratory Litigation Packages	1	each	0	0.00
4. Scheduled Clinic Visits	20			ð
Urine Collections	900	per test	39.33	35,397.00
Alcohol (Breath) Test	20	per test	39.33	786.60
5. Reasonable Suspicion Testing -No notice given				
Urine Collection - during business hours	5	per test	34.5	172.50
Urine Colleciton - after hours	5	per test	34.5	172.50
Alcohol(Breath)Test - during business hours	5	per test	10	50.00
Alcohol (Breath) Test - after hours	5	per test	10	50.00
			Total	\$ 87,873.60

Include the name of the software or internet-based result reporting:		
ETEST	W	

The vendor shall be required to perform all of the services named above. Failure to provide the services and bid prices shall result in disqualification of the bid.

RN Eporter

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ADDENDUM ACKNOWLEDGEMENT FORM SOLICITATION NO.: CRFQ DOT2400000085

Instructions: Please acknowledge receipt of all addenda issued with this solicitation by completing this addendum acknowledgment form. Check the box next to each addendum received and sign below. Failure to acknowledge addenda may result in bid disqualification.

Acknowledgment: I hereby acknowledge receipt of the following addenda and have made the necessary revisions to my proposal, plans and/or specification, etc.

Addendum Numbers Received:

(Check the box next to each addendum received)

[X]	Addendum No. 1	ſ]	Addendum No. 6
[]	Addendum No. 2	ſ]	Addendum No. 7
[]	Addendum No. 3	ſ)	Addendum No. 8
[]	Addendum No. 4	1]	Addendum No. 9
[]	Addendum No. 5	ſ	1	Addendum No. 10

I understand that failure to confirm the receipt of addenda may be cause for rejection of this bid. I further understand that that any verbal representation made or assumed to be made during any oral discussion held between Vendor's representatives and any state personnel is not binding. Only the information issued in writing and added to the specifications by an official addendum is binding.

Company

Authorized Signature

Date

NOTE: This addendum acknowledgement should be submitted with the bid to expedite document processing.

AGREED:	
Name of Agency:	Name of Associate: W Cresty
Signature:	Signature: Www. XWW
Title:	Title: President
Date:	Date: 4/28/24

Form - WVBAA-012004 Amended 06.26.2013

APPROVED AS TO FORM THIS 20 17

Retrick Morrisey
Astomey General



Theodore F. Shults, MS, JD Chairman (919) 489-9588

American Association of Medical Review Officers

February 14, 2023

Emily Vives M.D.
Sunny Medical PL
11183 S Orange Blossom TrlSte D
Orlando, FL 32837

Dear Dr. Vives:

Thank you for participating in the AAMRO recertification examination. I am pleased to inform you that based on your examination results; you have met AAMRO's criteria for recertification.

Your AAMRO number is the same:
Your new certification expiration date is: 02/09/2028

If your subscription/membership to the online MRO Center at http://www.aamro.com is up-to-date please make sure it is activated! If not contact AAMRO to pay for the annual membership. You will find a scarchable database with back issues of MRO ALERT, state laws, federal regulations and guidance. This will be a valuable resource for your MRO practice.

If you do not know or have a Username to access the MRO Center, please send an email to bbrandon@aamro.com or call 800-489-1839. We can provide your Username only, because we do not have access to your Password.

Your name and phone number you provided are listed in the AAMRO Registry of Certified MROs on our website at www.samro.com. Enclosed is verification letter and CME documents. If you wish to make changes, you can contact us or change your record online using your MRO Center sign-in information.

An update sticker for your AAMRO wall certificate is attached to this letter. The verification letter, showing the dates of your certification and recertification, will be useful to present to employers, laboratories, and others who need to verify your MRO status.

Sincerely,

Theodore F. Shults, J.D., M.S.

Andre of Hufts

Chairman

Enclosures

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Center for Advancing Translational Sciences; Notice of Closed Meeting

Pursuant to section 1009 of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), title 5 U.S.C., as amended. The contract proposals and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the contract proposals, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Center for Advancing Translational Sciences Special Emphasis Panel; TRND 5: Pharmacology Studies, Animal Model Development & Related Services for Drug Development.

Date: November 18, 2023.
Time: 1:00 p.m. to 4:30 p.m.
Agenda: To review and evaluate contract

proposals.

Place: National Center for Advancing Translational Sciences, National Institutes of Health, 6701 Democracy Bivd., Room 1073, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: M. Lourdes Ponce, Ph.D., Scientific Review Officer, Office of Scientific Review, National Center for Advancing Translational Sciences, National Institutes of Health, 6701 Democrecy Blvd., Room 1073, Bethesda, MD 20892, 301–435–0810, lourdes.ponce@nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.859, Pharmacology, Physiology, and Biological Chemistry Research; 93.350, B—Cooperative Agreements; 93.859, Biomedical Research and Research Training, National Institutes of Health, HHS)

Dated: September 26, 2023. Melanis J. Pantoja,

Program Analyst, Office of Federal Advisory Committee Policy.

(FR Doc. 2023-21589 Filed 9-29-23; 8:45 am) BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Current List of HHS-Certified Laboratories and instrumented initial Testing Facilities Which Meet Minimum Standards To Engage in Urine and Oral Fluid Orag Testing for Federal Agencies

AGENCY: Substance Abuse and Mental Health Services Administration, HHS. ACTION: Notice.

SUMMARY: The Department of Health and Human Services (HHS) notifies federal agencies of the laboratories and Instrumented Initial Testing Facilities (IITFs) currently certified to meet the standards of the Mandatory Guidelines for Federal Workplace Drug Testing Programs using Urine or Oral Fluid (Mandatory Guidelines).

FOR FURTHER INFORMATION CONTACT:
Anastasia Flanagan, Division of
Workplace Programs, SAMHSA/CSAP,
5800 Fishers Lane, Room 16N06B,
Rockville, Maryland 20857; 240–276–
2600 (voice); Anastasia.Flanagan@
samhsa.hhs.gov (email).

SUPPLEMENTARY ENFORMATION: In accordance with Section 9.19 of the Mandatory Guidelines, a notice listing all currently HHS-certified laboratories and ITFs is published in the Federal Register during the first week of each month. If any laboratory or ITF certification is suspended or revoked, the laboratory or ITF will be omitted from subsequent lists until such time as it is restored to full certification under the Mandatory Guidelines.

if any laboratory or IITF has withdrawn from the HHS National Laboratory Certification Program (NLCP) during the past month, it will be listed at the end and will be omitted from the monthly listing thereafter.

This notice is also evailable on the internet et https://www.samhsa.gov/workplace/resources/drug-testing/certified-lab-list.

The Department of Health and Human Services (HHS) notifies federal agencies of the laboratories and Instrumented Initial Testing Facilities (ITFs) currently certified to meet the standards of the Mandatory Guidelines for Federal Workplace Drug Testing Programs (Mandatory Guidelines) using Urine and of the laboratories currently certified to meet the standards of the Mandatory Guidelines using Oral Fluid.

The Mandatory Guidelines using Urine were first published in the Federal Register on April 11, 1988 (53

FR 11970), and subsequently revised in the Federal Register on June 9, 1984 (59 FR 29908); September 30, 1997 (62 FR 51118); April 13, 2004 (69 FR 19644); November 25, 2008 (73 FR 71858); December 10, 2008 (73 FR 75122); April 30, 2010 (75 FR 22809); and on January 23, 2017 (82 FR 7920).

The Mandatory Guidelines using Oral Fluid were first published in the Federal Register on October 25, 2019 (84 FR 57554) with an effective date of

January 1, 2020.

The Mandatory Guidelines were initially developed in accordance with Executive Order 12564 and section 503 of Public Law 100–71 and allowed urine drug testing only. The Mandatory Guidelines using Urine have since been revised, and new Mandatory Guidelines allowing for oral fluid drug testing have been published. The Mandatory Guidelines require strict standards that laboratories and ITTs must meet in order to conduct drug and specimen validity tests on specimens for federal agencies. HHS does not allow ITTs to conduct oral fluid testing.

To become certified, an applicant laboratory or HTF must undergo three rounds of performance testing plus an on-site inspection. To maintain that certification, a laboratory or HTF must participate in a quarterly performance testing program plus undergo periodic,

on-site inspections.
Laboratories and ITFs in the applicant stage of certification are not to be considered as meeting the minimum requirements described in the HHS Mandatory Guidelines using Urine and/or Oral Fluid. An HHS-certified laboratory or ITF must have its letter of certification from HHS/SAMHSA (formerly: HHS/NIDA), which attests that the test facility has met minimum standards. HHS does not allow liTFs to conduct oral fluid testing.

HHS-Certified Laboratories Approved To Conduct Oral Fluid Drug Testing

In accordance with the Mandatory Guidelines using Oral Fluid dated October 25, 2019 (84 FR 57554), the following HHS-certified laboratories meet the minimum standards to conduct drug and specimen validity tests on oral fluid specimens:

At this time, there are no laboratories certified to conduct drug and specimen validity tests on oral fluid specimens.

HHS-Certified Instrumented Initial Testing Facilities Approved To Conduct Urine Drug Testing

In accordance with the Mandatory Guidelines using Urine dated January 23, 2017 (82 FR 7920), the following HHS-certified IITFs meet the minimum standards to conduct drug and specimen validity tests on urine specimens:

Dynacare, 6628 50th Street NW, Edmonton, AB Canada T6B 2N7, 780–784–1190 (Formerly: Gamma-Dynacare Medical Laboratories)

HHS-Certified Laboratories Approved To Conduct Urine Drug Testing

In accordance with the Mandatory Guidelines using Urine dated January 23, 2017 (82 FR 7920), the following HHS-certified laboratories meet the minimum standards to conduct drug and specimen validity tests on urine specimens:

Alere Toxicology Services, 1111 Newton St., Gretna, LA 70053, 504–361– 6969/800–433–3823 (Formerly: Kroll Laboratory Specialists, Inc., Laboratory Specialists, Inc.)

Alere Toxicology Services, 450
Southlake Blvd., Richmond, VA
23238, 804–378–9130 (Formerly:
Kroll Laboratory Specialists, Inc.,
Scientific Testing Laboratories, Inc.;
Kroll Scientific Testing
Laboratories, Inc.)

Clinical Reference Laboratory, Inc., 6433 Quivira Road, Lenexa, KS 66215— 2802, 800—445—6917

Desert Tox, LLC, 5425 E Bell Rd, Suite 125, Scottsdale, AZ 85254, 802-457-5411/623-748-5045

DrugScan, Inc., 200 Precision Road, Suite 200, Horsham, PA 19044, 800-235-4890

Dynacare*, 245 Pall Mall Street, London, ONT, Canada N6A 1P4, 519-679-1630 (Formerly: Gamma-Dynacare Medical Laboratories)

ElSohly Laboratories, Inc., 5 Industrial Park Drive, Oxford, MS 38655, 662-236-2609

Laboratory Corporation of America Holdings, 7207 N. Gessner Road, Houston, TX 77040, 713-858-8288/ 800-800-2387

Laboratory Corporation of America Holdings, 69 First Ave., Raritan, NJ 08669, 908-526-2400/600-437-4986 (Formerly: Roche Biomedical Laboratories, Inc.)

Laboratory Corporation of America
Holdings, 1904 TW Alexander
Drive, Research Triangle Park, NC
27709, 919-572-6900/800-8333984 (Formerly: LabCorp
Occupational Testing Services, Inc.,
CompuChem Laboratories, Inc., A
Subsidiary of Roche Biomedical
Laboratory; Roche CompuChem
Laboratories, Inc., A Member of the
Roche Group)

Laboratory Corporation of America Holdings, 1120 Main Street, Southaven, MS 36671, 866-8278042/800-233-6339 (Formerly; LabCorp Occupational Testing Services, Inc.; MedExpress/National Laboratory Center)

Laboratory Center)
LabOne, Inc. d/b/a Quest Diagnostics,
10101 Renner Blvd., Lenexa, KS
66219, 913-688-3927/800-8738645 (Formerly: Quest Diagnostics
Incorporated; LabOne, Inc.; Center
for Laboratory Services, a Division
of LabOne, Inc.)

Legacy Laboratory Services Toxicology, 1225 NE 2nd Ave., Portland, OR 97232, 503-413-5295/800-950-5295

MedTox Laboratories, Inc., 402 W. County Road D, St. Paul, MN 55112, 651-636-7466/800-832-3244

Minneapolis Veterans Affairs Medical Center, Forensic Toxicology Laboratory, 1 Veterans Drive, Minneapolis, MN 55417, 612–725– 2088. Testing for Veterans Affairs (VA) Employees Only

Pacific Toxicology Laboratories, 9348
DeSoto Ave., Chatsworth, CA
91311, 800–328–6942 (Formerly:
Centinela Hospital Airport
Toxicology Laboratory)

Phamatech, Inc., 15175 Innovation Drive, San Diego, CA 92128, 888-635-5840

Quest Diagnostics Incorporated, 400
Egypt Road, Norristown, PA 19403,
610-631-4600/677-642-2216
(Formerly; SmithKline Beecham
Clinical Laboratories; SmithKline
Bio-Science Laboratories)

US Army Forensic Toxicology Drug Testing Laboratory, 2490 Wilson St., Fort George G. Meade, MD 20755-5235, 301-677-7085, Testing for Department of Defense (DoD) Employees Only

 The Standards Council of Canada (SCC) voted to end its Laboratory Accreditation Program for Substance Abuse (LAPSA) effective May 12, 1998. Laboratories certified through that program were accredited to conduct forensic urine drug testing as required by U.S. Department of Transportation (DOT) regulations. As of that date, the certification of those accredited Canadian laboratories will continue under DOT authority. The responsibility for conducting quarterly performance testing plus periodic on-site inspections of those LAPSA-accredited laboratories was transferred to the U.S. HHS, with the HHS' NLCP contractor continuing to have an active role in the performance testing and laboratory inspection processes. Other Canadian laboratories wishing to be considered for the NLCP mey apply directly to the NLCP contractor just as U.S. laboratories do.

Upon finding a Canadian laboratory to be qualified, HHS will recommend that DOT certify the laboratory (Federal Register, July 16, 1998) as meeting the minimum standards of the Mandatory Guidelines published in the Federal Register on January 23, 2017 (82 FR 7920). After receiving DOT certification, the laboratory will be included in the monthly list of HHS-certified laboratories and participate in the NLCP certification maintenance program.

Anastasia D. Flanagan,

Public Health Advisor, Division of Workplace Programs.

[FR Dag. 2023-21669 Filed 9-29-29; 8:45 am] BILLING CODE 4168-89-P

DEPARTMENT OF HOMELAND SECURITY

Federal Emergency Management Agency

[Docket ID FEMA-2023-0002]

Changes in Flood Hazard Determinations

AGENCY: Federal Emergency
Management Agency, Department of
Homeland Security.
ACTION: Notice.

SUMMARY: New or modified Base (1percent annual chance) Flood Elevations (BFEs), base flood depths, Special Flood Hazard Area (SFHA) boundaries or zone designations, and/or regulatory floodways (hereinafter referred to as flood hazard determinations) as shown on the indicated Letter of Map Revision (LOMR) for each of the communities listed in the table below are finalized. Each LOMR revises the Flood Insurance Rate Maps (FIRMs), and in some cases the Flood Insurance Study (FIS) reports, currently in effect for the listed communities.

DATES: Each LOMR was finalized as in the table below.

ADDRESSES: Each LOMR is available for inspection at both the respective Community Map Repusitory address listed in the table below and online through the FEMA Map Service Center at https://msc.fema.gov.

FOR FURTHER NFORMATION CONTACT: Rick Sacbibit, Chief, Engineering Services Branch, Federal Insurance and Mitigation Administration, FEMA, 400 C Street SW, Washington, DC 20472, (202) 646–7659, or (email) patrick.sacbibit@fema.dhs.gov; or visit the FEMA Mapping and Insurance eXchange (FMIX) online at https://www.floodmaps.fema.gov/fhm/fmx_main.html.

REQUEST FOR QUOTATION Open End Contract for Drug and Alcohol Testing Services

- 8.3 Reports: Vendor shall provide to the Agency quarterly random testing reports showing the selected employees and shall provide by February 15th each year an annual summary, on a calendar year basis, reporting the number of drug and alcohol tests within each testing category. Annual drug and alcohol testing reports must show DOT-regulated employees' information separately from non-regulated employees' testing information. Failure to supply such reports may be grounds for cancellation of this Contract.
- 8.4 Contract Manager: During its performance of this Contract, Vendor shall designate and maintain a primary contract manager responsible for overseeing Vendor's responsibilities under this Contract. The Contract manager must be available during regular business hours to address any customer service or other issues related to this Contract. Vendor is to list its Contract manager and his or her contact information below.

Contract Manager: Telephone Number:

Fax Number:

Email Address:

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DESIGNATED CONTACT: Vendor appoints the individual identified in this Section as the Contract Administrator and the initial point of contact for matters relating to this Contract.
(Printed Name and Title) Aristone Steel
(Address) 214 Hickman Dn St. 102 SAMMy fr 327
(Phone Number) / (Fax Number) 407-32+8611 / 407 32/-6/11 (email address) RN 4100 (a ao 1.00m)
(email address) 12N SINC @ aolocom
CERTIFICATION AND SIGNATURE: By signing below, or submitting documentation through wvOASIS, I certify that: I have reviewed this Solicitation/Contract in its entirety; that I understand the requirements, terms and conditions, and other information contained herein; that this bid, offer or proposal constitutes an offer to the State that cannot be unilaterally withdrawn; that the product or service proposed meets the mandatory requirements contained in the Solicitation/Contract for that product or service, unless otherwise stated herein; that the Vendor accepts the terms and conditions contained in the Solicitation, unless otherwise stated herein; that I am submitting this bid, offer or proposal for review and consideration; that this bid or offer was made without prior understanding, agreement, or connection with any entity submitting a bid or offer for the same material, supplies, equipment or services; that this bid or offer is in all respects fair and without collusion or fraud; that this Contract is accepted or entered into without any prior understanding, agreement, or connection to any other entity that could be considered a violation of law; that I am authorized by the Vendor to execute and submit this bid, offer, or proposal, or any documents related thereto on Vendor's behalf; that I am authorized to bind the vendor in a contractual relationship; and that to the best of my knowledge, the vendor has properly registered with any State agency that may require registration.
By signing below, I further certify that I understand this Contract is subject to the provisions of West Virginia Code § 5A-3-62, which automatically voids certain contract clauses that violate State law; and that pursuant to W. Va. Code 5A-3-63, the entity
entering into this contract is prohibited from engaging in a boycott against Israel.
(Company) Met The
(Signature of Authorized Representative) Anshire Skeek President 4-2724
(Printed Name and Title of Authorized Representative) (Date)
(Phone Number) (Fax Number)
(Email Address)