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

JAN 8 - 2007

Request for Quotation

RFQ NUMBER	PAGE
LBS70426	1

ADDRESS CORRESPONDENCE TO ATTENTION OF:
ROBERTA WAGNER
304-558-0067

VENDOR
*709065635 800-224-6723
BIO RAD LABORATORIES
~~2000 ALFRED NOBEL DRIVE~~
6565 185th Avenue NE
~~HERCULES CA 94547~~
Redmond, WA 98052

1-22-07

SHIP TO
HEALTH AND HUMAN RESOURCES
BPH - LABORATORY SERVICES

167-ELEVENTH AVENUE
SOUTH CHARLESTON, WV
25303 304-558-3530

DATE PRINTED	TERMS OF SALE	SHIP VIA	F.O.B	FREIGHT TERMS
12/26/2006	Net 30 Days	Best Way	Destination	

BID OPENING DATE: 01/25/2007 BID OPENING TIME 01:30PM

LINE	QUANTITY	UOP	CAT NO	ITEM NUMBER	UNIT PRICE	AMOUNT
0001	100	EA		475-00-99-001	NO BID	NO BID

TEST FOR DETECTION OF HEPATITIS A (IGM ANTI-HAV)

(NOTE: BID WILL BE EVALUATED ON ESTIMATED QUANTITY X COST PER INDIVIDUAL TEST = THE ESTIMATED ANNUAL COST.)
BRAND/ITEM#:.....
UNIT PACKAGE:.....TEST KIT PRICE $.....

OPEN-END CONTRACT

CONTRACTOR SHALL PROVIDE REAGENTS AND TESTING EQUIPMENT FOR THE DETECTION OF HEPATITIS A, B, C AND/OR HIV-1 AND HIV-2 PLUS GROUP O IN SERUM SPECIMEN. CONTRACTOR SHALL PROVIDE A FULLY AUTOMATED ANALYZER AT NO ADDITIONAL CHARGE FOR USE WITH THE REAGENTS. THE ANALYZER SYSTEM INCLUDES A COMPUTER, MONITOR AND PRINTER, ETC. WHICH WILL BE RETAINED AND MAINTAINED BY THE VENDOR AND MUST MUST HAVE THE CAPABILITY OF INTERFACING WITH THE LIMS (LABORATORY INFORMATION MANAGEMENT SYSTEM).
NOTE: SEE ATTACHED SPECIFICATIONS

DELIVERY ADDRESS OF REAGENTS AND EQUIPMENT:
OFFICE OF LABORATORY SERVICES
ATTN: DONNA D. SMITH
167 11TH AVENUE
SOUTH CHARLESTON, WV 25303

RECEIVED
2007 JAN 23 A 11:43
PURCHASING DIVISION
STATE OF WV

0002	2,000	EA		475-00-99-001	$1,200.00	$4,800.00

TEST FOR DETECTION OF HEPATITIS B SURFACE ANTIGEN

4 - 480 Test/Kit

SEE REVERSE SIDE FOR TERMS AND CONDITIONS

SIGNATURE	TELEPHONE	DATE
Janette J. Stockert	800-666-8111 x1761	1/22/07

TITLE	FEIN	ADDRESS CHANGES TO BE NOTED ABOVE
Bid Coordination Supervisor	94-1381833	

WHEN RESPONDING TO RFQ, INSERT NAME AND ADDRESS IN SPACE ABOVE LABELED 'VENDOR'

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BID OPENING DATE: 01/25/2007 BID OPENING TIME 01:30PM

LINE	QUANTITY	UOP	CAT NO	ITEM NUMBER	UNIT PRICE	AMOUNT
				(HBS AG)		
				BRAND/ITEM#: Bio-Rad/Catalog #32591, 480 Test/Kit		
0003	10	EA		475-00-99-001	$225.00	$2,250.00
				TEST FOR DETECTION OF ANTIBODY TO HEPATITIS B		
				UNIT PACKAGE: 25 TEST KIT PRICE $225.00		
				SURFACE ANTIGEN (HUMAN) CONFIRMATORY ASSAY IN HUMAN SERUM, PLASMA OR CADAVERIC SERUM.		
				BRAND/ITEM#: Bio-Rad/Catalog #32594		
				UNIT PACKAGE: 25 TEST KIT PRICE $225.00		
0004	2,000	EA		475-00-99-001	NO BID	NO BID
				TEST FOR DETECTION OF TOTAL ANTIBODIES TO HEPATITIS		
				CORE ANTIGEN (ANTI-HBC)		
				BRAND/ITEM#:		
				UNIT PACKAGE: TEST KIT PRICE $		

SEE REVERSE SIDE FOR TERMS AND CONDITIONS

SIGNATURE	TELEPHONE	DATE
Janette J. Stockert	800-666-111 x1716	1/22/07
TITLE: Bid Coordination SUPERVISOR	FEIN: 94-1381833	ADDRESS CHANGES TO BE NOTED ABOVE

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LINE	QUANTITY	UOP	CAT NO.	ITEM NUMBER	UNIT PRICE	AMOUNT
0005	100	EA		475-00-99-001	NO BID	NO BID
	TEST FOR DETECTION OF IGM ANTIBODY TO HEPATITIS B CORE ANTIGEN (IGM ANTI-HBC)					
	BRAND/ITEM#:.....					
	UNIT PACKAGE:.....TEST				KIT PRICE $.....	
0006	1,000	EA		475-00-99-001	NO BID	NO BID
	TEST FOR DETECTION OF ANTIBODY TO HEPATITIS B SURFACE ANTIGEN (ANTI-HBS)					
	BRAND/ITEM#:.....					
	UNIT PACKAGE:.....TEST				KIT PRICE $.....	
0007	2,000	EA		475-00-99-001	$3,240.00	$12,960.00
	TEST FOR DETECTION OF ANTIBODY TO HEPATITIS C VIRUS (ANTIHCV)					4 - 480 Test/Kit
	BRAND/ITEM#:	Ortho/Cat.		#930740		
	UNIT PACKAGE:.....480 TEST				KIT PRICE $3,240.00	

SEE REVERSE SIDE FOR TERMS AND CONDITIONS

SIGNATURE	TELEPHONE	DATE
Janette J. Stockert	800-666-8111 x1761	1/22/07

TITLE	FEIN	ADDRESS CHANGES TO BE NOTED ABOVE
BID COORDINATION SUPERVISOR	94-1381833	

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0008	7,000	EA		475-00-99-001	$1,800.00	$27,000.00

TEST FOR DETECTION OF HIV-1 AND HIV-2 PLUS O GROUP

15 - 480 Test/Kit

BRAND/ITEM#: Bio-Rad/Catalog #32588
UNIT PACKAGE: 480 TEST KIT PRICE $ 1,800.00
THE NUMBER OF TESTS REQUESTED ARE FOR BIDDING PURPOSES ONLY, AND THE VENDOR WILL BE REQUIRED TO PROVIDE ONLY THE QUANITITY NEEDED, BE IT MORE OR LESS. TEST KITS AR TO BE SHIPPED AS REQUESTED. REAGENTS ARE TO BE SHIPPED WITHIN THREE (3) DAYS OF RECEIVING AN ORDER.

THE TEST KIT MUST HAVE A MINIMUM SHELF LIFE OF NINETY (90) DAYS OR MORE BEYOND DATE OF RECEIPT.

ALL PRODUCTS AND EQUIPMENT ARE TO BE QUOTED FOB DESTINATION, UNLESS OTHERWISE STATED IN VENDOR'S QUOTATION.

EXHIBIT 3

LIFE OF CONTRACT: THIS CONTRACT BECOMES EFFECTIVE ON ..... AND EXTENDS FOR A PERIOD OF ONE (1) YEAR OR UNTIL SUCH "REASONABLE TIME" THEREAFTER AS IS NECESSARY TO OBTAIN A NEW CONTRACT OR RENEW THE ORIGINAL CONTRACT. THE "REASONABLE TIME" PERIOD SHALL NOT EXCEED TWELVE (12) MONTHS. DURING THIS "REASONABL TIME" THE VENDOR MAY TERMINATE THIS CONTRACT FOR ANY REASON UPON GIVING THE DIRECTOR OF PURCHASING 30 DAYS WRITTEN NOTICE.

UNLESS SPECIFIC PROVISIONS ARE STIPULATED ELSEWHERE IN THIS CONTRACT DOCUMENT, THE TERMS, CONDITIONS AND

SEE REVERSE SIDE FOR TERMS AND CONDITIONS

SIGNATURE		TELEPHONE	DATE
Janette Stockert	JANETTE J. STOCKERT	800-666-8111 x1761	1/22/07

TITLE	FEIN	
BID COORDINATION SUPERVISOR	94-1381833	ADDRESS CHANGES TO BE NOTED ABOVE

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PRICING SET HEREIN ARE FIRM FOR THE LIFE OF THE CONTRACT.

RENEWAL: THIS CONTRACT MAY BE RENEWED UPON THE MUTUAL WRITTEN CONSENT OF THE SPENDING UNIT AND VENDOR, SUBMITTED TO THE DIRECTOR OF PURCHASING THIRTY (30) DAYS PRIOR TO THE EXPIRATION DATE. SUCH RENEWAL SHALL BE IN ACCORDANCE WITH THE TERMS AND CONDITIONS OF THE ORIGINAL CONTRACT AND SHALL BE LIMITED TO TWO (2) ONE (1) YEAR PERIODS.

CANCELLATION: THE DIRECTOR OF PURCHASING RESERVES THE RIGHT TO CANCEL THIS CONTRACT IMMEDIATELY UPON WRITTEN NOTICE TO THE VENDOR IF THE COMMODITIES AND/OR SERVICES SUPPLIED ARE OF AN INFERIOR QUALITY OR DO NOT CONFORM TO THE SPECIFICATIONS OF THE BID AND CONTRACT HEREIN.

OPEN MARKET CLAUSE: THE DIRECTOR OF PURCHASING MAY AUTHORIZE A SPENDING UNIT TO PURCHASE ON THE OPEN MARKET, WITHOUT THE FILING OF A REQUISITION OR COST ESTIMATE, ITEMS SPECIFIED ON THIS CONTRACT FOR IMMEDIATE DELIVERY IN EMERGENCIES DUE TO UNFORESEEN CAUSES (INCLUDING BUT NOT LIMITED TO DELAYS IN TRANSPORTATION OR AN UNANTICIPATED INCREASE IN THE VOLUME OF WORK.)

QUANTITIES: QUANTITIES LISTED IN THE REQUISITION ARE APPROXIMATIONS ONLY, BASED ON ESTIMATES SUPPLIED BY THE STATE SPENDING UNIT. IT IS UNDERSTOOD AND AGREED THAT THE CONTRACT SHALL COVER THE QUANTITIES ACTUALLY ORDERED FOR DELIVERY DURING THE TERM OF THE CONTRACT, WHETHER MORE OR LESS THAN THE QUANTITIES SHOWN.

ORDERING PROCEDURE: SPENDING UNIT SHALL ISSUE A WRITTEN STATE CONTRACT ORDER (FORM NUMBER WV-39) TO THE VENDOR FOR COMMODITIES COVERED BY THIS CONTRACT,

SEE REVERSE SIDE FOR TERMS AND CONDITIONS

SIGNATURE		TELEPHONE	DATE
Janette Stockert	JANETTE J. STOCKERT	800-666-8111 x1761	1/22/07

TITLE	FEIN	ADDRESS CHANGES TO BE NOTED ABOVE
BID COORDINATION SUPERVISOR	94-1381833	

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P-2207

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EXCEPT WHEN PURCHASES ARE OF A DOLLAR AMOUNT ALLOWABLE TO BE MADE WITH THE WV STATE CREDIT CARD (P-CARD).

BANKRUPTCY: IN THE EVENT THE VENDOR/CONTRACTOR FILES FOR BANKRUPTCY PROTECTION, THIS CONTRACT IS AUTOMATICALLY NULL AND VOID, AND IS TERMINATED WITHOUT FURTHER ORDER.

THE TERMS AND CONDITIONS CONTAINED IN THIS CONTRACT SHALL SUPERSEDE ANY AND ALL SUBSEQUENT TERMS AND CONDITIONS WHICH MAY APPEAR ON ANY ATTACHED PRINTED DOCUMENTS SUCH AS PRICE LISTS, ORDER FORMS, SALES AGREEMENTS OR MAINTENANCE AGREEMENTS, INCLUDING ANY ELECTRONIC MEDIUM SUCH AS CD-ROM.

REV. 04/11/2001

VENDOR PREFERENCE CERTIFICATE

CERTIFICATION AND APPLICATION* IS HEREBY MADE FOR PREFERENCE IN ACCORDANCE WITH WEST VIRGINIA CODE, 5A-3-37 (DOES NOT APPLY TO CONSTRUCTION CONTRACTS).

A. APPLICATION IS MADE FOR 2.5% PREFERENCE FOR THE REASON CHECKED:

( ) BIDDER IS AN INDIVIDUAL RESIDENT VENDOR AND HAS RESIDED CONTINUOUSLY IN WEST VIRGINIA FOR FOUR (4) YEARS IMMEDIATELY PRECEDING THE DATE OF THIS CERTIFICATION; OR

SEE REVERSE SIDE FOR TERMS AND CONDITIONS

SIGNATURE	TELEPHONE	DATE
Janette Stockert JANETTE J. STOCKERT	800-666-8111 x1761	1/22/07

TITLE	FEIN	ADDRESS CHANGES TO BE NOTED ABOVE
BID COORDINATION SUPERVISOR	94-1381833	

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( ) BIDDER IS A PARTNERSHIP, ASSOCIATION OR CORPORATION RESIDENT VENDOR AND HAS MAINTAINED ITS HEADQUARTERS OR PRINCIPAL PLACE OF BUSINESS CONTINUOUSLY IN WEST VIRGINIA FOR FOUR (4) YEARS IMMEDIATELY PRECEDING THE DATE OF THIS CERTIFICATION; OR 80% OF THE OWNERSHIP INTEREST OF BIDDER IS HELD BY ANOTHER INDIVIDUAL, PARTNERSHIP, ASSOCIATION OR CORPORATION RESIDENT VENDOR WHO HAS MAINTAINED ITS HEADQUARTERS OR PRINCIPAL PLACE OF BUSINESS CONTINUOUSLY IN WEST VIRGINIA FOR FOUR (4) YEARS IMMEDIATELY PRECEDING THE DATE OF THIS CERTIFICATION; OR

( ) BIDDER IS A CORPORATION NONRESIDENT VENDOR WHICH HAS AN AFFILIATE OR SUBSIDIARY WHICH EMPLOYS A MINIMUM OF ONE HUNDRED STATE RESIDENTS AND WHICH HAS MAINTAINED ITS HEAQUARTERS OR PRINCIPAL PLACE OF JUSINESS WITHIN WEST VIRGINIA CONTINUOUSLY FOR THE FOUR (4) YEARS IMMEDIATELY PRECEDING THE DATE OF THIS CERTIFICATION.

B. APPLICATION IS MADE FOR 2.5% PREFERENCE FOR THE REASON CHECKED:

( ) BIDDER IS A RESIDENT VENDOR WHO CERTIFIES THAT, DURING THE LIFE OF THE CONTRACT, ON AVERAGE AT LEAST 75% OF THE EMPLOYEES WORKING ON THE PROJECT BEING BID ARE RESIDENTS OF WEST VIRGINIA WHO HAVE RESIDED IN THE STATE CONTINUOUSLY FOR THE TWO YEARS IMMEDIATELY PRECEDING SUBMISSION OF THIS BID;
OR
( ) BIDDER IS A NONRESIDENT VENDOR EMPLOYING A MINIMUM OF ONE HUNDRED STATE RESIDENTS OR IS A NONRESIDENT VENDOR WITH AN AFFILIATE OR SUBSIDIARY WHICH MAINTAINS ITS HEADQUARTERS OR PRINCIPAL PLACE OF BUSINESS WITHIN WEST VIRGINIA EMPLOYING A MINIMUM

SEE REVERSE SIDE FOR TERMS AND CONDITIONS

SIGNATURE	TELEPHONE	DATE
Janette Stockert JANETTE J. STOCKERT	800-666-8111 x1761	1/22/07

TITLE	FEIN	ADDRESS CHANGES TO BE NOTED ABOVE
BID COORDINATION SUPERVISOR	94-1381833	

WHEN RESPONDING TO RFQ, INSERT NAME AND ADDRESS IN SPACE ABOVE LABELED 'VENDOR'

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OF ONE HUNDRED STATE RESIDENTS WHO CERTIFIES THAT, DURING THE LIFE OF THE CONTRACT, ON AVERAGE AT LEAST 75% OF THE EMPLOYEES OR BIDDERS' AFFILIATE'S OR SUBSIDIARY'S EMPLOYEES ARE RESIDENTS OF WEST VIRGINIA WHO HAVE RESIDED IN THE STATE CONTINUOUSLY FOR THE TWO YEARS IMMEDIATELY PRECEDING SUBMISSION OF THIS BID.

BIDDER UNDERSTANDS IF THE SECRETARY OF TAX & REVENUE DETERMINES THAT A BIDDER RECEIVING PREFERENCE HAS FAILED TO CONTINUE TO MEET THE REQUIREMENTS FOR SUCH PREFERENCE, THE SECRETARY MAY ORDER THE DIRECTOR OF PURCHASING TO: (A) RESCIND THE CONTRACT OR PURCHASE ORDER ISSUED; OR (B) ASSESS A PENALTY AGAINST SUCH BIDDER IN AN AMOUNT NOT TO EXCEED 5% OF THE BID AMOUNT AND THAT SUCH PENALTY WILL BE PAID TO THE CONTRACTING AGENCY OR DEDUCTED FROM ANY UNPAID BALANCE ON THE CONTRACT OR PURCHASE ORDER.

BY SUBMISSION OF THIS CERTIFICATE, BIDDER AGREES TO DISCLOSE ANY REASONABLY REQUESTED INFORMATION TO THE PURCHASING DIVISION AND AUTHORIZES THE DEPARTMENT OF TAX AND REVENUE TO DISCLOSE TO THE DIRECTOR OF PURCHASING APPROPRIATE INFORMATION VERIFYING THAT BIDDER HAS PAID THE REQUIRED BUSINESS TAXES, PROVIDED THAT SUCH INFORMATION DOES NOT CONTAIN THE AMOUNTS OF TAXES PAID NOR ANY OTHER INFORMATION DEEMED BY THE TAX COMMISSIONER TO BE CONFIDENTIAL.

UNDER PENALTY OF LAW FOR FALSE SWEARING (WEST VIRGINIA CODE 61-5-3), BIDDER HEREBY CERTIFIES THAT THIS CERTIFICATE IS TRUE AND ACCURATE IN ALL RESPECTS; AND THAT IF A CONTRACT IS ISSUED TO BIDDER AND IF ANYTHING CONTAINED WITHIN THIS CERTIFICATE CHANGES DURING THE TERM OF THE CONTRACT, BIDDER WILL NOTIFY THE PURCHASING DIVISION IN WRITING IMMEDIATELY.

SEE REVERSE SIDE FOR TERMS AND CONDITIONS

SIGNATURE	TELEPHONE	DATE
Janette Stockert JANETTE J. STOCKERT	800-666-8111x1761	1/22/07
TITLE bid coordination supervisor / FEIN 94-1381833		ADDRESS CHANGES TO BE NOTED ABOVE

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BIDDER: Bio-Rad Laboratories, Inc.

DATE: 1/22/07

SIGNED: Janette J. Stockert

Janette J. Stockert

TITLE: Bid Coordination Supervisor

* CHECK ANY COMBINATION OF PREFERENCE CONSIDERATION(S) IN EITHER "A" OR "B", OR BOTH "A" AND "B" WHICH YOU ARE ENTITLED TO RECEIVE. YOU MAY REQUEST UP TO THE MAXIMUM 5% PREFERENCE FOR BOTH "A" AND "B".
(REV. 12/00)

NOTICE

AN ORIGINAL, SIGNED BID MUST BE SUBMITTED TO:

DEPARTMENT OF ADMINISTRATION
PURCHASING DIVISION
BUILDING 15
2019 WASHINGTON STREET, EAST
CHARLESTON, WV 25305-0130

SEALED BID

SEE REVERSE SIDE FOR TERMS AND CONDITIONS

SIGNATURE	TELEPHONE	DATE
Janette J. Stockert JANETTE J. STOCKERT	800-666-8111 x1761	1/22/07

TITLE	FEIN	ADDRESS CHANGES TO BE NOTED ABOVE
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BUYER: RW-22
RFQ. NO.: LBS70426
BID OPENING DATE: 01/25/2007
BID OPENING TIME: 1:30 P.M.

PLEASE PROVIDE A FAX NUMBER IN CASE IT IS NECESSARY TO CONTACT YOU REGARDING YOUR BID:

FAX #.....425-498-1757.....

CONTACT PERSON:.....Janette J. Stockert.....
...800-666-8111 x1761.....

THE MODEL/BRAND/SPECIFICATIONS NAMED HEREIN ESTABLISH THE ACCEPTABLE LEVEL OF QUALITY ONLY AND ARE NOT INTENDED TO REFLECT A PREFERENCE OR FAVOR ANY PARTICULAR BRAND OR VENDOR. VENDORS WHO ARE BIDDING ALTERNATES SHOULD SO STATE AND INCLUDE PERTINENT LITERATURE AND SPECIFICATIONS. FAILURE TO PROVIDE INFORMATION FOR ANY ALTERNATES MAY BE GROUNDS FOR REJECTION OF THE BID. THE STATE RESERVES THE RIGHT TO WAIVE MINOR IRREGULARITIES IN BIDS OR SPECIFICATIONS IN ACCORDANCE WITH SECTION 148-1-4(F) OF THE WEST VIRGINIA LEGISLATIVE RULES AND REGULATIONS.

INQUIRIES
WRITTEN QUESTIONS SHALL BE ACCEPTED THROUGH CLOSE OF BUSINESS ON WEDNESDAY, JAN. 10, 2007. QUESTIONS MAY BE SENT VIA USPS, FAX, COURIER OR EMAIL. IN ORDER TO ASSURE NO VENDOR RECEIVES AN UNFAIR ADVANTAGE, NO

SEE REVERSE SIDE FOR TERMS AND CONDITIONS

SIGNATURE		TELEPHONE	DATE
Janette J. Stockert	JANETTE J. STOCKERT	800-666-8111 x1761	1/22/07
TITLE: BID COORDINATION SUPERVISOR	FEIN: 94-1381833	ADDRESS CHANGES TO BE NOTED ABOVE	

WHEN RESPONDING TO RFQ, INSERT NAME AND ADDRESS IN SPACE ABOVE LABELED 'VENDOR'

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

Request for Quotation

RFQ NUMBER	PAGE
LBS70426	11

ADDRESS CORRESPONDENCE TO ATTENTION OF:
ROBERTA WAGNER
304-558-0067

VENDOR
*709065635 800-224-6723
BIO RAD LABORATORIES
~~2000 ALFRED NOBEL DRIVE~~
6565 185th Avenue NE
~~HERCULES CA 94547~~
Redmond, WA 98052
1-22-07

SHIP TO
HEALTH AND HUMAN RESOURCES
BPH - LABORATORY SERVICES

167-ELEVENTH AVENUE
SOUTH CHARLESTON, WV
25303 304-558-3530

DATE PRINTED	TERMS OF SALE	SHIP VIA	F.O.B.	FREIGHT TERMS
12/26/2006	Net 30 Days	Best WAy	Destination	

BID OPENING DATE: 01/25/2007 BID OPENING TIME 01:30PM

LINE	QUANTITY	UOP	CAT NO	ITEM NUMBER	UNIT PRICE	AMOUNT
	SUBSTANTIVE QUESTIONS WILL BE ANSWERED ORALLY. IF POSSIBLE, E-MAIL QUESTIONS ARE PREFERRED. ADDRESS INQUIRIES TO: ROBERTA WAGNER DEPARTMENT OF ADMINISTRATION PURCHASING DIVISION 2019 WASHINGTON STREET, EAST CHARLESTON, WV 25311 FAX: 304-558-4115 ~~E-MAIL:~~ RWAGNER@WVADMIN.GOV					
	PLEASE NOTE THAT WE PREFER FOB DESTINATION, IF YOU WISH TO BID OTHER THAN FOB DESTINATION, YOU MUST PROVIDE THE MAXIMUM SHIPPING COST SO THAT IT CAN BE ADDED TO THE BID. PREPAY TRANSPORTATION AND ADD TO INVOICE AS A SEPARATE ITEM. IF BY FREIGHT, ATTACH ORIGINAL PREPAID FREIGHT BILL.					
	****** THIS IS THE END OF RFQ LBS70426 ****** TOTAL:					$47,010.00

SEE REVERSE SIDE FOR TERMS AND CONDITIONS

SIGNATURE		TELEPHONE	DATE
Janette J. Stockert	JANETTE J. STOCKERT	800-666-8111 x1761	1/22/07

TITLE	FEIN	
BID COORDINATION SUPERVISOR	94-1381833	ADDRESS CHANGES TO BE NOTED ABOVE

WHEN RESPONDING TO RFQ, INSERT NAME AND ADDRESS IN SPACE ABOVE LABELED 'VENDOR'

A. REAGENT SPECIFICATIONS

1. Reagents must be bar-coded.
2. Must have ability to support up to 31 controls and calibrators.
3. Must be able to support multi-size reagent bottle volumes (milliliters) 3, 8, 15, 30, 60 and 125.
4. Must have lot and expiration management for reagents.
5. Must have disposable graphite tips, 300-1000 microliters.
6. Must have at least an on-board capacity of 480 tips (5 boxes of 96 tips).
7. Must be able to track tip usage on instrument and use partially filled tip racks.
8. Must have tip waste capacity of >1000 tips.
9. Must have mechanical tip detection.
10. Must have liquid level and clot detection.
11. Positive sample identification must be on plates and reagents.
12. The test kit must have a minimum shelf life of 90 days or more beyond date of receipt.

INDIVIDUAL TEST TYPE & SPECIFICATIONS FOR EACH TEST

Item # 1 Assay specifications for detection of Hepatitis A (IgM anti-HAV)

a. Must detect IgM Antibodies to Hepatitis A virus in human serum or plasma.
b. Must meet the following criteria for a qualitative assay of IgM Antibody to Hepatitis A Virus.
    i. Must have a 96-well test plate.
    ii. Must use monoclonal antibodies to human IgM as the basis for the enzyme immunoassay.
    iii. Must be an FDA approved method.
    iv. Must have positive agreement for acute profile samples of 100% with FDA approved comparator methods.
    v. Must have negative agreement for past infection profile samples of 100% with FDA approved comparator methods.
c. The principle must be direct antibody, sandwich EIA in a solid phase microwell.
d. Sample size must not be greater than 100 microliters.
e. All steps in the method must be adaptable to automation.

Item # 2 Assay specifications for detection of Hepatitis B Surface Antigen (HBsAg)

a. Must detect Hepatitis B Surface Antigen in human serum, plasma, or cadaveric serum.
b. Must meet the following criteria for a qualitative enzyme immunoassay of Hepatitis B Surface Antigen.
    i. Must have a 96-well test plate.
    ii. Must use a monoclonal antibody to Hepatitis B Surface Antigen.
    iii. Must be a FDA approved method.
    iv. Must have a specificity of at least 99%.
c. The principle must be direct antibody, sandwich EIA in a solid phase microwell.
d. Sample size must not be greater than 100 microliters.
e. All steps in the method must be adaptable to automation.

Item # 3 Assay specifications for detection of antibody to Hepatitis B Surface Antigen (human) confirmatory assay in human serum, plasma or cadaveric serum.

a. Must be a confirmatory neutralization of HBsAg enzyme immunoassay reactive specimens.
b. Must be compatible with the enzyme immunoassay used to detect antibody to Hepatitis B Surface Antigen assay.
c. Must be FDA approved.
d. All steps in the method must be adaptable to automation.

Item # 4 Assay specifications for detection of total antibodies to Hepatitis B Core Antigen (Anti-HBc).

a. Must detect total antibodies to Hepatitis B Core Antigen in human serum or plasma.
b. Must meet the following criteria for a qualitative assay of total antibodies to Hepatitis B Core Antigen.
    i. Must have a 96-well test plate.
    ii. Must use a monoclonal antibody directed to the immunodominant of Hepatitis B Core Antigen as the basis for the enzyme immunoassay.
    iii. Must be a FDA approved method.
    iv. Must have an analytical sensitivity of 0.4 u/ml using single-point serial dilutions of a standard preparation from the Paul-Ehrlich-Institute (PEI) or better.
c. The principle must be direct antibody, sandwich EIA in a solid phase microwell.
d. Sample size must not be greater than 100 microliters.
e. All steps in the method must be adaptable to automation.

Item # 5 Assay specifications for the detection of IgM Antibody to Hepatitis B Core Antigen (IgM anti-HBc)

a. Must detect IgM Antibody to Hepatitis B Core Antigen in human serum or plasma.
b. Must meet the following criteria for a qualitative assay of IgM antibody to Hepatitis B Core Antigen.
    i. Must have a 96-well test plate.
    ii. Must use monoclonal antibody to human IgM as the basis for the enzyme immunoassay.
    iii. Must be a FDA approved method.
    iv. Must agree with 95% exact confidence intervals with reference IgM Anti-HBc assay results.
    v. Must have an analytical sensitivity of 25 u/ml using single-point serial dilutions of a standard preparation from the Paul-Ehrlich-Institute (PEI) or better.
c. The principle must be direct antibody, sandwich EIA in a solid phase microwell. Sample size must not be greater than 100 microliters.
d. All steps in the method must be adaptable to automation.

Item # 6 Assay specifications for detection of antibody to Hepatitis B Surface Antigen (Anti-HBs)

a. Must detect antibodies to Hepatitis B Surface Antigen in human serum or plasma.
b. Must meet the following criteria for a qualitative assay of antibodies to Hepatitis B Surface Antigen.
    i. Must have a 96-well test plate.
    ii. Must be a direct, non-competitive enzyme inmmunoassay based on the use of polystyrene microwells coated with recombinant HBsAg.
    iii. Must be a FDA approved method.
    iv. Must show percent agreement with 95% exact confidence intervals with the reference Anti-HBs Assay results.
c. Sample size must not be greater than 100 microliters.
d. All steps in the method must be adaptable to automation.

Item # 7 Assay specifications for the detection of antibody to Hepatitis C Virus (Anti-HCV).

a. Must detect antibodies to Hepatitis C Virus in human serum or plasma.
b. Must meet the following criteria for a qualitative assay of antibody to Hepatitis C Virus.
    i. Must utilize microwells coated with recombinant Hepatitis C Virus encoded antigens as a solid phase in a 96-well plate.
    ii. Must use HCV recombinant proteins derived from the core, NS3, NS4 and NS5 region of the HCVgenome.
    iii. Must be a FDA approved method.
    iv. Must have a specificity of 99.95% or better in low prevalence population.
c. Must be an enzyme-linked, immunosorbent assay.
d. Must not require sample size greater than 100 microliters.
e. All steps in the method must be adaptable to automation.

Item #8 Assay Specifications for HIV-1 and HIV-2 plus O Group

a. Must detect antibodies to HIV-1 and 2 plus Group O in serum, plasma and cadaveric samples.
b. All vendors must provide supporting documentation that qualifies they are bidding an approved test for detection of both HIV-1 and 2.
c. Must meet the following criteria for HIV ½ recombinant DNA/synthetic peptide assay.
d. Must have a 96-well test plate.
e. HIV ½ peptide kit must have the following:
    i. Synthetic Peptide Immunoassay for the detection of the antibody to HIV-1 and HIV-2.
    ii. The microtiter wells are coated with a mixture of peptides; env and pol sequences for HIV-1 and HIV-2.
    iii. Sample dilution 1/10.
    iv. Must be FDA licensed recombinant peptide EIA for HIV-1 and HIV-2 plus Group O.
f. The principle must be direct antibody, sandwich Elisa in a solid phase microwell.
g. Sample size must not be greater than 75ul.
h. Turnaround time must not be greater than 3 hours for the HIV assay.
i. All steps in the method must be automated, including data reduction on one primary microplate instrument.
j. Incubation times (on the instrument) must not exceed (in minutes) 60-30-30.

k. Chromogen should not be lot specific for kit.
l. Stop solution must be ready to use.

B. INSTRUMENT SPECIFICATIONS

General Instrument Specifications

1. Must have primary sample capacity of 180 samples.
2. Must have 20 tubes per sample linear rack.
3. Must have Positive Identification for samples, microplates and reagents.
4. Must be able to sample from tubes up to 16mm diameter
5. Must be able to sample from tubes up to 100mm height.
6. Dead volume cannot be greater than 200 microliters.
7. All sample positions must be bar-coded on the sample tube and sample rack.
8. All reagent and quality control racks must be bar-coded.
9. Must have dilution capacity via tubes and microplate.
10. Sample dilution must be 1:10,000 or less.
11. Must have the capability to load continuously throughout the sample processing
12. Must include computer system and software.
13. Must be able to shake assays for variable times.
14. Must be able to process blood virus, infectious disease and autoimmune in a single run.
15. Must be able to incubate assays at Room Temperature (R.T.) and at 37 degrees C.

Washer Specifications

1. Must have an 8 channel manifold.
2. Must be able to use flat, U and V shaped plate bottom shapes.
3. Must have a plate and strip wash mode.
4. Must have variable wash cycles of 1 to 9.
5. Must have plate soak time of 0-999 secs.
6. Must have wash buffers with level sensors of 2 x 2L and 2 x 1L.
7. Must have waste capacity with level sensors of 1 x 10L.

Reader Specifications

1. Must have 8 channel read head.
2. Read time for full plate must not be greater than 15 seconds.
3. Must have a halogen light source.
4. Must have a reading range of up to 3.5 Optical Density (O.D.).
5. Must be equipped with at least 8 filter wheels to include 405, 450,492, 550, 620, 690 nm.
6. Must have an over-range filter.
7. Must have linearity (0-3.0 O.D.) to 1%; Precision (0-2.0 O.D.) to 2.5%.

C. COMPUTER INTERFACE SPECIFICATIONS

1. Must have ability to connect multiple instruments (up to 8) to a LAN (Local Area Network) and use one computer interface to interface to the facilities LIMS (Laboratory Information Management System) provider.
2. Interface cost must be included in cost per test.
3. Must have bi-directional interface with ASTM or ASCH file format.
4. User interface must be Windows 2000 operating system.
5. Must be able to track reactive results and perform duplicate assays prior to confirmation.

6. Must create a Primary Sample Validation Screen and Worklist for reactive samples.
7. Must be able to process and send repeat Worklist to instrument for analysis.
8. Must be able to check assay results to see if they are final (repeat reactive) or require repeating (initially reactive).
9. Once all results (both initial and repeat reactive) have been validated by the system, the final results must have the capability of being exported to the LIMS.
10. Vendor must be willing to assist in transition process to the LIMS.

D. TRAINING / INSTALLATION REQUIREMENTS

1. Vendor must provide a company representative for installation and training. Subcontracting of these services shall not be acceptable to the State of West Virginia. Any vendor responding to this contract that proposes to utilize a subcontractor shall not be considered during the award process.
2. Installation and training for equipment must be completed within 6 weeks of delivery date and must include one (1) key operator training at vendor's training site at vendor's expense.

E. EQUIPMENT OWNERSHIP / MAINTENANCE / TECHNICAL ASSISTANCE REQUIREMENTS

1. Vendor will retain ownership of all instrumentation.
2. All instrumentation provided by the selected vendor must be maintained at vendor's expense during the term of this contract. One (1) annual preventive maintenance visit at the laboratory site must be provided at no additional charge.
3. Vendor must provide a company representative for technical service, repairs, maintenance, etc. Any vendor responding to this contract that proposes to utilize a subcontractor shall not be considered during the award process.
4. Technical assistance must be available by telephone during normal business hours, 8:00 a.m. to 5:00 p.m. EST, Monday through Friday. If technical assistance does not resolve problems, replacement parts or loaner modules must be provided or on-site representative presence must be made available within 24 hours, except on weekends.

DELIVERY / SHIPPING REQUIREMENTS:

1. F.O.B. Destination unless vendor states otherwise in submitted quotation.
2. Reagents must be shipped no more than 3 days after receiving order.

ORDERING PROCEDURE:

Spending unit shall issue a written state contract order (Form Number WV-39) to the vendor for commodities covered by this contract, except when purchases are of a dollar amount allowable to be made with the WV State Credit Card (P-Card).

ATTACHMENT
P.O.# LBS70426

This agreement constitutes the entire agreement between the parties, and there are no other terms and conditions applicable to the licenses granted hereunder.

Agreed

1/22/07	
Signature Date	Signature Date
Janette J. Stockert	
Bid Coordination Supervisor	
Title	Title
Bio-Rad Laboratories, Inc.	
Company Name	Agency-Division

RFQ No. LBS70426

AFFIDAVIT

West Virginia Code §5A-3-10a states:
No contract or renewal of any contract may be awarded by the state or any of its political subdivisions to any vendor or prospective vendor when the vendor or prospective vendor or a related party to the vendor or prospective vendor is a debtor and the debt owned is an amount greater than one thousand dollars in the aggregate

DEFINITIONS:
"Debt" means any assessment, premium, penalty, fine, tax or other amount of money owed to the state or any of its political subdivisions because of a judgment, fine, permit violation, license assessment, defaulted workers' compensation premium, penalty or other assessment presently delinquent or due and required to be paid to the state or any of its political subdivisions, including any interest or additional penalties accrued thereon.

"Debtor" means any individual, corporation, partnership, association, limited liability company or any other form or business association owing a debt to the state or any of its political subdivisions. "Political subdivision" means any county commission; municipality; county board of education; any instrumentality established by a county or municipality; any separate corporation or instrumentality established by one or more counties or municipalities, as permitted by law; or any public body charged by law with the performance of a government function or whose jurisdiction is coextensive with one or more counties or municipalities. "Related party" means a party, whether an individual, corporation, partnership, association, limited liability company or any other form or business association or other entity whatsoever, related to any vendor by blood, marriage, ownership or contract through which the party has a relationship of ownership or other interest with the vendor so that the party will actually or by effect receive or control a portion of the benefit, profit or other consideration from performance of a vendor contract with the party receiving an amount that meets or exceed five percent of the total contract amount.

EXCEPTION:
The prohibition of this section does not apply where a vendor has contested any tax administered pursuant to chapter eleven of this code, workers' compensation premium, permit fee or environmental fee or assessment and the matter has not become final or where the vendor has entered into a payment plan or agreement and the vendor is not in default of any of the provisions of such plan or agreement.

LICENSING:
Vendors must be licensed and in good standing in accordance with any and all state and local laws and requirements by any state or local agency of West Virginia, including, but not limited to, the West Virginia Secretary of State's Office, the West Virginia Tax Department, West Virginia Insurance Commission, or any other state agencies or political subdivision. Furthermore, the vendor must provide all necessary releases to obtain information to enable the Director or spending unit to verify that the vendor is licensed and in good standing with the above entities.

CONFIDENTIALITY:
The vendor agrees that he or she will not disclose to anyone, directly or indirectly, any such personally identifiable information or other confidential information gained from the agency, unless the individual who is the subject of the information consents to the disclosure in writing or the disclosure is made pursuant to the agency's policies, procedures and rules. Vendors should visit **www.state.wv.us/admin/purchase/privacy** for the Notice of Agency Confidentiality Policies.

Under penalty of law for false swearing (West Virginia Code, §61-5-3), it is hereby certified that the vendor acknowledges the information in this said affidavit and are in compliance with the requirements as stated.

Vendor's Name: Bio-Rad Laboratories, Inc.

Authorized Signature: Janette J. Stockert Date: 1/22/07

Janette J. Stockert
Bid Coordination Supervisor

No Debt Affidavit (Revised 10/13/06)

Bio-Rad Laboratories

Diagnostics Group
6565 185th Avenue NE
Redmond, Washington 98052
Telephone: (425) 881-8300
Facsimile: (425) 861-5011

Date: January 22, 2007

ATTACHMENT "A"
INSTRUMENTATION AGREEMENT
RFQ Number LBS70424
Bid Tracking No. 4949

USER

State of West Virginia
Health and Human Resources
BPH – Laboratory Services
167 Eleventh Avenue
South Charleston, WV 25303
Attention: Roberta Wagner

SUPPLIER OF INSTRUMENTATION

Bio-Rad Laboratories, Inc.
6565 185th Avenue NE
Redmond, WA 98052
Attention: Janette J. Stockert

INSTRUMENTATION SUPPLIED
The following equipment is included in the above mentioned bid at no charge.

CATALOG NO.	QTY	DESCRIPTION	LIST PRICE
89700	1	Evolis™, (includes Evolis™ analyzer, PC, monitor and printer)	$92,500.00
25167	1	Evolis™, Workstation Table	$4,165.00
203301	1	Uninterrupted Power Supply 1.4KVA	$4,200.00

LOCATION OF INSTRUMENTATION
(IF OTHER THAN ABOVE ADDRESS OF USER)

Department_____

No. & Street_____

City_____

State & Zip_____

TERMS AND CONDITIONS

Bio-Rad Laboratories grants to The User possession of the instrumentation listed above and on any attached schedule, together with any replacements, duplicate parts, repairs, additions, devices, and accessories incorporated therein and/or affixed thereto, hereinafter referred to as the Instrumentation, to be used by The User at the specific location recited above subject to the following terms and conditions.

1. The Instrumentation shall at all times remain the property of Bio-Rad and The User shall have no right or property interest therein but only the right to use the Instrumentation. Bio-Rad shall have the right to display notice of ownership by affixing to the Instrumentation an identifying plate, stencil or other indication of ownership.

2. There will be no charges for the use of the Instrumentation, and this agreement does not require User to purchase supplies or services whatsoever from Bio-Rad for the use of this Instrumentation.

3. The User shall not permit or allow any attachment, lien, security interest, or other encumbrance to be filed against the Instrumentation by any individual, company, corporation, or other form of business organization with the exception of Bio-Rad or its assigns.

4. The User shall take proper care of the Instrumentation and shall not make any alterations, additions, or improvements to the Instrumentation without the prior written consent of Bio-Rad. The User shall not permit anyone other than a Bio-Rad Representative to service or repair the Instrumentation without the prior written consent of Bio-Rad.

5. Service Coverage

   5.1 At no additional cost to User, Bio-Rad will provide telephone assistance 24 hours per day, 365 days per year.

   5.2 As part of this Agreement, Bio –Rad or Bio-Rad appointed personnel will provide on-site or depot (returned to Bio-Rad) service, as needed, to keep the Equipment in good working order. On-site or depot service will be provided, at no cost to User, Monday through Friday, 8:00 a.m. to 5:00 p.m. (local time), excluding national holidays. On-site extended service coverage (Saturday, Sunday, and/or holidays) is available, but is not included in this Agreement. See Signature Service Agreement Rate Schedule currently effective for "Extended Reagent Rental Service Coverage" charges.

   5.3 Bio-Rad will not be required to pay the cost of any damage to the Equipment caused by User's negligence, abuse, or alteration of the Equipment, or by any service performed by unauthorized personnel.

   5.4 User agrees that only Bio-Rad appointed personnel are to service the Equipment.

   5.5 User agrees to utilize only Bio-Rad approved reagents, calibrators, and disposables on the Equipment.

6 Either party upon giving 60 days written notice to the other party can terminate this agreement at any time. After such termination, Bio-Rad may enter upon The User's premises and without any court order or other process of law, repossess and remove the Instrumentation with or without notice to The User.

7 Transportation charges to (and where applicable from) the place of business of the User for the Instrumentation shall be borne by Bio-Rad.

AGREED TO AND ACCEPTED BY:

The User	Bio-Rad Laboratories, Inc.
_____	Janette J. Stockert
Authorized Representative	
_____	Bid Coordination Supervisor
Title	Title
_____	January 22, 2007
Date	Date

Bio-Rad Laboratories

Diagnostics Group
6565 185th Avenue NE
Redmond, Washington 98052
Telephone: (425) 881-8300
Facsimile: (425) 861-5011

January 22, 2007

Roberta Wagner
Department of Administration
Purchasing Division
2019 Washington Street, East
Charleston, WV 25311

Re: HIPAA Request

Dear Ms. Wagner

In response to your letter asking Bio-Rad Laboratories to sign a Business Associate Agreement with respect to HIPAA, please be advised that Bio-Rad is not a Business Associate under the definition of HIPAA, as it does not receive protected health information from you. Please be advised that Abington Memorial Hospital is prohibited from producing any personal health care information to Bio-Rad. If Bio-Rad inadvertently receives such information, Bio-Rad will return this information immediately without review and will not retain any copies. I appreciate your cooperation in this matter and look forward to our continuing relationship. Please sign and return this letter to my attention.

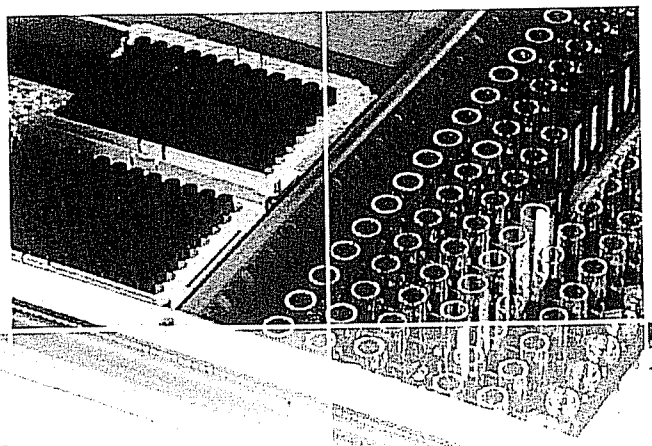
Very truly yours,

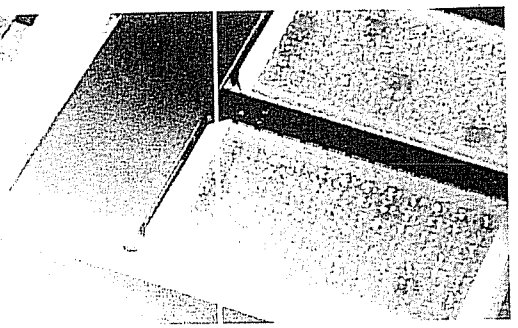
Janette J. Stockert
Bid Coordination Supervisor
Telephone: 666-8111 x1761
Fax: 425-498-1757

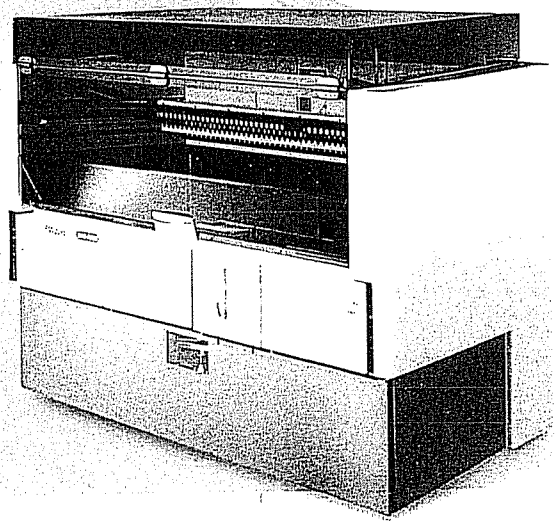
I agree to the foregoing.

Dated: _____

_____
Roberta Wagner

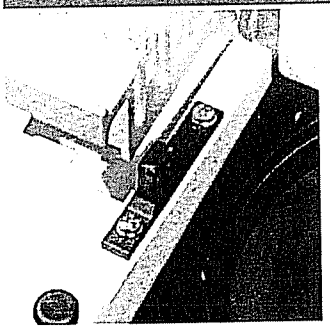






Evolis™

Automation meets throughput



Advanced laboratory automation from Bio-Rad Laboratories

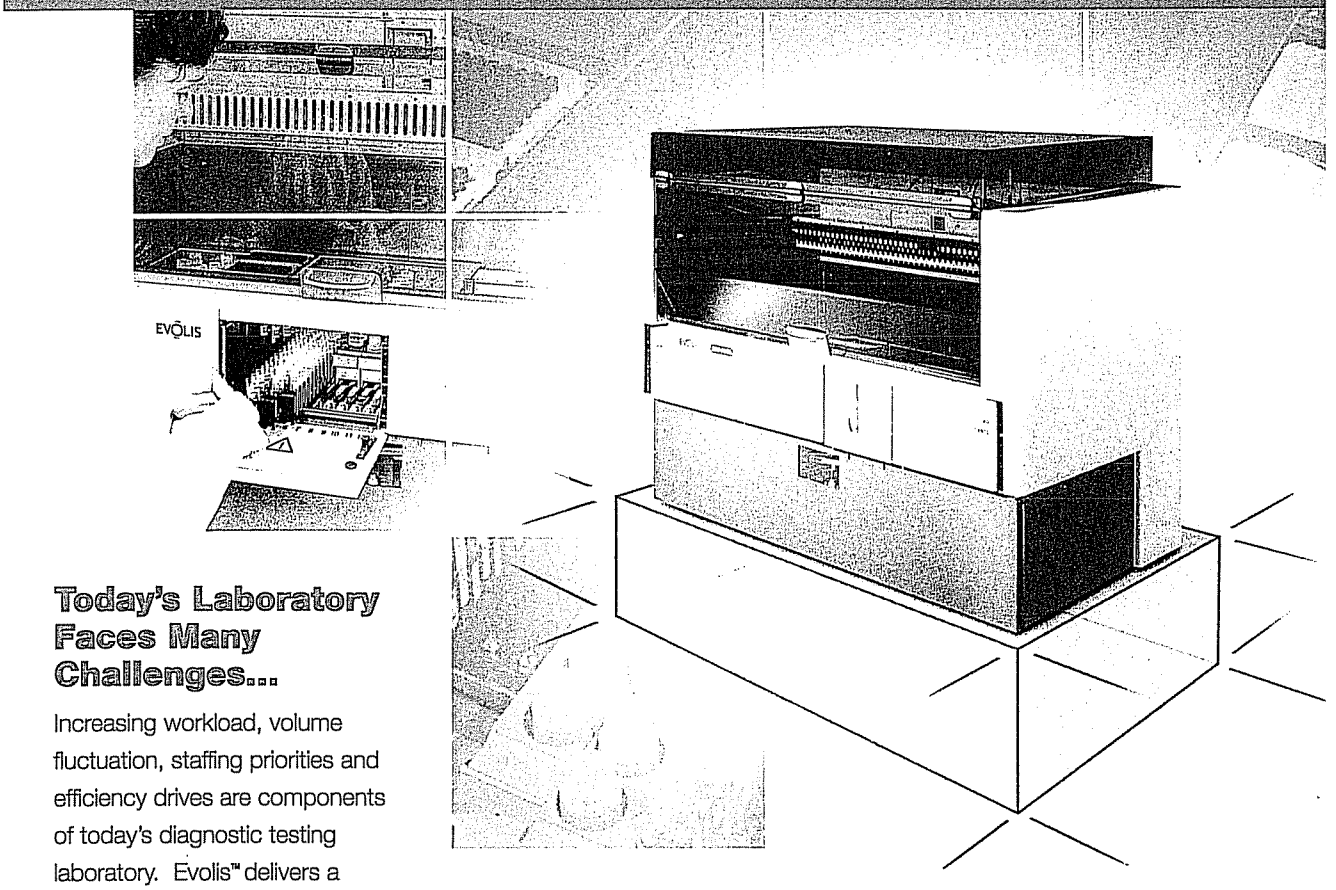
INSTRUMENTATION

BIO-RAD

Evolis™

SIDE VIEW OF LOADING DECK - Sample queue and reagent loading de

Where automation meets throughput.



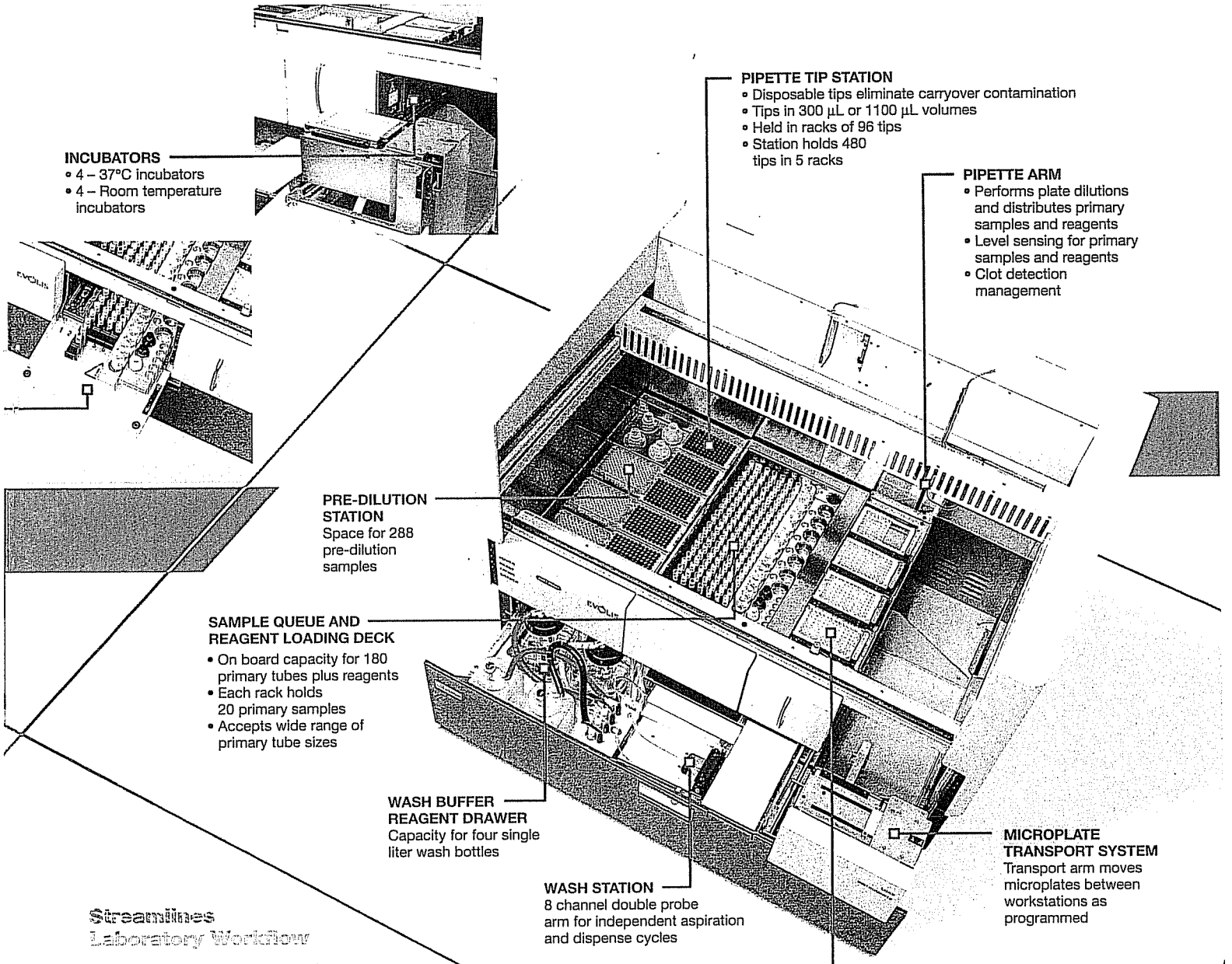
INSTRUMENTATION

Today's Laboratory Faces Many Challenges...

Increasing workload, volume fluctuation, staffing priorities and efficiency drives are components of today's diagnostic testing laboratory. Evolis™ delivers a new level of immunoassay microplate productivity to meet the challenge.

Evolis™ Microplate Automation Provides the Solution

Evolis™ is a benchtop automated pipettor, incubator, washer, reader, and data management system for EIA microplate assays. Evolis™ streamlines workflow for low-to-high volumes and optimizes productivity using a scheduler program to simplify instrument operation.



Streamlines Laboratory Workflow

- 10 minute daily start up—then load samples and reagents
- On board capacity for up to 180 samples
- Batch or multi-panel testing protocols
- Simultaneous processing for 37°C and room temperature assays
- Intuitive easy-to-use software program

Optimizes Productivity

- Software automates each assay step to reduce operator intervention
- Minimal hands-on time to add additional samples and reagents
- Sample throughput to 500 tests per shift depending on assay
- Bi-directional interface to exchange information with LIS
- Ability to link multiple units to a single LIS interface*
- CSN™ for remote technical support*

* Available soon.

Data Management

- Documents QC results
- Archives and retrieves data and sample information
- Manages repeat reactive samples

Intuitive scheduler program to prioritize instrument workflow

- Operator chooses assay order
- Scheduler prioritizes protocol steps
- Evolis™ alerts operator for additional samples or reagents

Ideal for low-to-high test volumes

- Evolis™ throughput determined by assay protocol and volume
- Batch samples to maximize throughput
- Run panels for optimal flexibility

Evolis™ Microplate Automation

Sample Processing

Primary sample capacity	180
Dilution capacity (via tubes/plate)	90/288
Container (diameter & height)	10-16 mm/50-100 mm
Sample dilution	up to 1/10,000

Reagent Processing

Calibrator & Control capacity	31
Reagent container	Bio-Rad reagent bottles
Reagent bottle volume (mL)	3, 8, 15, 30, 60, 125
Reagent identification	internal BCR
Lot & expiration management	yes

Disposable Tips

Carbon tips	300-1000 µL
Volume (useable)	10-1000 µL
Liquid level & Clot detection	capacitive
On-board capacity (walk-away)	480 tips (5 x 96)
Waste capacity	> 1000 tips
Tip detection	yes (mechanical)

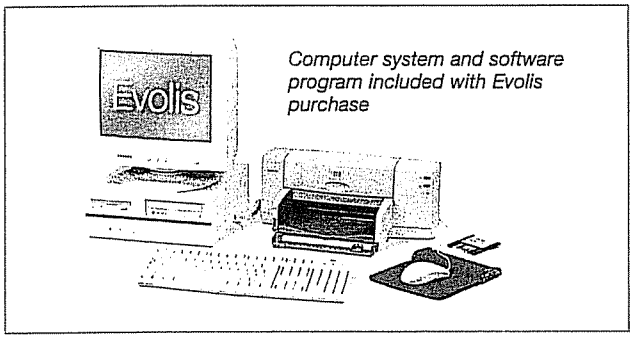
Processing Features

Patient samples (100 µl/well)	~ 16 minutes/full plate
Reagent (100 µl/well)	~ 4 minutes/full plate
Dilution (1:10)	~ 23 minutes/full plate
Precision (at 10 µl)	< 5% CV
Accuracy (at 10 µl)	< 5%
Precision (at 100 µl)	< 1.5% CV
Accuracy (at 100 µl)	< 2.5%

Internal Bar Code Scanner

Positive sample ID	yes
Reagent ID	yes
Plate ID	yes

Reads: Interleaved 2 of 5, Code 3 of 9, IATA 2 of 5, Industrial 2 of 5, UPC A & E, EAN 8 or 13, Code 128, Pharmacode, Codabar & EAN Addendum 2 or 5.



Computer system and software program included with Evolis purchase

Ordering Information

Catalog No.	Description
89601	Evolis™ Analyzer ..... 1 system
89771	Evolis™ Instrument Table ..... 1 unit

Back End Function

Work Table

Capacity	4 plates
Shaking (linear)	variable time

Incubation Tower

Loading slots	4 x ambient T°
Chambers (independent)	4 x RT + 5° to 50°C
Precision	± 1.4°C
Accuracy	± 1.0°C
Overshot	maximum 1°C
Heating-up (from RT to 37°C)	30 minutes

Washer Tool

Manifold	8 channels
Plate type (bottom shape)	flat, U- & V- shaped
Wash mode	plate and strip
Cycles/Methods (cross-wise)	1-9
Soak time (plate mode)	0-999 sec.
Wash buffers (with level sensor)	2 x 2L, 2 x 1L
Waste Capacity (with level sensor)	1 x 10L

Reader Tool

Reading head	8 channels
Read time (full plate)	~15 seconds
Light source	halogen
Reading range	0 - 3.5 O.D.
Filter wheel	8 (equipped 405, 450, 492, 550, 620, 690 nm)
Reading method	(beam) single/double
Over-range filter	yes
Linearity (0 - 2.0 O.D.)	1%
Precision (0 - 2.0 O.D.)	2.5%

Data Processing

User Interface

Software Environment	Windows® 2000 operating system

Communication

RS-232 Serial Port

On-line (bi-directional)	ASTM or ASCII format

Dimensions

Processor module

Size in inches (H x W x D)	38.0" x 45.2" x 30.4"
Weight	209 lbs.

Computer Module

Size in inches (H x W x D)	21.6" x 18.4" x 20.0"

N.B.: computer, monitor and printer are supplied

Table Module

Size in inches (H x W x D)	22.0" x 48.0" x 32.0"

Electrical

Line voltage (adjustable)	110 /260 V
Frequency	47-63 Hz
Power consumption	500 VA

Note: Specifications are subject to change without prior notice.
Evolis is a trademark of Bio-Rad Laboratories, Inc.
Windows is a trademark of Microsoft Corporation.

Bio-Rad Laboratories

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Clinical Diagnostics Group

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