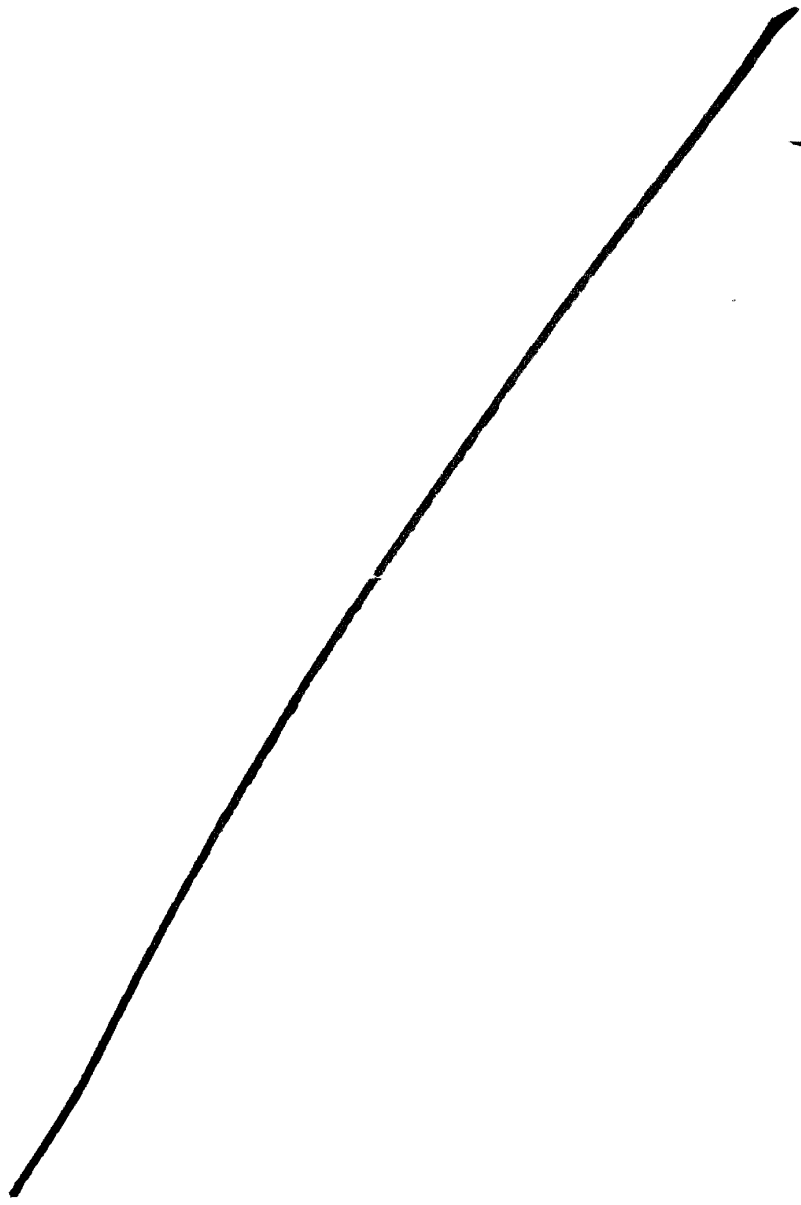


Validity Testing Information Part I

Laboratory Name: Express Analytical Laboratories
Address: 1301 18th Ave NW Suite 110 Austin, MN 55912
Responsible Person: Joseph P. Wetson (Printed Name)





RESEARCH TRIANGLE INSTITUTE

National Laboratory Certification Program

December 27, 2000

0360
Mr. Joseph P. Watson
Express Analytical Laboratory
1301 18th Avenue, NW
Suite 110
Austin, MN 55912

Dear Mr. Watson:

The enclosed critique was developed from the inspection report associated with the December 7, 2000, specimen validity testing inspection of your laboratory under the National Laboratory Certification Program (NLCP). The laboratory's procedures appeared to be in compliance with program guidance issued in Program Document 035 (September 28, 1998) and Program Document 037 (July 28, 1999).

The laboratory must ensure compliance with these program requirements. This will be reviewed at the next inspection.

The laboratory must review the enclosed critique and take all necessary corrective actions. All corrective actions must be implemented within 30 days receipt of this correspondence. **Failure to comply may result in the laboratory's suspension to perform specimen validity testing on federally regulated specimens.** All corrective actions will be reviewed at the next inspection.

Mr. Watson
Page 2 of 2
12/27/00

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
Susan Crumpton
NLCP Technical Analyst

Enclosure

cc: Project Files/svt360

NATIONAL LABORATORY CERTIFICATION PROGRAM

Document Review and Critique

Laboratory I.D. Number: 0360
Document No. Final

Laboratory: Express Analytical Labs

Location: Austin, MN

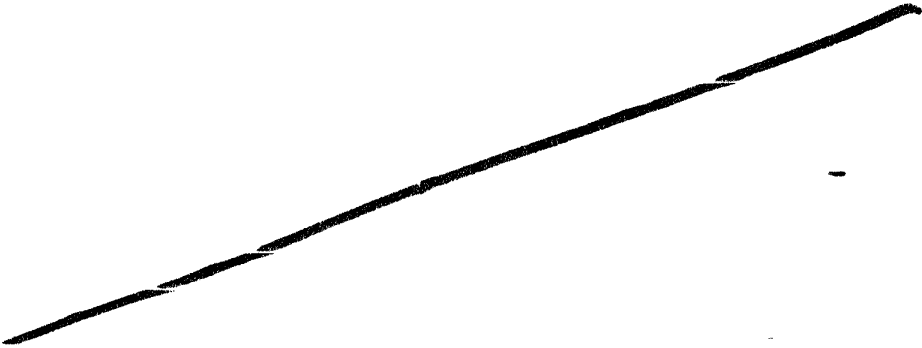
Document Reviewed: Specimen Validity Testing Inspection Report

Date: 7 December 2000


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

