



Designation: D6522 – 20

Standard Test Method for Determination of Nitrogen Oxides, Carbon Monoxide, and Oxygen Concentrations in Emissions from Natural Gas-Fired Reciprocating Engines, Combustion Turbines, Boilers, and Process Heaters Using Portable Analyzers¹

This standard is issued under the fixed designation D6522; the number immediately following the designation indicates the year of original adoption or, in the case of revision, the year of last revision. A number in parentheses indicates the year of last reappraisal. A superscript epsilon (ϵ) indicates an editorial change since the last revision or reappraisal.

1. Scope

1.1 This test method covers the determination of nitrogen oxides (NO and NO₂), carbon monoxide (CO), and oxygen (O₂) concentrations in controlled and uncontrolled emissions from natural gas-fired reciprocating engines, combustion turbines, boilers, and process heaters using portable analyzers with electrochemical sensors. Due to the inherent cross sensitivities of the electrochemical cells, this test method should not be applied to other pollutants or emission sources without a complete investigation of possible analytical interferences and a comparative evaluation with EPA test methods.

1.1.1 The procedures and specifications of this test method were originally developed during laboratory and field tests funded by the Gas Research Institute (GRI).² Comparative emission tests were conducted only on natural gas-fired combustion sources. Subsequently, the U.S. Environmental Protection Agency (EPA) sponsored Environmental Technology Verification (ETV) program conducted further evaluations of electrochemical cell analyzers, which included laboratory tests and field tests on natural gas and diesel-fueled generators. The EPA has reviewed the ETV test results, published additional information, and provided technical input that has been considered in the update of this test method.³

1.2 This test method contains notes that are explanatory and are not part of the mandatory requirements of the standard.

1.3 The values stated in SI units are to be regarded as standard. No other units of measurement are included in this standard.

¹ This test method is under the jurisdiction of ASTM Committee D22 on Air Quality and is the direct responsibility of Subcommittee D22.03 on Ambient Atmospheres and Source Emissions.

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² Juneau, P. Peeler, J. W., “Development of an Electrochemical Cell Emission Analyzer Test Method,” Gas Research Institute Topical Report prepared by Emission Monitoring Inc., GRI-96/0008, July 1997.

³ Shanklin, S., Wesson, K., and Kellar, P., “Evaluation of Portable Analyzers for Use in Quality Assuring Predictive Emission Monitoring Systems for NO_x,” prepared by The Cadmus Group, EPA Contract No. 68-W-03-033, September 2004.

1.4 This standard does not purport to address all of the safety concerns, if any, associated with its use. It is the responsibility of the user of this standard to establish appropriate safety, health, and environmental practices and determine the applicability of regulatory limitations prior to use.

1.5 This international standard was developed in accordance with internationally recognized principles on standardization established in the Decision on Principles for the Development of International Standards, Guides and Recommendations issued by the World Trade Organization Technical Barriers to Trade (TBT) Committee.

2. Referenced Documents

2.1 ASTM Standards:⁴

[D1356 Terminology Relating to Sampling and Analysis of Atmospheres](#)

[2.2 EPA Methods from 40 CFR Part 60, Appendix A:⁵ Method 3A Determination of Oxygen and Carbon Dioxide Concentrations in Emissions from Stationary Sources \(Instrumental Analyzer Procedure\)](#)

[Method 7E Determination of Nitrogen Oxides Emissions from Stationary Sources \(Instrumental Analyzer Procedure\)](#)

[Method 10 Determination of Carbon Monoxide Emissions from Stationary Source](#)

[Method 20 Determination of Nitrogen Oxides, Sulfur Dioxide, and Diluent Emissions from Stationary Gas Turbines](#)

[2.3 EPA Methods from 40 CFR Part 63, Appendix A:⁵ Method 301 Field Validation of Pollutant Measurement Methods from Various Waste Media](#)

⁴ For referenced ASTM standards, visit the ASTM website, www.astm.org, or contact ASTM Customer Service at service@astm.org. For *Annual Book of ASTM Standards* volume information, refer to the standard's Document Summary page on the ASTM website.

⁵ Available from U.S. Government Printing Office, Superintendent of Documents, 732 N. Capitol St., NW, Washington, DC 20401-0001, <http://www.access.gpo.gov>.

2.4 EPA Methods from 40 CFR Part 75, Appendix H:⁶
 Protocol G1 Procedures
 Protocol G2 Procedures

3. Terminology

3.1 Definitions:

3.1.1 For terminology relevant to this test method, see Terminology **D1356**.

3.2 Definitions of Terms Specific to This Standard:

3.2.1 *measurement system, n*—total equipment required for the determination of gas concentration. The measurement system consists of the following major subsystems:

3.2.1.1 *data recorder, n*—a strip chart recorder, computer, or digital recorder for recording measurement data.

3.2.1.2 *electrochemical cell, n*—that portion of the system that senses the gas to be measured and generates an output proportional to its concentration, or any cell that uses diffusion-limited oxidation and reduction reactions to produce an electrical potential between a sensing electrode and a counter electrode.

3.2.1.3 *external interference gas scrubber, n*—device filled with scrubbing agent used to remove interfering compounds upstream of some electrochemical cells.

3.2.1.4 *sample interface, n*—that portion of a system used for one or more of the following: sample acquisition, sample transport, sample conditioning, or protection of the electrochemical cells from particulate matter and condensed moisture.

3.2.2 *initial NO cell temperature, n*—temperature of the NO cell that is recorded during the most recent pretest calibration error check.

3.2.2.1 *Discussion*—Since the NO cell can experience significant zero drift with temperature changes in some situations, the temperature must be monitored if the analyzer does not display negative concentration results. Nitric oxide cell temperature monitoring is not required if the analyzer can display negative concentrations. Drift due to temperature changes will be identified in the post calibration check for these analyzers where negative concentrations will be observed.

3.2.3 *interference check, n*—method of quantifying analytical interferences from components in the stack gas other than the analyte.

3.2.4 *linearity check, n*—method of demonstrating the ability of a gas analyzer to respond consistently over a range of gas concentrations.

3.2.4.1 *Discussion*—Linearity checks are not required for analyzers where the electrochemical sensor manufacturer has published data demonstrating linearity through the sensor range.

3.2.5 *nominal range, n*—range of concentrations over which each cell is operated (25 % to 125 % of upscale calibration gas value).

3.2.5.1 *Discussion*—Several nominal ranges may be used

for any given cell as long as the linearity and stability check results remain within specification.

3.2.6 *response time, n*—amount of time required for the measurement system to display 95 % of a step change in gas concentration on the data recorder.

3.2.7 *upscale calibration error, n*—difference between the gas concentration exhibited by the gas analyzer and the known concentration of the upscale calibration gas.

3.2.8 *upscale calibration gas, n*—known concentration of a gas in an appropriate diluent gas.

3.2.9 *stability check, n*—method of demonstrating that an electrochemical cell operated over a given nominal range provides a stable response and is not significantly affected by prolonged exposure to the analyte.

3.2.10 *stability time, n*—elapsed time from the start of the gas injection to the start of the 15-min or 30-min stability check period, as determined during the stability check.

3.2.11 *zero calibration error, n*—gas concentration exhibited by the gas analyzer in response to zero-level calibration gas.

4. Summary of Test Method

4.1 A gas sample is continuously extracted from a duct and conveyed to a portable analyzer for determination of NO, NO₂, CO, and O₂ gas concentrations using electrochemical cells. Analyzer design specifications, performance specifications, and test procedures are provided to ensure reliable data.

4.2 Additions to or modifications of some vendor-supplied analyzers (for example, sample systems which chill the sample at the probe and transports cold, dry gas to the analyzer, heated sample line, flow meters, and so forth) may be necessary to meet the design specifications of this test method.

5. Significance and Use

5.1 The results of this test method may be used to determine nitrogen oxides and carbon monoxide emission concentrations from natural gas combustion at stationary sources.

5.2 This test method may also be used to monitor emissions during short-term emission tests or periodically in order to optimize process operation for nitrogen oxides and carbon monoxide control.

6. Interferences

6.1 NO and NO₂ can interfere with CO concentration measurements, and NO₂ can interfere with NO concentration measurements. The interference effects for the CO and NO emission measurements are quantified in 9.2 and shall not exceed 5 % of the measurement.

7. Apparatus

7.1 The minimum detectable limit depends on the nominal range of the electrochemical cell, calibration drift, and signal-to-noise ratio of the measurement system. For a well-designed system, the minimum detectable limit should be less than 2 % of the nominal range.

⁶ EPA 600/R-12/531, EPA Traceability Protocol for Assay and Certification of Gaseous Calibration Standards, May 2012. Available from <https://www.epa.gov/ord>.

7.2 Any measurement system that meets the performance and design specifications in Section 9 and 10.4.11 of this test method may be used. For systems employing a heated sampling line, the line shall maintain the gas sample at a temperature above the dew point up to the moisture removal system. The sample conditioning system shall be designed so that there are no entrained water droplets in the gas sample when it contacts the electrochemical cells. A schematic of an acceptable measurement system is shown in Fig. 1. The essential components of the measurement system are described below:

7.2.1 *Sample Probe*, glass, stainless steel, or other nonreactive material, of sufficient length to traverse the sample points, and, if necessary, heated to prevent condensation.

7.2.2 *Heated Sample Line*, heated (sufficient to prevent condensation), nonreactive tubing, to transport the sample gas to the moisture removal system.

7.2.3 *Sample Transport Lines*, nonreactive tubing to transport the sample from the moisture removal system to the sample pump, sample flow rate control, and electrochemical cells.

7.2.4 *Calibration Assembly*, a tee-fitting to attach to the probe tip for introducing calibration gases at ambient pressure during the calibration error checks. The vented end of the tee should have a flow indicator to ensure sufficient calibration gas flow. Any other method that introduces calibration gases at the probe at atmospheric pressure may be used. It is critical that all calibration gases and sample gases are delivered to the analyzer at $\pm 5\%$ the same flow rates and at $\pm 5\%$ the same absolute pressure as electrochemical cell measurements are very sensitive to pressure differences. These parameters are typically verified using a digital pressure gauge with an accuracy of $\pm 2\%$.

7.2.5 *Digital Pressure Gauge*, of appropriate pressure range and an accuracy of $\pm 2\%$ or better.

7.2.6 *Moisture Removal System*, a chilled condenser or similar device (for example, permeation dryer), to remove

condensate continuously from the sample gas while maintaining minimal contact between the condensate and the sample gas.

7.2.7 *Particulate Filters*—Filters at the probe or the inlet or outlet of the moisture removal system and inlet of the analyzer may be used to prevent accumulation of particulate material in the measurement system and extend the useful life of the components. All filters shall be fabricated of materials that are nonreactive to the gas being sampled.

7.2.8 *Sample Pump*, a leak-free pump, to pull the sample gas through the system at a flow rate sufficient to minimize the response time of the measurement system. The pump must be constructed of any material that is nonreactive to the gas being sampled.

7.2.9 *Sample Flow Rate Control*, a sample flow rate control valve and rotameter, or equivalent, to maintain a constant sampling rate within 10% during sampling and calibration error checks. The components shall be fabricated of materials that are nonreactive to the gas being sampled.

7.2.10 *Gas Analyzer*, a device containing electrochemical cells to determine the NO, NO₂, CO, and O₂ concentrations in the sample gas stream and, if necessary, to correct for interference effects. The analyzer shall meet the applicable performance specifications of Section 9.

7.2.10.1 A means of controlling the analyzer flow rate and a device for determining proper sample flow rate shall be provided at the analyzer. For example, a needle valve and precision rotameter, or pressure gauge downstream of all flow controls, or equivalent can be used.

7.2.10.2 The electrochemical cell analyzer should have a minimum upscale calibration level appropriate to the stack gas concentration being measured. For example, if the stack gas NO_x concentration is less than 10 ppm, the analyzer should have the capability to analyze a 10-ppm (or less) upscale calibration gas for the NO and NO₂ cells.

NOTE 1—Housing the analyzer in a clean, thermally-stable, vibration-free environment will minimize drift in the analyzer calibration.

NOTE 2—If the NO_x analyzer resolution is 0.1 ppm, it will be more likely to pass the performance specifications when testing at sources with low stack gas concentrations.

NOTE 3—It is recommended that analyzer manufacturer's maintenance procedures be followed.

7.2.11 *Data Recorder*, a strip chart recorder, computer, or digital recorder, for recording measurement data. The data recorder resolution (that is, readability) shall be at least 1 ppm for CO, NO, and NO₂; 0.1% O₂ for O₂; and 1°C for temperature. Alternatively, a digital or analog meter having the same resolution may be used to obtain the analyzer responses and the readings may be recorded manually.

NOTE 4—Some analyzers incorporate a digital data logger. Such a recorder may be used provided it meets the resolution requirements of 7.2.11.

7.2.12 *External Interference Gas Scrubber*, a device used by some analyzers to remove interfering compounds upstream of a CO electrochemical cell. The measurement system should provide the operator with a means of determining when the scrubbing agent is exhausted (that is, visible color change indication, or electronic ppm hour counter, or equivalent).

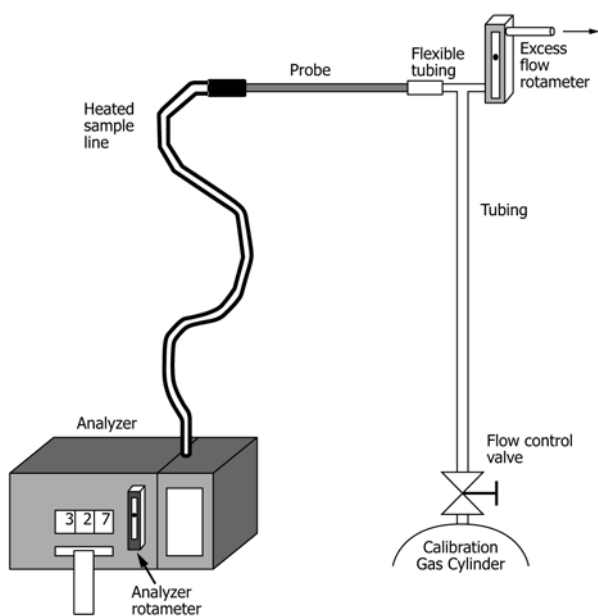


FIG. 1 Calibration System Schematic

7.2.13 *NO Cell Temperature Indicator*, a thermocouple, thermistor, or other device must be used to monitor the temperature of the NO electrochemical cell. The temperature may be monitored at the surface or within the cell. This is not required if the analyzer is capable of displaying negative concentrations.

8. Reagents and Materials

8.1 The analytical range for each gas component is determined by the electrochemical cell design. A portion of the analytical range is selected by choosing an upscale calibration gas concentration approximating the flue gas concentrations.

8.2 *Calibration Gases*—The calibration gases for the gas analyzer shall be CO in nitrogen or CO in air, NO in nitrogen, NO₂ in air, and O₂ in nitrogen.

8.2.1 For the mid-level and upscale calibration gases, use calibration gases certified according to EPA Protocol G1 Procedures or Protocol G2 Procedures.

8.2.2 Alternative certification techniques may be used, if approved in writing by the applicable regulatory agency.

8.3 *Upscale Calibration Gases*—Use these gases for calibration error, linearity, and interference checks of each nominal range of each cell. Select concentrations as follows:

8.3.1 *CO and NO Upscale Calibration Gases*—Choose an upscale calibration gas concentration such that the average stack gas reading for each test run is greater than 25 % of the upscale calibration gas concentration. Alternatively, choose the upscale calibration gas such that it is not greater than twice the concentration equivalent to the emission standard. If concentration results exceed 125 % of the upscale calibration gas at any time during the sampling run, then the test run for that channel is not valid.

8.3.2 *NO₂ Upscale Calibration Gas*—Choose an upscale calibration gas concentration such that the average stack gas reading for each test run is greater than 25 % of the upscale calibration gas concentration. Alternatively, choose the upscale calibration gas concentration such that it is not greater than the ppm concentration value of the NO upscale calibration gas. The tester should be aware that NO₂ cells are generally designed to measure much lower concentrations than NO cells and the upscale calibration gas should be chosen accordingly. If concentration results exceed 125 % of the upscale gas at any time during the sampling run then the test run for that channel is not valid.

8.3.3 *O₂ Upscale Calibration Gas*—Choose an upscale calibration gas concentration such that the difference between the upscale calibration gas concentration and the average stack gas reading for each run is less than 10 % O₂. Where the stack oxygen is high, dry ambient air having a dew point less than 20°C may be used and assumed to have a concentration of 20.9 % O₂.

8.4 *Mid-Level Gases*—Select mid-level gas concentrations that are 40 to 60 % of the upscale calibration gas concentrations.

8.5 *Zero Gas*—Zero gas must have concentrations of less than 0.25 % of the upscale calibration gas for each component.

Ambient air may be used in a well-ventilated area provided the upscale calibration level is >25 ppm.

9. Preparation of Apparatus

9.1 *Linearity Check*—The procedures in this subsection are not required if the manufacturer of the sensors used in the particular analyzer has published information clearly demonstrating the linearity of the sensor throughout the sensor range, and explicitly states the minimum and maximum measurement ranges for which the sensor can be shown to exhibit a linear response meeting or exceeding the requirements of this test method. It is the responsibility of the person performing this test method to acquire such information from the sensor manufacturer or portable analyzer manufacturer and have this prior to performing the test. If this information is not available from the manufacturer at the time of the test, the following procedures in this section shall be conducted.

9.1.1 Conduct the linearity check once for each nominal range that is to be used on each electrochemical cell (NO, NO₂, CO, and O₂) before each field test program.

9.1.1.1 Repeat the linearity check immediately after 5 days of analyzer operation, if a field test program lasts longer than 5 days.

9.1.1.2 Repeat the linearity check whenever a cell is replaced.

9.1.2 If the analyzer uses an external interference gas scrubber with a color indicator or other depletion indicator, verify that the scrubbing agent is not depleted, following the analyzer manufacturer's recommended procedure.

9.1.3 Calibrate the analyzer with zero and upscale calibration gases.

9.1.4 Inject the zero, mid-level, and upscale calibration gases that are appropriate for each nominal range to be used on each cell. Gases need not be injected through the entire sample handling system.

9.1.5 Purge the analyzer, briefly with ambient air between gas injections.

9.1.6 For each gas injection, verify that the flow rate is constant and that the analyzer responses have stabilized.

9.1.7 Record all of the responses (stabilized) on a form similar to Fig. 2.

9.1.8 For the zero, mid-level, and upscale calibration gases, calculate the absolute value of the difference between the gas value and the analyzer response.

9.1.9 *Linearity Specifications:*

9.1.9.1 *NO, CO, and O₂ Cells*—≤2.5 % of the upscale calibration gas concentration or <1 ppm difference from the upscale calibration gas concentration, whichever is less restrictive.

9.1.9.2 *NO₂ Cells*—≤3.0 % of the upscale calibration gas concentration or <1 ppm difference from the upscale calibration gas concentration, whichever is less restrictive.

9.2 *Interference Check:*

9.2.1 Determine interference responses for the CO and NO cells, using the results from the upscale calibration gas injections (see 11.2 and 11.3 for calculations).

9.2.2 *Interference Specifications*—CO and NO interference responses: ≤5 %.

Date _____ Analyst _____

Analyzer Manufacturer/Model No. _____

Analyzer Serial Number _____

Calibration Gas Conc.	ppm NO Response	ppm NO ₂ Response	ppm CO Response	% Oxygen Response

iTeh Standards
<https://standards.itih.ai>
 Document Preview

<https://standards.itih.ai/catalog/standards/sist/4e9a71ba-5079-4b77-bfd0-dee1a7805e99/astm-d6522-20>

FIG. 2 Linearity Check Data Sheet

9.3 Stability Check:

9.3.1 Conduct the stability check once for each nominal range that is to be used on each pollutant electrochemical cell (NO, NO₂, and CO) before each field test program. The analyzer should be purged with ambient air between gas injections. Otherwise, the cells will be exposed to high NO and

NO₂ concentrations for an extended time which can cause a cell's performance to degrade (this is the so-called "O₂-starved exposure").

9.3.1.1 Repeat the stability check immediately after 5 days of analyzer operation, if a field test program lasts longer than 5 days.

9.3.1.2 Repeat the stability check if a cell is replaced or if a cell is exposed to gas concentrations greater than 125 % of the upscale calibration gas concentration.

9.3.2 Inject the upscale calibration gas into the analyzer and record the analyzer response at least once per minute until the conclusion of the test. One-minute average values may be used instead of instantaneous readings.

9.3.3 After the analyzer response has stabilized, continue to flow the upscale calibration gas for at least 30 minutes.

9.3.3.1 Alternatively, if the concentration reaches a maximum value within 5 minutes, the data may be recorded for at least 15 minutes (rather than 30 minutes) following the initial maximum value.

9.3.3.2 The more stringent specification in 9.3.8 shall apply when the 15-minute test period is used.

9.3.4 Make no adjustments to the analyzer during the test except to maintain constant flow.

9.3.5 Record the stabilization time as the number of minutes elapsed between the start of the gas injection and the start of the 30-minute (or 15-minute) stability check period.

9.3.6 Determine the highest and lowest concentrations recorded during the 30-minute (or 15-minute) period and record the results on a form similar to Fig. 3.

9.3.7 Calculate the absolute value of the difference between the maximum and minimum values recorded during the 30-minute period (or the 15-minute period) for the CO, NO, and NO₂ upscale calibration gases.

9.3.8 *Stability Specifications:*

9.3.8.1 *Thirty-Minute Period*— ≤ 2.0 % of the upscale calibration gas concentration or < 1 ppm difference, whichever is less restrictive.

9.3.8.2 *Fifteen-Minute Period*— ≤ 1.0 % of the upscale calibration gas concentration or < 1 ppm difference, whichever is less restrictive.

10. Procedure

10.1 *Reciprocating Engines—Selection of Sampling Site and Sampling Points:*

10.1.1 Select a sampling site located at least five stack, duct, or pipe diameters downstream of any turbocharger exhaust, crossover junction, or recirculation take-offs and upstream of any dilution air inlet.

10.1.2 Locate the sampling site no closer than 1 m or three stack, duct, or pipe diameters (whichever is less) upstream of the gas discharge to the atmosphere.

10.1.3 Use a minimum of three sampling points located at positions of 16.7, 50, and 83.3 % of the stack, duct, or pipe diameter.

10.1.4 The tester may choose an alternative sampling location or sample from a single point in the center of the stack, duct, or pipe, if previous test data demonstrate that the stack, duct, or pipe gas concentration does not vary significantly across the duct diameter, or both.

10.2 *Combustion Turbines—Selection of Sampling Site and Sampling Points:*

10.2.1 Select a sampling site and sample points according to the procedures in EPA Method 20.

10.2.2 The tester may choose an alternative sampling location or sample from a single point in the center of the stack, duct, or pipe if previous test data demonstrate that the stack, duct, or pipe gas concentrations of CO, NO_x, and O₂ do not vary significantly across the duct diameter, or both.

10.3 *Warm Up Period:*

10.3.1 Assemble the sampling system and allow the analyzer and sample interface to warm up and adjust to ambient temperature at the location where the stack measurements will take place.

10.3.2 The warm-up period ensures that excessive calibration drift does not occur due to temperature changes. If the pretest and post test calibration error check results are within the specifications of the method and the NO cell temperature meets the requirements of 10.5.3 (for analyzers that cannot display negative values), then the duration of the warm-up period is sufficient.

10.4 *Pretest Calibration Error Check:*

10.4.1 Conduct the calibration error check at the sampling location (near the sampling port) just prior to the start of an emissions test or test run. Keep the analyzer in the same location until the post test calibration error check is conducted.

10.4.2 For analyzers that have an external interference gas scrubber tube, inspect the condition of the scrubbing agent and ensure that it will not be exhausted during sampling.

10.4.3 Inject the zero and upscale calibration gases at the probe tip using the calibration assembly.

10.4.4 Ensure that the calibration gases flow through all parts of the sample interface (including any exhaust lines).

10.4.5 During this check, make no adjustments to the system except those necessary to achieve the correct calibration gas flow rate at the analyzer.

10.4.6 Set the analyzer flow rate to the value recommended by the analyzer manufacturer.

10.4.7 Allow each reading to stabilize (no less than the stability time noted during the stability check) before recording the final response on a form similar to Fig. 4.

10.4.8 After achieving a stable response, disconnect the gas and briefly purge with ambient air.

10.4.9 Determine the NO and CO response times by observing the time required to respond to 95 % of a step change in the analyzer response for both the zero and upscale calibration gases. Note the longer of the two times as the response time. For NO₂ upscale calibration gas record the time required to respond to 90 % of a step change.

10.4.10 Calibrate all electrochemical cells in the analyzer if the analyzer uses an internal calculation method to compensate for interferences.

10.4.11 If the zero and upscale calibration error test results are not within the specifications stated below, take corrective action and repeat the calibration error check until acceptable performance is achieved.

10.4.11.1 *Zero Calibration Error Specifications*— ≤ 3 % of the upscale calibration gas value or < 0.5 ppm difference from the upscale calibration gas value, whichever is less restrictive, for NO, NO₂, and CO channels; ≤ 0.3 % O₂ for the O₂ channel.

10.4.11.2 *Upscale Calibration Error Specifications*— ≤ 5 % of the upscale calibration gas value or < 1 ppm difference from

Date _____

Analyst _____

Analyzer Manufacturer/Model No. _____

Analyzer Serial Number _____

Pollutant Channel _____ Upscale Calibration Gas Concentration _____

Elapsed Time (Minutes)	Analyzer Response	Elapsed Time (Cont.)	Analyzer Response	Elapsed Time (Cont.)	Analyzer Response
1		17		33	
2		18		34	
3		19		35	
4		20		36	
5		21		37	
6		22		38	
7		23		39	
8		24		40	
9		25		41	
10		26		42	
11		27		43	
12		28		44	
13		29		45	
14		30		46	
15		31		47	
16		32		48	

For 30-minute stability period: _____ maximum _____ minimum

For 15-minute stability period: _____ maximum _____ minimum

Maximum Deviation = $100 \times (\text{max.} - \text{min.}) / \text{upscale calibration gas conc.} =$ _____

Stability Time _____

FIG. 3 Stability Check Data Sheet

the upscale calibration gas value, whichever is less restrictive, for NO, NO₂, and CO channels; ≤0.5 % O₂ for the O₂ channel.

10.5 *NO Cell Temperature Monitoring*—Nitric oxide (NO) cell temperature monitoring is required if the analyzer does not display negative concentrations.

10.5.1 Record the initial NO cell temperature during the pretest calibration error check.

10.5.2 Monitor and record the temperature regularly (at least once each 5 min) during the sample collection period.

10.5.3 If at any time during sampling the NO cell temperature is ≥30°C and has increased or decreased by more than 3°C since the pretest calibration, do the following:

10.5.3.1 Stop sampling immediately.

10.5.3.2 Conduct a post test calibration error check according to 10.7.

10.5.3.3 Re-zero the analyzer.

10.5.3.4 Then conduct another pretest calibration error check before continuing.