APPENDIX A

STANDARD OPERATING PROCEDURES FOR NATIONAL PARK SERVICE FILTER PREPARATION, EXTRACTION, AND ANION ANALYSIS
Standard Operating Procedure for
National Park Service Filter Preparation, Extraction, and Anion Analysis

Environmental Chemistry Department
Research Triangle Institute
Research Triangle Park, North Carolina

Prepared by: _______________ Date: ________

Reviewed by: _______________ Date: ________

Approved by: _______________ Date: ________
1.0 PROCEDURAL SECTION

1.1 Purpose and Applicability

This document outlines procedures for the filter preparation and extraction, and the subsequent determination of anions in filter extracts. Analytical procedures outlined are specific to the Dionex Model DX-500 Ion Chromatographs located in Research Triangle Institute’s Center for Environmental Measurements and Quality Assurance.

1.2 Summary of Method

Nylon filters for collection of anions do not require pretreatment. Quartz fiber filters are impregnated with K$_2$CO$_3$ glycerol solution for collection of gaseous SO$_2$. Exposed filter samples are extracted by a method appropriate for the analyte(s) of interest. Nylon filters are extracted with deionized water. Extraction with deionized water makes it possible to analyze for both anions and cations. Carbonate-impregnated quartz fiber filters are extracted with a 0.1% peroxide solution that oxidizes the SO$_2$ to sulfate ion.

Sample extracts are passed through a column of ion chromatographic resin consisting of polymer beads coated with quaternary ammonium active sites. During passage through the column, anion separation occurs due to the different affinities of the anions for the active resin sites. Following separation, the anions pass through a suppressor column which exchanges all cations for H$^+$ ions. An eluent which yields a low conducting acid is used. Species are detected and quantified as their acids (e.g., HCl) by a conductivity meter.

1.3 Health and Safety Warnings

The NPS ion analysis operations do not involve unusual risks from electrical equipment or chemical exposures. Standard RTI laboratory health and safety precautions will be followed.

1.4 Cautions

Laboratory personnel should always wear clean clothes and wash hands thoroughly before performing filter handling and analysis procedures. The use of cleanroom gloves is required and will minimize the potential for laboratory contamination.

1.5 Interferences

Large amounts of anions eluting close to the ions of interest will result in an interference. No interferences have been observed in nylon filters samples analyzed to date. If interferences are observed, several steps to increase separation can be taken, such as reducing eluent strength and/or flow rate or replacing the columns.
Analysis of the extracts of \( \text{K}_2\text{CO}_3 \)/glycerol coated filters, which are used for \( \text{SO}_2 \) collection, requires careful quantitation of the sulfate peak. The sulfate peaks of undiluted extracts are broad and asymmetrical compared to the sulfate peaks of the standards, resulting in low concentrations when peak height is used for quantitation. A 5-fold dilution of the extract reduces the \( \text{K}_2\text{CO}_3 \)/glycerol matrix interference sufficiently to produce a sulfate peak that is symmetrical and quantifiable. This method is used for all extracts of \( \text{K}_2\text{CO}_3 \)/glycerol coated filters, with careful monitoring of recoveries through the use of spiked extracts.

1.6 Personnel Qualifications

Personnel employed to perform ion analysis operations will have at least a Bachelor's Degree in a laboratory science, and will be trained by a supervisor before being allowed to process client samples for the NPS program.

1.7 Apparatus and Materials

1.7.1 Carbonate-Impregnated Filter Preparation

Quartz fiber filters (Pallflex NO. 3700 AT-UP, 37 mm, or equivalent

- Volumetric flask, 1000 mL
- Crystallizing dish (flat bottom and straight sides), 150 mm x 75 mm
- Tweezers
- Glass rod drying racks
- Vacuum drying oven

1.7.2 Filter Extraction and Analysis

- Disposable centrifuge tubes with screw caps, 50 mL, polypropylene
- Tweezers
- Calibrated Rainin electronic pipette
- Adjustable Eppendorf pipette
- Ultrasonic bath fitted with epoxy-coated test tube rack to hold centrifuge tubes
NPS Anion Analysis
Revision 3
Date: October 26, 2005
Page 5 of 16

ANALYSIS OF FILTERS FROM THE IMPROVE AIR QUALITY NETWORK

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- Ion chromatograph (Dionex Model DX-500 with LC20 chromatography module, one IP25 isocratic pump, one GP50 gradient pump, two CD20 conductivity detectors, two Dionex AS40 automated samplers and PeakNet Control Windows 2000 Workstation) with Dionex AG12A anion guard column, Dionex AS12A anion separator column, and Dionex ASRS-ULTRA anion self-regenerating suppressor column or AMMS-III chemical suppressor column for each system.
- Pressurized eluent reservoirs.
- Volumetric flasks - an assortment of sizes
- Dionex autosampler vials with filter caps

1.8 Reagents

Use ACS reagent grade chemicals for the preparation of all solutions. Dry chemicals used for the preparation of calibration standards at 105°C for 2 hours and cool in a desiccator immediately before weighing.

1.8.1 Carbonate-Impregnated Filter Preparation

1.8.1.1 K₂CO₃/2% glycerol solution: Mix 200 grams reagent grade potassium carbonate, 100 mL glycerol, and dilute to 1000 mL in a volumetric flask.

1.8.2 Carbonate-Impregnated Filter Extraction

1.8.2.1 0.1% H₂O₂: Dilute 6.68 mL 30% reagent grade H₂O₂ to 2000 mL using deionized water. (Prepare fresh each extraction day.)

1.8.3 Anion Analysis

1.8.3.1 Eluent, 0.3mM NaHCO₃/2.7mM Na₂CO₃: Dissolve 0.5041 g NaHCO₃ and 5.7235 g Na₂CO₃ in 20 liters deionized water (DI H₂O).

1.8.3.2 Mixed Stock Solution, 1000 mg/L NO₂⁻, NO₃⁻, and SO₄²⁻, and 200 mg/L Cl⁻: Dissolve 1.4998 g NaNO₂, 1.6305 g KNO₃, 1.8142 g K₂SO₄, and 0.3297 g NaCl in 1 liter DI H₂O.

1.8.3.3 Standard Solution A: Dilute 10 mL mixed stock solution to 100 mL with DI H₂O (100 mg/L NO₂⁻, NO₃⁻, SO₄²⁻, and 20 mg/L Cl⁻).

1.8.3.4 Standard Solution B: Dilute 10 mL Standard Solution A to 100 mL with DI H₂O (10 mg/L NO₂⁻, NO₃⁻, and SO₄²⁻, and 2 mg/L Cl⁻).
1.8.3.5 Calibration Standards: Using Standard Solutions A and B, prepare calibration standards with deionized water in 100 mL volumetric flasks as shown in Table 1. Prepare fresh calibration standards weekly and refrigerate when not in use.

1.8.3.6 Quality Assurance Stock Solutions: Purchase from CPI International, GFS (Columbus, OH), and/or EM Science (Gibbstown, NJ).

TABLE 1. PREPARATION OF ANION CALIBRATION STANDARDS

<table>
<thead>
<tr>
<th>Standard</th>
<th>Cl (mg/L)</th>
<th>NO$_2^-$, NO$_3^-$, SO$_4^{2-}$ (mg/L)</th>
<th>mL of Standard Solution/100 mL</th>
</tr>
</thead>
<tbody>
<tr>
<td>STANDARD SOLUTION A</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>5.0</td>
<td>25.0</td>
<td>25.0</td>
</tr>
<tr>
<td>2</td>
<td>2.0</td>
<td>10.0</td>
<td>10.0</td>
</tr>
<tr>
<td>3</td>
<td>0.6</td>
<td>3.0</td>
<td>3.0</td>
</tr>
<tr>
<td>STANDARD SOLUTION B</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>0.2</td>
<td>1.0</td>
<td>10.0</td>
</tr>
<tr>
<td>6</td>
<td>0.1</td>
<td>0.5</td>
<td>5.0</td>
</tr>
<tr>
<td>7</td>
<td>0.04</td>
<td>0.2</td>
<td>2.0</td>
</tr>
<tr>
<td>1 ppm STANDARD</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>8</td>
<td>0.02</td>
<td>0.1</td>
<td>10.0</td>
</tr>
<tr>
<td>9</td>
<td>0.01</td>
<td>0.05</td>
<td>5.0</td>
</tr>
</tbody>
</table>

NOTE: Higher concentration standards can be prepared from Standard A or from the mixed stock solution if needed.

1.9 Carbonate-Impregnated Filter Preparation

Use the procedures described below to prepare filters for collection of sulfur dioxide. The chemist will wear cleanroom gloves and handle the filter with forceps or tweezers.

(1) Purchase quartz fiber filters (Pallflex No. 3700 AT-UP, 37 mM, or equivalent for 24-hour sampling).

(2) Inspect all filters for imperfections. Analyze at least one blank filter from every 100 for sulfate to verify the manufacturer's pre-established specifications. Return rejected filters to the manufacturer for credit.
(3) Pour approximately 250 mL of filter coating solution (Section 1.8.1.1) into a 150 mm x 75 mm crystallizing dish (with a flat bottom and straight sides).

(4) Saturate a batch of 50, 37-mm quartz filters in the filter-coating solution. Using forceps or tweezers, remove the filters one at a time from the coating solution and drain for a few seconds. Place the filters on a clean glass rod drying rack and dry the filters in an SO$_2$-free oven under vacuum at 60°C for 30 minutes, or until dry. Place the filters in a clean, resealable polyethylene bag, then store the bag in a refrigerator.

(5) Extract and analyze 1% of the newly-made filters for SO$_4^{2-}$ background. Reject the batch if the standard deviation of the mean sulfate concentration is more than half the lower quantifiable limit of sulfate.

1.10 Sample Collection

Sample collection is not applicable to this SOP because samples are acquired by the PMCC.

1.11 Sample Handling

RTI will provide chain-of-custody documentation with all sample shipments to track and ensure that filter samples are collected, transferred, stored, and analyzed by authorized personnel; sample integrity is maintained during all phases of sample handling and analysis; and an accurate written record is maintained of sample handling and treatment from the time of its collection, through the laboratory analytical process, to the eventual relinquishing of all data to the NPS.

1.12 Filter Extraction Procedure

1.12.1 Nylon Filters

NOTE: Nylon filters to be analyzed for anions are extracted with water. Extraction with deionized water makes it possible to analyze for both anions and cations. The anion eluent produces a large sodium peak in the cation chromatogram that precludes quantitation of the sodium ion in the filter extract and interferes with the quantitation of ammonium ion.

To extract the filters, the analyst will do the following:

(1) Remove filters to be extracted from the freezer and allow them to equilibrate to room temperature.

(2) Using gloved hands and tweezers, place each filter in a centrifuge tube that has been labeled with the sample I.D. (The label is carefully taped near the top of the centrifuge tube to prevent loss during sonication.)

(3) Add exactly 20 mL of deionized water using a calibrated Rainin electronic pipette.

(4) Screw the cap tightly on the centrifuge tube.
(5) Place the batch of centrifuge tubes in an epoxy-coated wire test tube rack, expose them to ultrasonic energy in a bath for 30 minutes, and then allow them to sit at room temperature overnight. Refrigerate at least one additional night prior to analysis.

(6) Record the date of extraction on the RTI Sample Filter Processing Form.

Allow the samples to warm to room temperature just prior to analysis.

1.13 IC Procedure

(1) Fill the eluent reservoirs with the eluent.

(2) Start the eluent flow, activate the self-regenerating suppressor, and allow the baseline to stabilize.

(3) Inject two DI H₂O blanks to flush the system and to ensure that the system is operating properly.

(4) Using the calibration schedule, perform the daily multipoint calibration over the range 0.05 to 25.0 ppm NO₂⁻, NO₃⁻ and SO₄²⁻ (0.01 to 5.0 ppm Cl⁻) followed by quality assurance/quality control (QA/QC) samples listed below:

- QC sample containing concentrations of Cl⁻, NO₂⁻, NO₃⁻ and SO₄²⁻ typical of those found in the mid-range of actual filter extract concentrations.
- QC sample containing concentrations of Cl⁻, NO₂⁻, NO₃⁻ and SO₄²⁻ typical of those found at the lower end of actual filter extract concentrations.
- Commercially prepared, NIST-traceable QA sample containing known concentrations of Cl⁻, NO₂⁻, NO₃⁻ and SO₄²⁻.

If the observed value for any ion differs by more than 10 percent from the known values, identify and correct the problem before analyzing samples.

(5) Load the sample extracts into the autosampler vials according to the schedule prepared for that day. Typically, fifty field samples are analyzed per day. The daily schedule includes, at a minimum, 3 duplicate samples, 2 spiked samples and 5 QA/QC samples.

(6) Begin the analysis run, occasionally checking to ensure that the system is operating properly.

(7) Examine the data at the end of the run. If the concentration of any ion exceeds the upper end of the calibration curve, dilute the sample appropriately and include with the samples to be analyzed the following day.
1.14 Calculations and Data Reduction

Peak areas are entered into the computer automatically by the PeakNet software where calculations are performed using a quadratic fit to the calibration data. The quadratic fit yields the following:

\[ y_i = ax_i^2 + bx_i + c \]

where:
- \( y \) = the calculated anion concentration, mg/L
- \( x \) = the instrument response

Initially, the calibration curves from 0.05 to 10.0 ppm NO$_2^-$, NO$_3^-$ and SO$_4^{2-}$ (0.01 to 2.0 ppm Cl$^-$) are used for the calculation of the extract anion concentrations. All nitrite, nitrate, and sulfate concentrations that exceed 10 ppm and all chloride concentrations that exceed 2 ppm are recalculated with the 25.0 ppm NO$_2^-$, NO$_3^-$ and SO$_4^{2-}$ (5 ppm Cl$^-$) standard added to the calibration curve. If a recalculated nitrite, nitrate or sulfate concentration exceeds 25 ppm or a recalculated chloride concentration exceeds 5 ppm, the extract is diluted appropriately (usually 5-fold) to bring the ion concentration into the calibration range and reanalyzed.

2.0 QUALITY ASSURANCE AND QUALITY CONTROL

If correlation coefficient does not exceed 0.998, stop the analysis and identify the problem.

Analyze QC samples (see Section 1.14) at the beginning of every analytical run. Compare the results with those obtained during previous QC tests. If the observed concentration of any ion differs from the known value by greater than 10%, stop the analysis until the problem is identified and corrected. Analyze a duplicate sample, a QA/QC sample, and a spiked sample after at least every 20 field samples.

3.0 REFERENCE

DRI Document No. 8068.1F4, Appendix D, Section 4.2
APPENDIX B

STANDARD OPERATING PROCEDURE
FOR NATIONAL PARK SERVICE
CATION ANALYSIS
Standard Operating Procedure for
National Park Service Cation Analysis

Environmental Chemistry Department
Research Triangle Institute
Research Triangle Park, North Carolina

Prepared by: _____________________ Date: __________

Reviewed by: _____________________ Date: __________

Approved by: _____________________ Date: __________
1.0 Procedural Section

1.1 Purpose and Applicability

This document outlines procedures for the extraction and subsequent determination of ammonium, sodium, potassium, calcium, and magnesium ions in nylon filter extracts. Analytical procedures outlined are specific to the Dionex Model DX-500 Ion Chromatograph located in Research Triangle Institute’s Center for Environmental Measurements and Quality Assurance.

1.2 Summary of Method

Nylon filters for collection of ions do not require pretreatment. Exposed filter samples are extracted by a method appropriate for the analyte(s) of interest. Nylon filters to be analyzed for cations as well as anions will be extracted with deionized water.

Cations in solution are separated when passed through an ion chromatographic column containing surface-sulfonated ion exchange resin; separation is caused by differing affinities of the cations for the active sites on the resin. After separation, the cations pass through a suppressor column which exchanges all anions for OH⁻ ions. Species are detected and quantified as their hydroxides by a conductivity meter. The eluent is sulfuric acid which yields deionized water when passed through the suppressor column.

1.3 Health and Safety Warnings

The NPS ion analysis operations do not involve unusual risks from electrical equipment or chemical exposures. Standard RTI laboratory health and safety precautions will be followed.

1.4 Cautions

Laboratory personnel should always wear clean clothes and wash hands thoroughly before performing filter handling and analysis procedures. The use of cleanroom gloves is required and will minimize the potential for laboratory contamination.

1.5 Interferences

Large amounts of extraneous cations eluting close to the cations of interest will result in an interference. No interferences have been observed in extracts analyzed by RTI to date. If interferences are observed, several steps to increase separation can be taken, such as reducing eluent strength and/or flow rate, or replacing the columns.

1.6 Personnel Qualifications

Personnel employed to perform ion analysis operations will have at least a Bachelor's degree in a laboratory science, and will be trained by a supervisor before being allowed to process client samples for the NPS program.
Apparatus and Materials

- Ion chromatograph (Dionex Model DX-500 with LC20 chromatography module, two IP20 isocratic pumps, two CD20 conductivity detectors, two Dionex AS40 automated samplers and PeakNet software) with Dionex CS12A cation separator column, and cation self-regenerating suppressor
- Pressurized eluent reservoirs
- Volumetric flasks - an assortment of sizes
- Dionex Autosampler Vials with filter caps

1.8 Reagents

Use ACS reagent grade chemicals for the preparation of all solutions

1. Concentrated Eluent Stock Solution: 1 N H$_2$SO$_4$ purchased from VWR Scientific

2. Eluent, 22mN H$_2$SO$_4$ Acid: Dilute 22mL 1N H$_2$SO$_4$ to 1 liter using deionized water.

3. Calibration Standard Stock Solutions, 1000 mg/L each.
   - NH$_4^+$: Dissolve 2.9654 g NH$_4$Cl in 1 liter deionized water
   - Na$^+$: Dissolve 2.5420 g NaCl in 1 liter deionized water
   - K$^+$: Dissolve 2.2284 g K$_2$SO$_4$ in 1 liter deionized water.
   - Ca$^{2+}$: Purchase NIST-traceable 1000 mg/L Ca$^{2+}$ standard solution from CPI International, Santa Rosa, CA.
   - Mg$^{2+}$: Purchase NIST-traceable 1000 mg/L Mg$^{2+}$ standard solution from CPI International, Santa Rosa, CA.

4. Standard Solution A: Add 10 mL of each stock solution (NH$_4^+$, Na$^+$, K$^+$, Ca$^{2+}$, and Mg$^{2+}$) to a 100 mL volumetric flask and dilute to the mark with deionized water and mix thoroughly (100 mg/L NH$_4^+$, Na$^+$, K$^+$, Ca$^{2+}$, and Mg$^{2+}$).

5. Standard Solution B: Dilute 10 mL Standard Solution A to 100 mL with deionized water (10 mg/L NH$_4^+$, Na$^+$, K$^+$, Ca$^{2+}$, and Mg$^{2+}$).
6. Calibration Standards: Using Standard Solutions A and B, prepare calibration standards with deionized water in 100 mL volumetric flasks as shown in Table 1. Prepare fresh calibration standards weekly.

**TABLE 1. PREPARATION OF CATION CALIBRATION STANDARDS**

<table>
<thead>
<tr>
<th>Standard</th>
<th>NH$_4^+$, Na$^+$, K$^+$, Ca$^{2+}$, Mg$^{2+}$ (mg/L each)</th>
<th>mL of Standard Solution/100 mL</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>STANDARD SOLUTION A</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>25.0</td>
<td>25.0</td>
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<tr>
<td>2</td>
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<td>3</td>
<td>3.0</td>
<td>3.0</td>
</tr>
<tr>
<td><strong>STANDARD SOLUTION B</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>1.0</td>
<td>10.0</td>
</tr>
<tr>
<td>5</td>
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<tr>
<td><strong>1 ppm STANDARD</strong></td>
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<td></td>
</tr>
<tr>
<td>6</td>
<td>0.1</td>
<td>10.0</td>
</tr>
<tr>
<td>7</td>
<td>0.05</td>
<td>5.0</td>
</tr>
</tbody>
</table>

7. Quality Assurance Stock Solutions: 1000 mg/L each NH$_4^+$, Na$^+$, K$^+$, Ca$^{2+}$, and Mg$^{2+}$, NIST Traceable: Purchase from GFS, Columbus, OH and/or EM Science, Gibbstown, NJ.

**Sample Collection**

Sample collection is not applicable to this SOP because samples are acquired by the PMCC.

**Sample Handling**

RTI will provide chain-of-custody documentation with all sample shipments to track and ensure that samples are collected, transferred, stored, and analyzed by authorized personnel; sample integrity is maintained during all phases of sample handling and analysis; and an accurate written record is maintained of sample handling and treatment from the time of its collection, through the laboratory analytical process, to the eventual relinquishing of all data to the client.
Filter Extraction Procedure

Anion and cation analyses will be performed on the same filter extract. Filters to be analyzed for anions and cations will be extracted with deionized water. The anion eluent (sodium carbonated, bicarbonate solution) produces a large sodium peak in the cation chromatogram that precludes quantitation of the sodium ion in the filter extract and interferes with the quantitation of ammonium ion. Extraction procedures are detailed in the RTI SOP “Standard Operating Procedure for NPS Filter Preparation, Extraction, and Anion Analysis.”

1.12 IC Procedure

1. Fill the eluent reservoirs with eluent.

2. Start the eluent flow, activate the self-regenerating suppressor, and allow the baseline to stabilize.

3. Inject two eluent blanks to flush the system and to ensure that the system is operating properly.

4. Using the calibration schedule, perform the daily multipoint calibration over the appropriate range followed by quality assurance/quality control (QC/QA) samples listed below:
   - a QC sample with NH$_4^+$, Na$^+$, K$^+$, Ca$^{2+}$, and Mg$^{2+}$ concentrations typical of the higher concentrations found in actual filter extracts.
   - a QC sample with NH$_4^+$, Na$^+$, K$^+$, Ca$^{2+}$, and Mg$^{2+}$ concentrations typical of the lower concentrations found in actual filter extracts.
   - a commercially prepared, NIST-traceable QA sample containing known concentrations of NH$_4^+$, Na$^+$, K$^+$, Ca$^{2+}$, and Mg$^{2+}$.

If the observed value for any cation differs by more than 10 percent from the known value, identify and correct the problem before analyzing samples.

5. Load the filter extracts into the autosampler vials according to the schedule prepared for that day. The daily schedule includes duplicate samples, spiked samples and QA/QC samples.

6. Begin the analysis run, occasionally checking to ensure that the system is operating properly.

7. Examine the data at the end of the run. If the NH$_4^+$, Na$^+$, K$^+$, Ca$^{2+}$, or Mg$^{2+}$ concentration of any extract exceeds the upper end of its calibration curve, dilute the extract appropriately and analyze that day or include with the samples to be analyzed the following day.
1.13 Calculations and Data Reduction

Peak areas are entered into the computer automatically where calculations are performed using a quadratic fit to the calibration data. The quadratic fit yields the following:

\[ y_i = ax_i^2 + bx_i + c \]

where:
- \( y \) = the calculated cation concentration, mg/L
- \( x \) = the instrument response

Initially, the calibration curve from 0.05 to 10.0 ppm is used for the calculation of the extract \( \text{NH}_4^+ \), \( \text{Na}^+ \), \( \text{K}^+ \), \( \text{Ca}^{2+} \), and \( \text{Mg}^{2+} \) concentrations. If a cation concentration exceeds 10 ppm, the extract is diluted appropriately (usually 5-fold) to bring the cation concentration into the calibration range and reanalyzed.

2.0 Quality Control and Quality Assurance

Compare the regression parameters (a, b, c and correlation coefficient) for the standard curves with those obtained in the past. If they exceed the control limits, stop the analysis and identify the problem.

Analyze QC data (see Section 1.12) at the beginning of every analytical run. Compare the results with those obtained during previous QC tests. If the observed concentration of any cation differs from the known value by greater than 10%, stop the analysis until the problem is identified and corrected. Analyze a duplicate sample, a QA/QC sample, and a spiked sample after at least every 20 field samples.

3.0 Reference

DRI Document No. 8068.1F4, Appendix D, Section 4.2