PRINCIPLE

Fluoxetine and norfluoxetine co-elute by our gas chromatographic methods unless chemically modified. Additionally, fluoxetine appears to break down on column. The TFAA (trifluoroacetic anhydride) derivatives of fluoxetine and norfluoxetine are stable and are well resolved from one another.

This derivitization method is used in cases where there are fluoxetine/norfluoxetine or other co-eluting compounds. For example, nortriptyline co-elutes with doxepin and desipramine co-elutes with cocaine. The resulting TFAA derivatives are then separated by routine gas chromatography analytical methods.

SAFETY

The handling of all samples, reagents and equipment is performed within the established laboratory safety guidelines detailed in the safety manual.

REAGENTS AND MATERIALS

All chemicals should be analytical reagent (AR) grade or higher. The chemical reagents required for the extraction procedure are prepared as indicated. In each case, the prepared reagent is stable for a minimum of six months. All new reagents are validated against the old ones prior to use.

1. Distilled or Deionized water
2. Buffer, pH 9.8
   Slowly add 212 g of Na₂CO₃ to 2500 mL of distilled water in a 4000 mL Erlenmeyer flask. Once this is completely dissolved, slowly add 168 g of NaHCO₃. Dilute to the mark with distilled water and mix on a stirring plate for one hour. Using a pH meter, verify that the pH is in fact 9.8. If the pH needs adjustment, use acid or base as needed to adjust to 9.8, document on the solution preparation sheet that the pH was checked. Date and initial preparation sheet.
3. n-Butyl chloride (1-Chlorobutane). Available from most chemical companies in Analytical Reagent Grade.
4. 0.5 N HCl.
   Slowly add 164 mL of concentrated HCl to 3500 mL of distilled water in a 4000 mL graduated cylinder. Dilute to the mark with distilled water and mix on a stirring plate for one hour.

   CAUTION: PREPARE THIS SOLUTION IN A FUME HOOD. USE APPROPRIATE SAFETY EQUIPMENT. ALWAYS ADD ACID TO WATER!
5. **Sodium carbonate**
   Slowly add 530 g of Na$_2$CO$_3$ to 3000 mL of distilled water in a 4000 mL Erlenmeyer flask. Dilute to the mark with distilled water and mix on a stirring plate for one hour.

6. **Toluene.** Available from chemical companies as AR grade.

7. **Trifluoroacetic Anhydride (TFAA).** Available from chemical companies.

**CALIBRATORS AND CONTROLS**

Calibrators and in-house controls are prepared according to the section “Preparation of GC Calibrators and Controls” with following drugs to be used in grouped pairs as needed.

Fluoxetine and norfluoxetine.

Amitriptyline and nortriptyline.

Doxepin and nordoxepin.

Imipramine and desipramine

**SPECIMEN PREPARATION**

The procedure is routinely applied to the following biological specimens and their aliquots unless otherwise specified:

- **Blood**: 2 mL of the undiluted specimen
- **Urine**: 2 mL for qualitative identification
- **If requested:**
  - **Bile**: 2 mL of a 1:5 dilution
  - **Brain**: 2 mL of a 1:3 homogenate
  - **Gastric Contents**: 2 mL of a 1:10 dilution
  - **Liver**: 2 mL of a 1:5 homogenate
  - **Vitreous Humor**: 2 mL of a 1:5 dilution

**DILUTION OF SPECIMENS**

Specimens are diluted as follows:

<table>
<thead>
<tr>
<th>Reviewed by:</th>
<th>Date:</th>
</tr>
</thead>
</table>
Bile 1:5  
1.0 mL of undiluted bile q.s. to 5 mL with deionized water.

Brain 1:3  
5.0 g of brain homogenized with 10 mL of deionized water.

Liver 1:5  
5.0 g of liver homogenized with 20 mL of deionized water.

Gastric Contents 1:10  
2.0 mL of liquid q.s. to 20 mL of deionized water, or 2.0 g of a solid specimen homogenized with 18 mL of deionized water.

**Note:** Do not use homogenates older than two weeks unless low sample size requires it. Discuss with supervisor and note in case record.

**Note:** The entire submitted amount of gastric contents must be weighed prior to homogenizing and sampling; total weight must be documented on reports.

**EXTRACTION PROCEDURE**

1. Obtain a worklist of TFAA derivative cases to be analyzed. Find and collect all samples on worklist and place in an empty rack. Allow samples to come to room temperature.

8. Obtain enough 16 x 150 mm screw-cap culture tubes (two per aliquot) and Teflon-lined caps to extract one blank sample, calibrator samples and all requested cases. Label the tubes appropriately. Tubes should bear the entire toxicology number (e.g. 97-1234, not 1234), the specimen type and any appropriate dilution.

9. Obtain the required calibrator and control working solutions from the refrigerator and allow them to equilibrate to room temperature. Record the lot numbers of the solutions used on the lot sheet.

10. Pipet 2.0 mL of a matching, negative matrix into each of the blank and calibrator tubes (the negative matrix used for gastric and vitreous humor is 2.0 mL of deionized water). Lightly cap these tubes to avoid possible contamination from other specimens.

11. Mix each specimen container by gentle inversion two to four times and pipet 2.0mL of each specimen into the appropriate tube. Make certain that the toxicology number on the specimen bottle exactly matches the number on the tube. Lightly cap each tube after pipetting to prevent contamination from other cases. If a blood specimen is being pipetted and a clot is encountered, homogenize the specimen via the Polytron or a glass manual homogenizer, document on the blood bottle that it was homogenized; in addition annotate the sequence list and the Case Analysis Form in the comments section. Continue in this fashion until all cases have been pipetted.

12. **Note:** Immediately re-file all case specimens and negative matrices. Do not leave this for a later time. Enter the date extracted in the Dataease database for each case picked up in the batch.

**NOTE: DO NOT WAIT UNTIL THE FOLLOWING DAY TO COMPLETE THE ABOVE!**

13. Spike the tubes with various calibrator solutions as indicated. Each drug analyzed in the batch must have its appropriate calibrators.
14. Spike the tubes with various control solutions as indicated.

<table>
<thead>
<tr>
<th>Final Concentration (mg/L)</th>
<th>Stock Solution Concentration (mg/L)</th>
<th>Spiking Volume (µL)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.05</td>
<td>100</td>
<td>1</td>
</tr>
<tr>
<td>0.2</td>
<td>100</td>
<td>4</td>
</tr>
<tr>
<td>1.0</td>
<td>100</td>
<td>20</td>
</tr>
<tr>
<td>2.0</td>
<td>100</td>
<td>40</td>
</tr>
</tbody>
</table>

15. Add 40 µL of 50 mg/L Methapyrilene Internal Standard (or the required internal standard) to each tube.

16. Pipet 1.0 mL of the pH 9.8 buffer into each tube. Mix thoroughly by vortexing contents of each tube briefly. Pipet 4.0 mL of n-butyl chloride into each tube and cap all tubes. Invert each tube to check for leaks. If a leak is found, transfer the tube contents to another appropriately labeled tube. Place all tubes on the mechanical shaker for 10 minutes using a low setting. Remove the tubes from the shaker and centrifuge for 10 minutes at ≈ 3000 rpm. If an emulsion is noticed, break up the emulsion by shaking or using a sterilized cotton tipped applicator, and re-centrifuge. If the emulsion still persists, see the supervisor for alternate methods.

17. Transfer the upper, organic layer of each tube to a new appropriately labeled tube which contains 2.0 mL of 0.5N HCl. Check that the number on the two tubes matches. Handle only one set of two tubes at once. Cap and briefly invert each tube to check for leaks. Place on the mechanical shaker for 10 minutes on a low setting. Remove the tubes from the shaker and centrifuge for 10 minutes at ≈ 3000 RPM.

18. Upon removal from the centrifuge, uncap each tube and aspirate the upper, organic layer to waste using a suction/aspiration apparatus. Pipet 1.0 mL of the sodium carbonate solution into the lower acid layer. Vortex briefly to mix.

**Note:** Addition of the sodium carbonate will cause release of CO₂ during the neutralization reaction which will result in foaming.

19. Pipet 200 µL of toluene into each tube. Cap all tubes and place on the mechanical shaker for ten minutes on a high setting. Remove the tubes from the shaker and centrifuge for 10 minutes at ≈ 3000 rpm.
20. Transfer the toluene layer into a clean, appropriately labeled 10 mL conical test tube. Add 100 µL of TFAA to each tube via Eppendorf. Immediately cap each tube after the addition of TFAA. Incubate tubes for 15 minutes at 90° C in a heating block or an oven.

21. Remove tubes from heat and allow tubes to cool to room temperature. Add 1.0 mL of pH 9.8 buffer and vortex for 5 seconds. Centrifuge for 10 minutes at ~3000 RPM. Using a Pasteur pipet, transfer the upper toluene layer to a glass insert in an appropriately labeled vial. After each transfer, immediately seal each vial with an aluminum seal, by using a hand crimper, to avoid possible contamination from other samples. Samples may also be transferred into screw cap vials and capped immediately. Do not wait until all transfers have been made to seal the vials. Transfer the vials to the GC Instrument and continue with analysis by gas chromatography.

INSTRUMENTATION ANALYSIS BY GAS CHROMATOGRAPHY

Instrumentation
The instrumentations used for analysis is the Agilent 6890 gas chromatograph equipped with an HP-17 column, a 7683 Series autosampler and a Agilent 6890 Series nitrogen-phosphorous detector. A computer with HP Chemstation software is used to gather the raw data and Chemstation Enhanced Data Analysis software is used to process the data.

HP-17 Column
The primary column used in the GC section is a 10.0 m x 0.53 mm x 2 µm HP-17 (50% phenyl methylsiloxane) megabore capillary column. This column is used for most routine work, both qualitative and quantitative.

INSTRUMENT SETUP

6890 Series II Gas Chromatogram
1. Ensure that the selected GC is operational and not in need of repair. If maintenance is required, consult the appropriate manual and notify the supervisor.
2. Make sure GC power on.
3. Check that the NPD bead voltage is set to current value as listed on the instrument.
4. The Chemstation method file for quantitation is usually named GCxNQNT.M, where the x refers to the instrument number. There should be a method file with the appropriate name in each method subdirectory. Load the appropriate method for the analysis. Consult supervisor if correct method is unclear.
5. On Chemstation verify that all other GC parameters are set correctly; go to Method, Edit Entire Method. Wait for “Ready”.

Method Section To Run:

(X) Save Copy of Method With Data
( ) Instrument Control Pre-Run Cmd/Macro =
( ) Data Analysis Pre-Run Cmd/Macro =
(X) Data Acquisition
( ) Data Analysis
( ) Instrument Control Post-Run Cmd/Macro =
( ) Data Analysis Post-Run Cmd/Macro =

**Inlet and Injection Parameters**

Sample Inlet: GC  
Injection Source: GC ALS

**GC Injector (7683 Series Injector)**

*Select Single or Dual*  
*Injector Parameters (Front or Back or Both)*

<p>| | | | | |</p>
<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Sample Washes</td>
<td>1</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sample Pumps</td>
<td>4</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Injection Volume</td>
<td>5.0 microliters</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Syringe Size</td>
<td>10.0 microliters</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PreInj Solvent A Washes</td>
<td>0</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PreInj Solvent B Washes</td>
<td>0</td>
<td></td>
<td></td>
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<tr>
<td>PostInj Solvent A Washes</td>
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<td></td>
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</tr>
<tr>
<td>PostInj Solvent A Washes</td>
<td>5</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Viscosity Delay</td>
<td>0 seconds</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Plunger Speed</td>
<td>Fast</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PreInjection Dwell</td>
<td>0 minutes</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PostInjection Dwell</td>
<td>0 minutes</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Inlet Parameters**

<table>
<thead>
<tr>
<th></th>
<th>Front Inlet</th>
<th>Back Inlet</th>
</tr>
</thead>
<tbody>
<tr>
<td>(Split/Splitless)</td>
<td>Mode</td>
<td>Mode</td>
</tr>
<tr>
<td>Mode</td>
<td>Splitless</td>
<td>Splitless</td>
</tr>
<tr>
<td>Initial Temp</td>
<td>275 °C (on)</td>
<td>275 °C (on)</td>
</tr>
<tr>
<td>Pressure</td>
<td>5.20 psi (on)</td>
<td>5.20 psi (on)</td>
</tr>
<tr>
<td>Pulse pressure</td>
<td>10.0 psi</td>
<td>Pulse pressure</td>
</tr>
<tr>
<td>Pulse time</td>
<td>0.60 min</td>
<td>Pulse time</td>
</tr>
<tr>
<td>Purge flow</td>
<td>30.3 mL/min</td>
<td>Purge flow</td>
</tr>
<tr>
<td>Purge time</td>
<td>2.00 min</td>
<td>Purge time</td>
</tr>
<tr>
<td>Total Flow</td>
<td>48.8 mL/min</td>
<td>Total Flow</td>
</tr>
<tr>
<td>Gas Saver</td>
<td>off</td>
<td>Gas Saver</td>
</tr>
<tr>
<td>Gas type</td>
<td>Helium</td>
<td>Gas type</td>
</tr>
</tbody>
</table>

Reviewed by: ___________________________  Date: ___________________________
Column Parameters

*HP-17 COLUMN (Column 1 & Column 2)*

### Capillary column

<p>| | |</p>
<table>
<thead>
<tr>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Model number</td>
<td>HP 19095L-121</td>
</tr>
<tr>
<td>HP-50+ 50% Phenyl Methyl Siloxane</td>
<td></td>
</tr>
<tr>
<td>Max Temp</td>
<td>310 °C</td>
</tr>
<tr>
<td>Nominal length</td>
<td>10.0m</td>
</tr>
<tr>
<td>Nominal diameter</td>
<td>530.00 um</td>
</tr>
<tr>
<td>Nominal film thickness</td>
<td>2.00um</td>
</tr>
<tr>
<td>Mode</td>
<td>Ramped Pressure</td>
</tr>
<tr>
<td>Initial Pressure</td>
<td>5.20 psi</td>
</tr>
<tr>
<td>Initial Time</td>
<td>12.00 min</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>#</th>
<th>Rate</th>
<th>Final Pressure</th>
<th>Final time</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>3.20</td>
<td>8.20</td>
<td>0.00</td>
</tr>
<tr>
<td>2</td>
<td>0.0</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Post flow 5.20 psi
Nominal initial flow 15.4 mL/min
Average velocity 131 cm/sec
Inlet Front, Back, or Both
Outlet (detector) Front, Back, or Both
Outlet pressure ambient

### Oven Parameters (6890)

- Initial temp: 120 °C
- Maximum temp: 280 °C
- Initial time: 2.00 min
- Equilibration time: 1.00 min

### Ramps:

<table>
<thead>
<tr>
<th>#</th>
<th>Rate (°C/min.)</th>
<th>Final Temp. (°C)</th>
<th>Final Time (min.)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>15.0</td>
<td>280</td>
<td>18.00</td>
</tr>
<tr>
<td>2</td>
<td>0.0 (off)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Post temp. 120 °C
Post time 1.00 min
Run time 30.67

### Detector Parameters

<table>
<thead>
<tr>
<th>Front detector (NPD)</th>
<th>Back detector(NPD)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Temperature</td>
<td>Temperature</td>
</tr>
<tr>
<td>325 °C mL/min(on)</td>
<td>325 °C (on)</td>
</tr>
</tbody>
</table>
Hydrogen flow | 3.0 mL/min (on) | Hydrogen flow | 3.0 mL/min (on)  
Air flow | 60.0 mL/min (on) | Air flow | 60.0 mL/min (on)  
Mode | Constant column+makeup flow | Mode | Constant column+makeup flow  
Combined flow | 30.0 mL/min | Combined flow | 30.0 mL/min  
Makeup flow | On | Makeup flow | On  
Makeup gas type | Helium | Makeup gas type | Helium  
Adjust offset | 30.00 | Adjust offset | 30.00  
Electrometer | On | Electrometer | On  
Bead | On | Bead | On  
Equilibration time | 0.00 | Equilibration time | 0.00  

### Signals
(Select Front, Back, or Both)

<table>
<thead>
<tr>
<th>Signal 1</th>
<th>Signal 2</th>
</tr>
</thead>
<tbody>
<tr>
<td>Data rate</td>
<td>20 Hz</td>
</tr>
<tr>
<td>Type</td>
<td>Front detector</td>
</tr>
<tr>
<td>Save Data</td>
<td>On</td>
</tr>
<tr>
<td>Start Save Time</td>
<td>1.00 min</td>
</tr>
<tr>
<td>Stop Save Time</td>
<td>30.00 min</td>
</tr>
<tr>
<td>Zero</td>
<td>0.0 (off)</td>
</tr>
<tr>
<td>Range</td>
<td>0</td>
</tr>
<tr>
<td>Fast Peaks</td>
<td>Off</td>
</tr>
<tr>
<td>Attenuation</td>
<td>0</td>
</tr>
</tbody>
</table>

6. After reviewing instrument parameters select a “Real Time Plot” for viewing on the instrument top screen prior to running the sequence.

7. Make sure that under “Select Reports” is all UNCHECKED.

8. Save the Method.

### SEQUENCE
Prepare a sequence by using the following steps:

1. Make sure the desired instrument session is up and running. Each GC has an instrument session, such as GC1, GC2, etc.

2. From the top menu select **Sequence → Load Sequence → Default.S**. Select **Sequence** again, then **Edit**. The **Sample Log Table** will open.

3. In the upper left hand corner on the **Sample Log Table**, select **Data Path**. Click on **Browse** to create a new folder under the **Data** folder of the instrument to be used.
4. In the upper right hand corner on the Sample Log Table, by clicking on Browse select the instrument Method to be used for analysis.

5. Type in all information under each column (i.e. Type, Vial, Sample, Method/Keyword, Datafile, Comments/Keywordstring, multiplier, multiple dilutions).

6. If the Sample is a calibrator, then under Comments/Keywordstring type-in the MIX or Filter it is for the batch; i.e. for all levels of CAL 1 type-in MIX [A], for levels of CAL 2 type-in MIX [B] etc. This step will be helpful during processing the data.

7. The Datafile name should be read GCXMMDDx001; where “X” is the instrument’s number, “MM” is the month, “DD” is the day, and “x” is the prefix of the sequence, such as “a”, “b” etc.

8. Once all the information is typed, select OK to close the Sample Log Table. Save the sequence by selecting Sequence followed by Save Sequence As . . . Name the sequence as follows: GCXMMDDYY; where “YY” defines the year.

9. Select Sequence then Simulate Sequence. In the window that opens, select Full Method, Inject Anyway, and Overwrite Existing DataFiles. For Sequence Comment enter the instrument # and detector used, the date of injection, the analyst initials, and the type of batch being injected. Select Run Sequence, verify all keywords by clicking “OK”.

10. Print sequence by selecting Sequence followed by Print . . . then select Brief. Add a “Chain of Custody”, a “Vials Loaded” label and a ECM documentation label to the hard copy of the sequence.

11. Empty solvent wash bottles on the instrument and replace it with fresh methanol before starting the sequence.

12. Use the printed sequence list to load the vials in the correct position on the autosampler tray. Edit the sequence as necessary to correct any errors; save the sequence. Reprint sequence if necessary.


15. Start sequence by selecting Sequence → Run Sequence.

16. When the sequence is complete, remove all vials using the sequence. Once again document the “Vials Loaded” label.

Setting-Up a Subsequence

On the Sample Log Table:

1. Select “Keyword” for “Type”

2. Select “DataPath” for “Method/Keyword”

3. Under “Comment/KeywordString” type in the new data path for your subsequence ie: C:\MSDCH2M\DATA\GCXMMMMDDYYx where “X” is the instrument’s number and “x” is the letter designated to the subsequence (it must be different than that of the original sequence).

4. Create a new folder for subsequence.
5. The suffix of the data files must be different from that of the original; ie: GCXMMDDx001; **the subsequence data files must start with 1 again.**
6. After typing in the entire sequence, save sequence accordingly.
7. Go to Sequence  Run Sequence.
8. A dialog box will pop-up: DataPath C:\MSDCHEM\2\DATA\GCXMMDDYYx does not exist. Edit Sample Log Table?  → Click No if the sequence was set up correctly.
9. A 2nd dialog box will pop-up: Create C:\MSDCHEM\2\DATA\GCXMMDDYYx?  → Click Yes.
10. A 3rd dialog box will pop-up: Sequence Verification Done! View it?  → Click Yes or No.
11. Save and Print Sequence.
12. Start sequence.

**DATA TRANSFER AND PROCESSING – exporting raw data for processing**

Retrieving Method from Data Acquiring Instrument (Raw Data)
1. Save the method to the batch from the acquiring instrument Chemstation.
2. Save method to batch by clicking ECM on the tool bar, then select **Save Method to ECM.** Locate the batch, select ok to save a copy of the method into the batch to be processed.
3. A copy of the method has now been saved to ECM and will accompany the batch when retrieved from ECM on any Data Analysis Processing Station.
4. Attach and document “ECM label”

**DATA REVIEW**

There are three levels of review; the first level of review is the transference and processing of the raw data, this may be performed by any trained analyst; the second level of review is performed by an experienced analyst who is trained and signed off in data review, he / she will review the processed data; the third level of review is considered the final level of review, this can only be performed by the Laboratory Manager. He/she will review the data for the entire case ensuring that screening, confirmatory and quantitative analysis on the case have been completed and reported accurately. As needed, he/she will also schedule additional analysis and contact the Medical Examiner on the case to discuss any findings and / or review case history.

**PROCESSING USING ENHANCED DATA ANALYSIS**

On any processing station:

<table>
<thead>
<tr>
<th>Reviewed by:</th>
<th>Date:</th>
</tr>
</thead>
</table>
1. Click on the Processing Data Analysis icon on the desktop and Log-in by using: analyst's OCME login for the username and their password.

2. From the tool bar menu select ECM → Retrieve Entire Sequence From ECM. Browse to locate and select the batch. Each data file will be retrieved along with the method to be used for processing.

3. Once the retrieval is complete the batch folder is located in the C:\msdchem\ECM\retrieve folder.

4. Select the method under the batch by right clicking on it and Load the method.

5. Select Calibrate → Clear; clear all responses and all calibrator levels.

6. Select Calibrate from the tool bar, then Update, Global Update, Set Curve Fit Type, Linear Regression Force (0, 0), select OK. If a different type of curve fit is used in place of Linear regression force (0,0), this must be annotated on the front page of the master copy of calibrators and controls that is appended to each case chromatogram.

7. Set up processing Filters (Filter A = Cal 1, Filter B = Cal 2, Filter C = Cal 3, Filter D = Cal 4, Filter E = Cal 5 etc.)
   a. Select Calibrate again, Edit Compounds; select Name, type in compound name in the space provided, click Find Compound.
   b. Change Compound Type from “T” to the required filter letter for that compound. All compounds for pools or Cals 1 & 2 are already assigned in all master methods (i.e. the compound type for meperidine is always A, doxylamine is always B etc.) and should not have to be assigned when processing each batch.
   c. Click OK to save all changes to the compounds that have been made.

8. In order to ensure that only compounds that are found are printed in the report select Quantitate in the menu bar, then Report Options. In the lower right hand box labeled “All Reports” make sure that the box labeled “Omit Target Compounds that are Missed” is checked.

9. Create the filters needed to process the batch:
   a. Select NYC_OCME in the menu bar, then select QDB Compound Type Filter.
   b. Select Create and type-in as many filters as needed for the batch one at a time (i.e. A, B, C, D etc). Filters are case sensitive.
   c. Once all filters are created a “qdb filter” window automatically pop-up with a list of all the compounds in the filters needed to process the batch.

10. Load the first calibrator at the highest level (i.e. 2.0 mg/L) by selecting File → Load Data File.

11. Load the filter needed for the data file:
   a. Select NYC_OCME in the menu bar, then select Compound Type Filter
   b. Type the # for that filter, where 1 is the first assigned filter.

12. In the space provided type in QT 1, then click on Execute; this processes the data file.
13. Select the **View** from the top menu, select **Easy ID** (this function allows for Retention times to be assigned for the entire method and batch). **Easy ID should only be used for your first calibrator level.**

14. To select a compound from the list double click, use the right click to integrate a peak for the compound, use the left click to control zooming on the chromatogram.

15. Once all of the retention times have been assigned, click on **Exit** and click on **Yes** to “update retention times to the compound list”.

16. Type in **QT 1**, then click on **Execute**.

17. Again select the **View** from the top menu, select **Q-Edit Quant Result** (this function to view the response and retention times for each compound.) **Cal 2.0 mg/L will not have an “x” next to each compound name because it is the first calibrator to be processed**, Click **Exit** when all compounds have been reviewed.

18. Assign the calibrator level:
   a. Select **Calibrate**, then **Update**, select **Update 1 Level**.
   b. Select **Add Level** and type in the concentration of the calibrator (i.e. 2.0, 1.0 etc.).
   c. Type-in the internal standard concentration (1.0).
   d. On the right hand side of the window type-in the Level ID i.e. 4A, 3A, and 2A, 1A; where the # is the calibrator level and the letter is the filter.
   e. Click on **Do Update** then select **OK**.

19. Load the next data file, Cal 1.0 mg/L. Type **QT 1** then click on **Execute**.

20. Select the **View**, select **Q-Edit Quant Result**. **Easy ID is not needed for this step, as retention times have already been set for this calibrator.** Double check each of the compound’s retention time and that each compound has a response. All auto-integrated compounds should have an “x” next to their name. Click **Exit** once when finished.

21. Assign the calibrator level.

22. Repeat steps 15-21 for the remaining levels (0.2 and 0.05mg/L) for the first calibrator.

23. Requant the first calibrator:
   a. Go to **Tools, DOLIST**.
   b. Remove any options that are on the right hand side of the box.
   c. On the left hand menu select “**Summary Quant No Report**” which is **QT 2**, with arrows move this command to the empty space on the right then click **OK**.
   d. Select the calibrators just processed and move them to the empty space on the right then click **Process**. Each file will be “requanted”.

Reviewed by:  
Date:
e. Review all files through Q-Edit Quant Result to verify quant results.

24. Process the corresponding QC’s for the calibrator that was just reviewed, with the same filter on:
   a. Go to Tools, DOLIST.
   b. Remove any options that are on the right hand side of the box.
   c. On the left hand menu select “Quant No Report” which is QT 1, with arrows move this command to the empty space on the right then click OK.
   d. Select the QC’s to be processed and move them to the empty space on the right then click Process. Each file will be processed.

25. Review each QC by selecting View, select Q-Edit Quant Result.

26. With the filter still on, print the Cal’s and QC’s for that filter:
   a. Go to Tools, DOLIST.
   b. Remove any options that are on the right hand side of the box.
   c. On the left hand menu select “Summary Quant w/o Calculations” which is QT 0, 1, “s”, with arrows move this command to the empty space on the right then click OK.
   d. Only select all Cal levels and QC’s just reviewed and quanted and move them to the empty space on the right then click Process. Each file will print.

27. Load the 2.0mg/L of the next calibrator to be processed and follow steps 10-26.

28. Once all calibrators and QC’s have been processed and printed, save the processing method to ECM then set the filter # to 0, which gives a list of all the compounds for the entire method, and process the all the blank injections and cases:
   a. Go to Tools, DOLIST.
   b. Remove any options that are on the right hand side of the box.
   c. On the left hand menu select “Quant No Report” which is QT 1, with arrows move this command to the empty space on the right then click OK.
   d. Select the QC’s to be processed and move them to the empty space on the right then click Process. Each file will be processed.

29. Review each blank and case through Q-Edit Quant Result:
   a. Deselect any compounds that are not needed in each blank or case by double clicking on it and clicking on QDel.
   b. Check the responses and retention times of the needed compounds.
   c. If a compound is needed but not integrated, double click on the compound and integrate.
   d. Click Exit once when finished, and click “yes” to the save changes dialogue box.

30. If a compound is too low to be integrated or no peak is present include an overlay:
   a. Go to Tools, Overlay Chromatogram.
   b. Select the calibrator or standard to overlay on the case; multiple overlays can also be set.
c. Select Process.
d. Go to File, Print then select Chromatogram

31. Once all blanks and cases are processed, print as directed in step # 26.

32. Print an $R^2$ report by:
   a. Go to Calibrate, List then select Calibration Report and OK.
   b. Compounds that were not used in the batch can be deleted by highlighting them and simply pressing delete on the keyboard. Caution should be used, do not remove compounds needed.
   c. Right click on the list and select Print, choose the correct printer.

33. Once everything is printed, save the all the data files from the batch back to ECM:
   a. Select ECM from the top tool bar; select Save Multiple Data Files to ECM.
   b. Select all data files on the left and move them to the right empty space, then click Process.
   c. Allow each data file to be successfully copied to ECM before exiting Data Analysis.

34. After uploading the files to ECM, delete the sequence from the C: Drive of the processing station:
   a. Right click on the Start on the lower left hand corner and select Explore
   b. Open the MSDCHEM folder on the C: Drive, then select 1 → ECM → Retrieve
   c. Locate the batch processed, right click on it and delete it.
   d. Document" ECM label" on sequence list.

RE-INJECTION CRITERIA

Occasionally, samples analyzed may need to be reinjected for a variety of reasons. The criteria for reinjection, other than poor chromatography or requests made by the appropriate supervisor, are listed below.

1. Individual blood samples may be re-injected once if internal standard (IS) area counts fall below that of the counts for the Cal’s and QC’s in the batch.

2. Analytes in a calibrator or control are not present in the chromatogram.

3. Additional peaks present in the chromatogram of a calibrator or control. See TROUBLESHOOTING OF AGILENT GAS CHROMATOGRAPHS section of the SOP manual.
4. QC failure: reinject the blank, calibrators QC’s and the associated case(s). If upon re-injection there is still a QC failure schedule the cases for re-extraction, make a note in DataEase and on the appropriate Case Analysis Form(s).

5. Examine peak shape for signs of overload. If peaks are still overloaded after reinjection, re-extract samples with overloaded peaks using appropriate dilutions. If a calibrator peak is still overloaded after re-injection, the batch must be re-extracted.

CARRYOVER

Two types of carryover are possible with the methods employed by GC. One is the “carryover” of late-eluting peaks from one injection into the next (or even next two or three) injection. This is usually apparent from the shape of the peak (broad).

The other possibility is “carryover” due to contamination from an excessive amount of drug in the previous sample, either through the syringe or by drug being retained in the injection port or column. Any samples with large concentration of drug are rescheduled with dilutions, to reduce the amount to within the linear range of the method. If it appears that carryover of this type has occurred, the reviewer will reschedule the case after the overload to verify that there was no contamination.

Frequent changing of the autosampler wash vials helps to control this type of potential carryover.

ACCEPTANCE CRITERIA

1. In-house controls must be analyzed in accordance with the standard laboratory procedures to determine the target and acceptable range before being incorporated into routine analysis of the specimens.

2. None of the target analytes should be detected in the negative control (blank).

3. Retention times of target analytes should match those of the calibrator or controls within ± 2%.

4. The positive controls should be within ± 20%, for blood batches, or ± 30%, for tissue batches, of the established target for all analytes detected in the samples. In other words, if fluoxetine and norfluoxetine are present in the samples, the respective components of the controls must meet the acceptance criteria, but it is not necessary for the other compounds in that same control to meet the acceptance criteria. If the QC fails for quantitation, but is correct qualitatively, a batch can be accepted for qualitative results only. Acceptance of controls outside of ± 20% blood or ± 30% for tissues, of the established target require authorization of the supervisor and documentation.

5. The analyst shall choose from the standard levels a level at their discretion. All internal standard areas/peak heights for that sequence shall be compared to the internal standard(s) in that level. The acceptance criteria for the internal standard recovery/response shall be ± 30% of the selected internal standard. If the internal standard recovery/response is outside these limits the specimen shall be repeated unless the analyte in question is not related to the internal standard that is outside the acceptable limits. If the internal standard is still
unacceptable in the repeat analysis in the same way (i.e. both times too high or both times too low), then the sample shall be repeated a third time using a “blank and spike” method. The “blank and spike” run shall be viewed as acceptable if the following conditions are met: All standards and QC materials are within acceptable ranges. The “blank and spike” samples internal standards are within ± 20% of their average responses. The calculated amount of the spiked sample is ± 20% of the theoretical amount. If any of these conditions are not met then consult with a toxicology manager for further direction.

DATA REVIEW – FIRST LEVEL REVIEW
6. Check chromatography of all injections. Examine the peak shape and note if any peaks show non-Gaussian shape. Evaluate unresolved peaks and peaks with shoulders on either side. Address any noted problems before proceeding. Consult a supervisor about any unusual events.

7. Check that all significant peaks in the chromatogram are integrated. If the printout allows, check if the baseline used to integrate is appropriate. The analyst should consult with a supervisor about any unusual events, such as the presence of overload peaks.

8. Check that all components of each calibrator are present and that each peak is properly assigned. Correct any errors. If any calibration compounds do not extract, or if expected multiple peaks are detected for a calibration compound, the analyst must consult with a supervisor.

9. Check the blank for significant peaks that might co-elute with a compound of interest. There is some room for judgment in this step, but the analyst should consult with a supervisor if there are ANY questions. A blank may be scheduled for mass spectrometry to discover the identity of peaks in a blank. If any significant peaks are present in a blank for the batch, the run must be rejected.

10. Review QC samples, determine if the controls meet all acceptance criteria and verify the concentration of the components.

11. A copy of the sequence calibrators and controls are forwarded to the QC officer, who will enter the results of the QC sample in the QC database. If the batch fails, annotate sample chromatograms, listing the reason for failure. Notify a supervisor and / or the QC manager.

12. Annotate the following information on the Calibrator chromatogram:
   a. Calibrators and blank reviewed and accepted (or not accepted). Any QC or calibrator failures are annotated on the proper chromatograms.
   b. Note any deviations from the SOP in a concise but detailed fashion.
   c. Initial and date.
   d. Regression coefficient ($r^2$) must be greater or equal to 0.98.

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8. Make sure that each control has the appropriate target concentration range label.

9. All data should be photocopied using double sided mode. A master copy of the sequence list, calibration, control and any other data is prepared. Arrange this master copy as follows (from top to bottom): Calibrators, all QC samples and blanks, r2 report, sequence list, internal standard recovery form and the lot sheet form. It is the responsibility of the analyst to verify that the master copy is legible and is an accurate copy of the originals, with no information cut off at the margins. Photocopies are made from this master, one for each case in the batch.

10. Submit data for Second Level Review.

DATA REVIEW – SECOND LEVEL REVIEW

Quantitative Results
The Second Level Reviewer will review the processed data in its entirety according to the criteria listed in the data review section and acceptance criteria section of this SOP. The second level reviewer will also ensure the following:

1. The reviewer should review the calibration, control, and sequence information; use the same criteria as listed in the data review section of this SOP, with the additional criteria that the $r^2$ of the four-point calibration curve of each component present in any case in the batch must be equal to or greater than 0.98. Make any necessary corrections before proceeding.

2. Retrieve the case file for each sample in the batch and associate it with its corresponding chromatogram(s). Ensure a copy of the reviewed calibration, control, and sequence information is placed with each original case chromatogram.

3. Review the case’s chromatogram. Determine if the results are consistent with all the other applicable case data to date. For quantitation, If the results are consistent, report the results on the Result Summary Sheet. Date and initial both the chromatogram and the Result Summary Sheet. If the results are not consistent, the inconsistency must be resolved. Reschedule the sample if necessary, or take other appropriate corrective action.

4. Repeating analysis may be necessary if the chromatogram shows overloaded peaks or poor recovery of the internal standard. If the problem is an overload, reschedule with appropriate dilutions. In cases of poor recovery, the case, controls, negative controls, calibrators may be re-injected once. If the criteria as listed in batch review are still not met, the sample is re-extracted. If any cases have an unknown or unidentified peak not observed in previous scan results, the case vials must be transferred to GC/MS for identification. Consult with the supervisor if there are any questions.

5. If no rescheduling is necessary submit the appropriate case file for third level review.

Note: Do not discard any sample paperwork.
REPORTING (QUANTITATIVE RESULTS)

After the batch has undergone second level review and has been printed, either the first level reviewer or the second level reviewer may report the data in the appropriate case file

1. All control acceptance criteria must be met.
2. Analytes detected at a concentration of 0.05 mg/L (lowest calibrator) or 0.025 mg/L (lowest control) are reported as “< (less than...)” the lowest calibrator or control (i.e. if the analytes concentration is < 0.025 mg/L, report as less than 0.025 mg/L).
   a. If 0.025 mg/L control passes and the analytes concentration is greater than the 0.025 mg/L control, report the numerical value.
   b. If 0.025 mg/L control fails and the lowest calibrator (0.05 mg/L) passes report the analytes concentration as less than 0.05 mg/L.
3. If the lowest calibrator and control fail the case must be repeated.
4. All positive findings are reported to two decimal points (e.g., “fluoxetine 0.82 mg/L”).
5. Drugs detected at a concentration above the upper limit of quantitation must be reported as “greater than...”, unless the concentration was determined by repeat analysis using appropriate dilutions. Dilutions are performed until the results are within the linear range of the calibration curve. If the sample is insufficient or other considerations preclude dilution, report as above.
6. Drug facilitated sexual assault cases and driving under the influence cases are reported in ng/mL

THIRD LEVEL REVIEW (FINAL REVIEW)

The third and final level review can only be performed by the Laboratory Managers. He/she will review the data for the entire case according to all established criteria. They will ensure that screening, confirmatory and quantitative analysis on the case have been completed and reported accurately. As needed, they will also schedule additional analysis and contact the Medical Examiner on the case to discuss any findings and / or review case history.
REVISION HISTORY

Ver 03.08.2013  1. Revision history implemented.
Ver 10.03.2013  1. Changed sample volume to 2.0 mL.
Ver 04.20.2015  1. Included a calibration level inadvertently omitted in the previous version.
                2. Minor grammatical & spelling corrections
Ver 08.31.2015  1. Added internal standard acceptance criteria.
Ver 09.04.2015  1. Defined the level of review(s).
Ver 09.16.2015  1. Minor grammatical errors.
                2. Removed calibrator level @ 3.0 mg/l
                3. Changed reporting criteria.