

## Section Five

### Quality Assurance

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#### 5.12 Solution Preparation

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##### 5.12.1 BACKGROUND

Refer to references.

##### 5.12.2 SCOPE

This section describes the proper preparation of solutions and buffers used in the extraction of drug compounds from blood and urine specimens.

##### 5.12.3 EQUIPMENT AND SUPPLIES

###### 5.12.3.1 Glassware

Adequately sized beakers, volumetric flasks, graduated cylinders and volumetric pipettes

###### 5.12.3.2 Laboratory balance

###### 5.12.3.3 pH Meter and/or Indicator Strips

###### 5.12.3.4 Appropriate buffer solutions for pH meter

###### 5.12.3.5 Stirring hotplate

###### 5.12.3.6 Magnetic stirrers

###### 5.12.3.7 Safety Equipment

- Chemical Fume Hood
- Acid Resistant Apron
- Laboratory Coat
- Safety Goggles and/or face Shield
- Laboratory Gloves

##### 5.12.3 REAGENTS

All chemicals must be ACS Grade or equivalent.

###### 5.12.4.1 Acids

- Acetic, Glacial
- Hydrochloric
- Phosphoric
- Sulfuric

###### 5.12.4.2 Salts

- Ammonium Chloride
- Potassium Hydroxide
- Potassium Phosphate Monobasic
- Potassium Phosphate Dibasic
- Sodium Acetate Trihydrate
- Sodium Bicarbonate
- Sodium Hydroxide

- Sodium Phosphate Monobasic
- Sodium Phosphate Dibasic
- Sodium Tetraborate Decahydrate

5.12.4.4 Solvents

- Methanol

## 5.12.5 PROCEDURES

Preparation of the following solutions must be recorded on corresponding preparation log. Solutions may be made in different volumes by adjusting reagent ratios.

*Note: Appropriate safety equipment must be worn during the preparation of solutions to prevent exposure to caustic/corrosive solutions. The order of the addition of chemicals may be crucial to prevent exothermic reactions. Refer to appropriate MSDS sheets for more information handling chemicals.*

### 5.12.5.1 Acetic Acid

#### 5.12.5.1.1 **1.0M Acetic Acid (500mL)**

Place approximately 400mL DI water into a 500mL volumetric flask. Add 29mL **glacial acetic acid**, mix. QS to 500mL.

*A positive and negative control will be run with each use. Remake as indicated by control data.*

#### 5.12.5.1.2 **20% Acetic Acid (500mL)**

Place approximately 300mL DI water into a 500mL volumetric flask. Add 100mL glacial acetic acid, mix. QS to 500mL.

*A positive and negative control will be run with each use. Remake as indicated by control data.*

### 5.12.5.2 Ammonium Chloride

#### 5.12.5.2.1 **Saturated Ammonium Chloride (500mL)**

Place approximately 300mL DI water in a beaker and heat/stir over low heat. Add **ammonium chloride** until the solution is saturated. QS to 500mL.

*A positive and negative control will be run with each use. Remake as indicated by control data.*

### 5.12.5.3 Borate Buffers

**5.12.5.3.1 Borate Buffer, pH 9.2**

Place  $\cong$ 500mL DI water into a 1000mL beaker. Heat and stir while adding 50g sodium tetraborate ( $\text{Na}_2\text{B}_4\text{O}_7 \cdot 10 \text{H}_2\text{O}$ ). Once dissolved, allow to cool. Bring volume up to  $\cong$ 950mL with DI water. Verify pH and adjust as necessary to  $\text{pH } 9.2 \pm 0.2$  with 1N KOH or 100mm HCl. Place solution in 1000mL volumetric flask and QS with DI water.

*Solution is stable for six months.*

**5.12.5.3.2 Borate Buffer, pH 12**

Place  $\cong$ 500mL DI water into a 1000mL beaker. Heat and stir while adding 50g sodium tetraborate ( $\text{Na}_2\text{B}_4\text{O}_7 \cdot 10 \text{H}_2\text{O}$ ). Once dissolved, allow to cool. Bring volume up to  $\cong$  900mL with DI water. Add 25mL 10N NaOH and stir. Verify pH and adjust as necessary to  $\text{pH } 12 \pm 0.2$  with 10N NaOH or 6N HCl. Place solution in 1000mL volumetric flask and QS with DI water.

*Solution is stable for six months.*

**5.12.5.4 Hydrochloric Acid****5.12.5.4.1 0.1M/100mM Hydrochloric Acid (500mL)**

Place approximately 300mL DI water into a 500mL volumetric flask. Add 4.2mL **concentrated hydrochloric acid**, mix. QS to 500mL.

*A positive and negative control will be run with each use. Remake as indicated by control data.*

**5.12.5.4.2 1% HCl in Methanol (10mL)**

Add approximately 5mL of methanol to a 10mL volumetric flask. Pipet 100 $\mu$ L of **concentrated HCl**, QS and mix. Store in a brown glass bottle.

*Solution is stable for three-months.*

**5.12.5.5 Potassium Hydroxide (KOH)**

*Note: The addition of KOH to water will generate significant heat, exercise due caution.*

**5.12.5.5.1 1M/1N Potassium Hydroxide (100mL)**

Dissolve 5.6g **potassium hydroxide** in approximately 80mL DI water in a 100mL volumetric flask. QS to 100mL.

*A positive and negative control will be run with each use. Remake as indicated by control data.*

5.12.5.5.2 **11.8N Potassium Hydroxide (1000mL)**

Gradually(!) add 662g **potassium hydroxide** to approximately 600mL DI water, stir on stir plate to dissolve. Allow to cool (this takes awhile) and QS in a 1000mL volumetric flask.

*A positive and negative control will be run with each use. Remake as indicated by control data.*

5.12.5.6 Potassium Phosphate Buffers

5.12.5.6.1 **Saturated Potassium Phosphate Buffer (1000mL)**

Place approximately 1000mL DI water in a beaker and heat/stir over low heat. Add **potassium phosphate monobasic** until the solution is saturated. Allow solution to cool. Adjust pH to approximately 1.8 with **concentrated phosphoric acid**.

*A positive and negative control will be run with each use. Remake as indicated by control data.*

5.12.5.6.2 **0.1M/100mM Potassium Phosphate Buffer (100mL) - Adjusted to pH 6**

Dissolve 1.36g **potassium phosphate monobasic** in  $\approx$ 90mL DI water in a 150mL beaker. Adjust to pH 6.0 with 1.0M **potassium hydroxide**. QS in a 100mL volumetric flask. *Store in brown glass container. Solution is stable for 6-months.*

5.12.5.7 Sodium Acetate Buffers

5.12.5.7.1 **0.1M/100mM Acetate Buffer, pH 4.5 (500mL)**

Dissolve 2.93g **sodium acetate trihydrate** in 400mL DI water in a 600mL beaker. Add 1.62mL **glacial acetic acid**, and mix well. Adjust to pH  $4.5 \pm 0.1$  with **glacial acetic acid** or **100mM acetic acid**. QS to 500mL in a 500mL volumetric flask.

*A positive and negative control will be run with each use. Remake as indicated by control data.*

5.12.5.7.2 **0.1M/100mM Acetate Buffer, pH 5.0 (500mL)**

Prepare as with pH 4.5 buffer (5.12.5.9.1). Adjust pH to  $5.0 \pm 0.1$ .

*A positive and negative control will be run with each use. Remake as indicated by control data.*

5.12.5.7.3 **2.0M Acetate Buffer, pH 4.8 (1000mL)**

Dissolve 141.4g **sodium acetate trihydrate** in  $\approx$ 800mL DI water. Add 55.2mL **glacial acetic acid**. Adjust to pH 4.8 and QS to 1000mL.

*A positive and negative control will be run with each use. Remake as indicated by control data.*

5.12.5.8 Sodium Bicarbonate

5.12.5.8.1 **50mM Sodium Bicarbonate, pH 11 (500mL)**

Dissolve 2.1g **sodium bicarbonate** in 500mL DI water.

*A positive and negative control will be run with each use. Remake as indicated by control data.*

5.12.5.9 Sodium Hydroxide (NaOH)

*Note: The addition of NaOH to water will generate heat, exercise due caution.*

5.12.5.9.1 **2N NaOH (1000mL)**

Place approximately 250mL DI water into a 1000mL beaker. Gradually add 80g **NaOH**. Transfer to 500mL volumetric flask and QS to 500mL. (Caution: Exothermic)

*A positive and negative control will be run with each use. Remake as indicated by control data.*

5.12.5.9.2 **10N NaOH (500mL)**

Place approximately 400mL DI water into a 1000mL beaker. Gradually add 200g **NaOH**. Transfer to 500mL volumetric flask and QS to 500mL. (Caution: Exothermic)

*This reagent is used in the preparation of other reagents those reagents are checked with each use.*

5.12.5.10 Sodium Phosphate

5.12.5.10.1 **100mM Sodium Phosphate Dibasic (200mL)**

Dissolve 2.84g **sodium phosphate dibasic** in  $\approx$ 160mL DI water. QS to 200mL and mix.

*Store in glass container. A positive and negative control will be run with each use. Remake as indicated by control data.*

**5.12.5.10.2 100mM Sodium Phosphate Monobasic (200mL)**

Dissolve 2.76g **sodium phosphate monobasic** in  $\approx$ 160mL DI water. QS to 200mL and mix.

*Store in glass container. A positive and negative control will be run with each use. Remake as indicated by control data.*

**5.12.5.11 Sodium Phosphate Buffers****5.12.5.11.1 0.1M/100mM Sodium Phosphate Buffer (1000mL) Adjusted to pH 6**

Dissolve 1.70g **sodium phosphate dibasic** ( $\text{Na}_2\text{HPO}_4$ ) and **12.14 sodium phosphate monobasic** ( $\text{NaH}_2\text{PO}_4 \cdot \text{H}_2\text{O}$ ) in approximately 800mL DI water in a 1000mL volumetric flask. QS to 1000mL. Adjust to pH 6.0  $\pm$ 0.1 with 100mM **monobasic sodium phosphate** (to lower pH) or 100mM **dibasic sodium phosphate** (to raise the pH).

*Check pH prior to use for blood toxicology casework; if pH outside preparation tolerance, remake buffer. Store in glass container. A positive and negative control will be run with each use. Remake as indicated by control data.*

**5.12.5.12 Sulfuric Acid****5.12.5.12.1 0.05M/0.1N Sulfuric Acid**

Place approximately 800mL distilled/deionized (DI) water into a 1L volumetric flask. Add 2.7mL **concentrated sulfuric acid**, mix. QS to 1L.

*A positive and negative control will be run with each use. Remake as indicated by control data.*

**5.12.6 QUALITY ASSURANCE**

5.12.6.1 Refer to toxicology Analytical Method 5.2 for balance intermediate check and calibration requirements.

Note: Balances properly monitored by drug discipline analysts fulfills quality assurance requirements. Additional check need not be performed.

**5.12.7 REFERENCES**

5.12.7.1 Shugar, G.J., Shugar, R.A. and Bauman, L. *Grades of Purity of Chemicals* pp. 145-154, *pH Measurement*. pp. 232-234. *in:*

Chemical Technicians' Ready Reference Handbook, McGraw Hill: New York, 1973.

5.12.7.2 Ansys, Inc. SPEC Extraction Methods

5.12.7.3 United Chemical Technologies, Inc. Applications Manual.

## Revision History

### Section Five

### Quality Assurance

## 5.12 Solution Preparation

Revision #	Issue Date	History
0	05-07-2007	Combined urine solution preparation (2.6) and blood solution preparation (3.8).
1	08-20-2008	Removed obsolete solutions, added reference for balance check requirements, clarifications.
2	7/8/2011	For solutions known to have a long shelf life, language was added to allow use longer than previously indicated. The following statement is in place of a definite expiration date. <i>A positive and negative control will be run with each use. Remake as indicated by control data.</i> Solutions that were no longer being used in toxicology analytical methods were removed. Solutions that were duplicated but listed in different volumes were removed. A statement allowing different volumes of solutions to be made was added. A statement was added to the safety note referencing MSDS sheets. Numbering updated.