NEW MODEL FOR BIOMONITORING OF OCCUPATIONAL CHEMICAL EXPOSURE TO VOLATILE ORGANIC COMPOUNDS

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- In the Years 2006-2010, we have tested a New Model for Biological Monitoring of Occupational Chemical Exposure to obtain more accurate and precise results, considering the low, medium or nonspecificity of some metabolites, used as bioindicators of exposure to Volatile Organic Compounds (VOCs).
- We have analyzed approximately 5, 347 urine samples of workers at risk of exposure to Benzene, Toluene, Xylene, Styrene and Ethylbenzene.
- These chemicals are produced, stored and transported by **PEMEX**.

Policy, Procedures and a Work Team that Make Possible the BOHRA

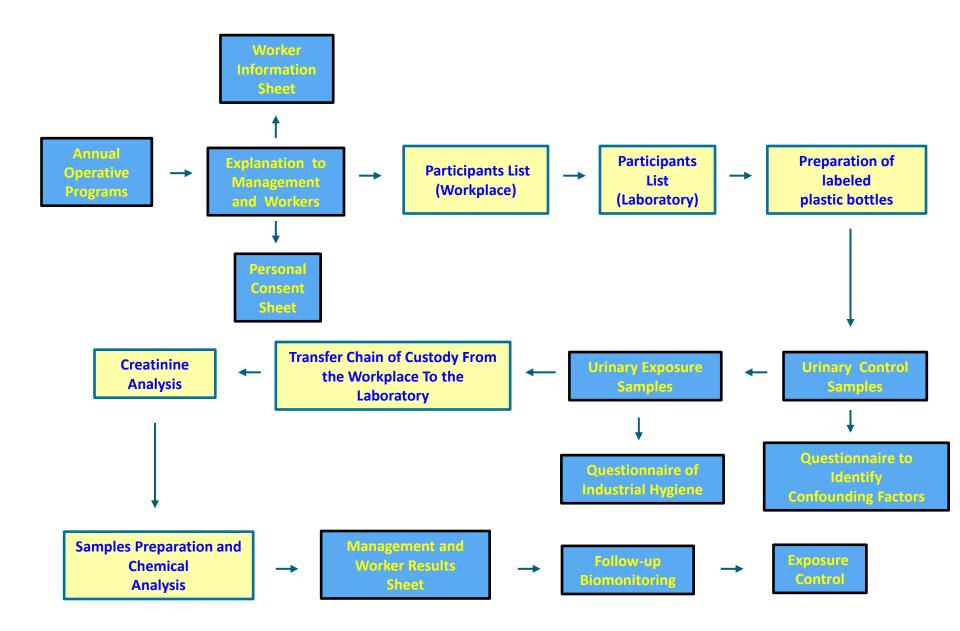
• Policy and Procedures of PEMEX:

- PEMEX Institutional «SSPA» : Administrative System for Industrial Safety, Occupational Health and Environmental Protection.
- PEMEX Institutional «**SAST**»: Occupational Health Administrative System.
- **Technical Guide** of the **BRANCH OF PEMEX HEALTH SERVICES** for Occupational Biomonitoring.
- **Procedure** of **PEMEX PETROQUIMICA** for Occupational Biomonitoring.
- Working Team that participates in the BOHRA:
 - Workplace Administration
 - Union Representatives
 - Workers themselves
 - Occupational Physicians
 - Industrial Hygiene Personnel
 - Industrial Nurses

Biomonitoring Practice in Our Industrial Toxicology Lab.

<u>Chemical Agent</u>	<u>Determinant</u>	<u>*Biological</u> <u>Exposure</u> <u>Index (</u> The underlined concentrations are used in our Laboratory)	<u>Specificity</u> Ns: <u>Nonspecific</u>	<u>Reference</u>
Benzene	Urinary Phenols (Used in years: 2006-2010) *Urinary S-	***50 mg/g Creat. **Tentative Maximum Permissible Concentration: <u>20 mg/g Creatinine</u>	Low, Ns	 ***Mexican Official Standard, NOM-047-SSA1-1993. **Lauwerys Robert., Casarett & Doull's Toxicology, Chapter 33: Occupational Toxicology, Edit. Klaassen D. Curtis, Mc Graw Hill, Fifth Edition, p. 999, 1996.
	Phenylmercapturic Acid (Used since 2011)	<u>*25 microg/g Creat.</u>	High	*ACGIH, 2009.
Toluene	Urinary Hippuric Acid	***2.5 g/g Creat <u>*1.6 g/g Creatinine</u>	Low, Ns	***Mexican Official Standard, NOM-047-SSA1-1993. *ACGIH, 2009, NIC.
Xylenes	Urinary Methylhippuric Acids	* ***1.5 g/g Creatinine	High	*** Mexican Official Standard, NOM-047-SSA1-1993. *ACGIH, 2009.
Styrene	Urinary Mandelic Acid plus Phenylglyoxylic Acid	*400 mg/g Creatinine	Medium, Ns	*ACGIH, 2009.
and Ethylbenzene		<u>*0.7 g/g Creatinine</u>	Medium, Ns	*ACGIH, 2009.

The Model Proposed for BOHRA of VOCs.



Explanation to Management and Workers

 It is very important to explain the objectives of Biomonitoring and make participate the Administration, the Workers and their Representatives. **Worker Information Sheet**

Contains information about:

- What is Biomonitoring of Occupational Chemical Exposure?
- Why is useful?
- With indications to the workers of:
 - not to smoke and avoid passive smoking
 - not to eat red meat
 - avoid canned food, plums, blueberries.
 - drink enough water
 - work as every day

Worker Consent Sheet

(International Code of Ethics for Occupational Health Professionals, International Commission on Occupational Health (ICOH), Point 12, 2002).

Contains:

- Information about the risks in the practice of Biomonitoring.
- **Consent of each worker through his signature** to participate in the Biomonitoring.
- The compromise to give to each worker the results with an understandable interpretation



- General Population Baseline
- Non Exposed Workers Baseline
- Individual Worker Baseline

Baseline of the Individual Worker

 The Baseline of the <u>Individual Worker</u> is obtained with the measurement of the concentration of the chosen bioindicator in the <u>Urinary Control Sample</u>, <u>collected prior to shift</u>.

• This is possible, in the case of VOCs, because the Biological Half Life of VOCs is <u>medium</u> (between 1.5 and 5.7 hrs.), and the Urinary Control Sample represents the recent exposure, but not, for instance, in the case of Metals or other Organic Compounds, where the Biological Half Life is long (between 10 and 100 hours or more).

Elimination Half Lives* of the Volatile Organic Compounds (VOCs) Analyzed in our Laboratory

Chemical	Metabolite	Hours
Benzene	Urinary Phenols	5.7
Toluene	Urinary Hippuric Acid	1.5
Xylene	Urinary Methylhippuric Acids	3.6
Styrene	Urinary Mandelic Acid	4
Ethylbenzene	Urinary Phenylglyoxylic Acid	4

*Documentation of the Threshold Limit Values and Biological Exposure Indices, American Conference of Governmental Industrial Hygienists (ACGIH), Sources of Variability in Biological Monitoring, pp. 5-6, 2001.

Classification of Elimination Half Lives**	Hours	
Short Half Life	X< 2	
<u>Medium Half</u> <u>Life</u>	<u>2<x< 10<="" u=""></x<></u>	
Long Half Live	10< X< 100	

**V. Fiserova, Scand. J. Work Environ. Health 11, 7-21 (1985).

Benefits of the Individual Worker Baseline

- To know if the worker had nonoccupational recent exposure to VOCs.
 - Active and passive smoking.
 - General environmental pollution.
 - Other sources of exposure, such as personal or family jobs or workshops.

To integrate personal variables and characteristics:

- Age,
- Weight,
- Diet,
- Medication,
- Metabolism (genotype and phenotype)

The Urinary Exposure Sample (UES) collected at the end of shift in relation to the Urinary Control Sample (UCS) collected prior to shift

• To measure the <u>exposure gained during the work</u> day, it is highly reccomended, to take into account both: UES and UCS

Urinary Exposure Sample – Urinary Control Sample

-

Exposure gained during the work day

First conclusion:

 <u>Urinary Control Sample</u> of the worker himself can be considered as the <u>Gold Standard</u> for <u>Biomonitoring of Occupational Exposure to</u> <u>Volatile Organic Compounds.</u>



Questionnaire to Identify Confounding Factors

Basic Questions about:

- Active and Passive Smoking

- **Diet** (Red Meat): Amino acids with benzene rings in their molecular structure: **Phenylalanine**,**Tryptophan**, **Tyrosine**.

<u>Medication</u>: We have found the following drugs or medicines ingested or administered to the workers with benzene rings in their molecular structure: Ampicillin, Amoxycillin, Benzetacyl, Bi-euglucon, Chlorpropamide, Diclofenac, Norvase, Pantoprazol and Paracetamol.

Questionnaire of Industrial Hygiene

Basic Questions about:

- <u>Types of Chemicals Used.</u>
- <u>Type of Activity</u> (painting, metal cleaning, equipment maintenance).
- <u>Use of the Personal Protective Equipment:</u>
 First, if they have or not. Second, if it is used or not. Third: Types of Respirators and Gloves Used.
- If the <u>activity has been made in an open or closed space.</u>

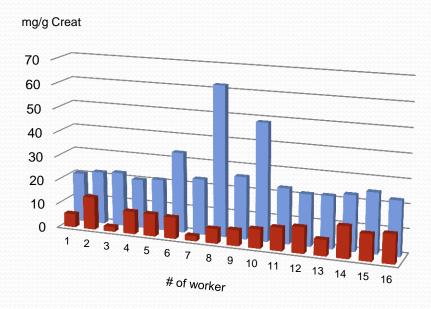


- Identification of False Positive.
- Identification of True Positive.
- Prevention and Control of Chemical Exposure through the Biological Monitoring Follow up.

True Positive Benzene and Toluene Results

True Positive Benzene Results

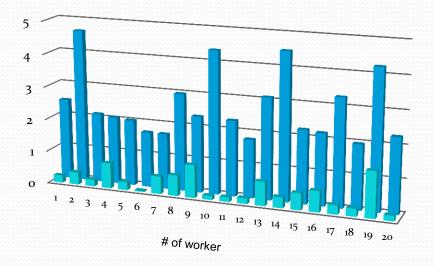
True Positive Toluene Results



Control Urine Phenols

Exposure Urine Phenols

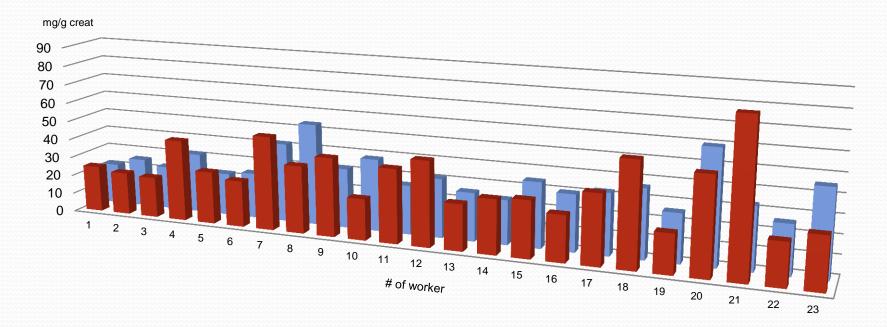
g/g Creat



Control Urine Hippuric AcidExposure Urine Hippuric Acid

False Positive Benzene Results

False Positive Benzene



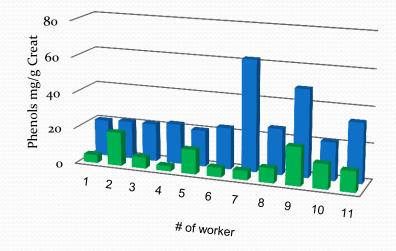
Control Urine Phenols Exposure Urine Phenols

Follow-up Biomonitoring Results of Workers

Found Overexposed

Follow-up Biological Monitoring of Benzene

Follow-up Biological Monitoring of Toluene



5.0000 Hippuric Acid g/g Creat 4.0000 3.0000 2.0000 1.0000 0.0000 1 2 3 4 5 6 7 8 9 10 # of worker

- Follow-up Biological Monitoring Average Results
- Annual Biological Monitoring

- Follow-up Biological Monitoring Average Results
- Annual Biological Monitoring

Final Conclusions:

1. We consider useful and suitable the proposed Model because allows us:

a) <u>To identify True Positive</u>: workers with exposure above the BEIs or Adopted Reference Values, overexposed.

b) <u>To identify False Positive</u>: workers with nonoccupational exposure, with high concentrations prior to shift.

c) <u>To verify the effectiveness of the Industrial Hygiene measures through the</u> <u>Follow-up Biomonitoring</u>.

2.- <u>There is a need to prove this Model with more specific and sensitive biomarkers</u> <u>of exposure.</u>

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