# TEACHING INTERMOLECULAR FORCES USING FORENSIC CHEMISTRY REBEKAH WILKINS, HONORS PROGRAM – MIDAMERICA NAZARENE UNIVERSITY, OLATHE KS



# Drug Sample from Crime Scene

## Abstract

The aim of my research was to find an effective way to teach Chemistry students about intermolecular forces. Forensic Chemistry is a growing field that requires instrumental analysis and the most common instrument used in a crime lab is the Gas Chromatography-Mass Spectrometer (GC-MS). I designed a lab that uses a Forensic scenario to teach about the effect of different intermolecular forces on GC-MS results. This lab was modeled after the method of drug identification used in a crime lab, outlined above. It requires students to integrate basic chemical principles and instrumental methods, all inside the context of Forensic Chemistry.

## Research Methods

# Step 1: Find appropriate compounds

The Forensic scenario calls on the students to correctly identify a drug and two adulterants from a single sample. needed to find compounds that would show different intermolecular forces clearly.

# Step 2: Adjust concentration ratio

The concentration of each compound is reflected in the height of the peaks that come out on the GC-MS spectrum. I had to find the concentration ratio that resulted in three peaks of approximately equal heights.

# **Step 3: Create temperature**

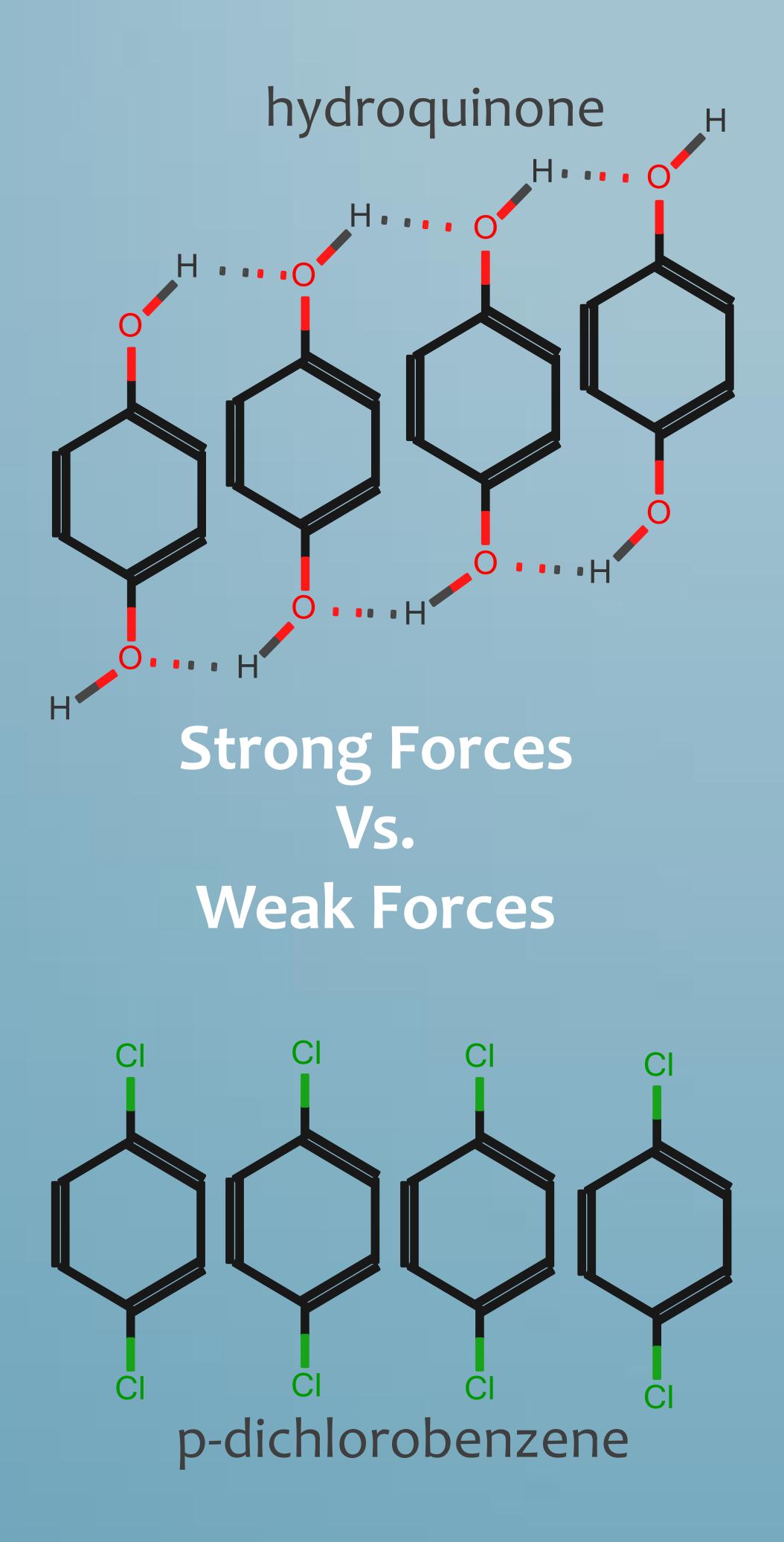
### program

The GC-MS separates compounds by running a sample (mobile phase) through a heated column (stationary phase). Each compound will come out at a different time, called a retention time. I needed to create a temperature program that resulted in three separated peaks.

# Method of Drug Analysis Used by Forensic Chemists

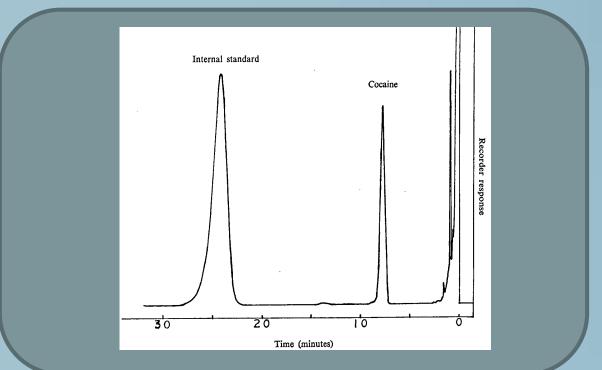


Data Acquisition/Instrumentation



## Conclusion

Through this lab, students will learn about intermolecular forces and have a chance to test their knowledge in a real-life scenario. The parameters I designed can be tailored to fit a General Chemistry, Organic Chemistry, or Intro to Chemistry class.

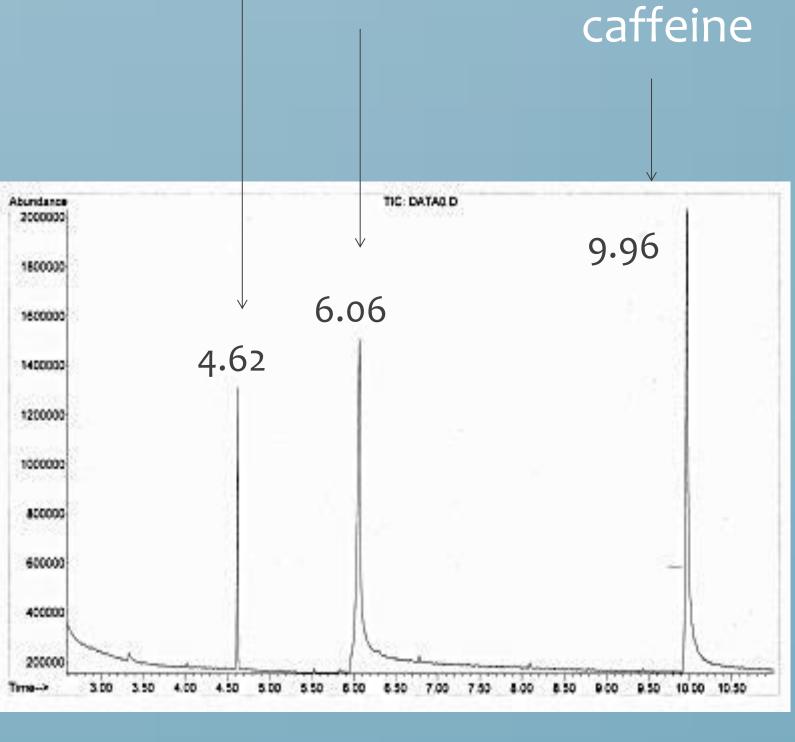


# Data Analysis

#### Table of Compounds Used

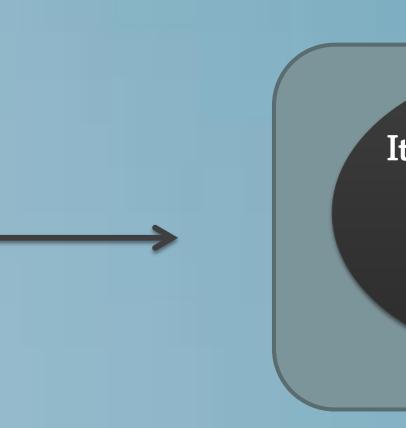
| Name              | Structure | <b>Boiling Point</b> | Molecular Weight |
|-------------------|-----------|----------------------|------------------|
| Caffeine          | N N       | 178                  | 194              |
| Hydroquinone      | но-{      | 287                  | 110              |
| p-Dichlorobenzene | CI        | 174                  | 147              |

### p-dichlorobenzene hydroquinone



# Strength of Forces

The GC-MS separates compounds based on, among other factors, the strength of their intermolecular forces.



## Interpretation/Conclusions

# Step 1

I chose caffeine as the mock drug because it is a standard chemical used in GC-MS. After comparing many different compounds, I found that hydroquinone and p-dichlorobenzene were exactly what I needed. They are similar in molecular weight and symmetry, but have different intermolecular forces. So students can compare the forces in a controlled manner.

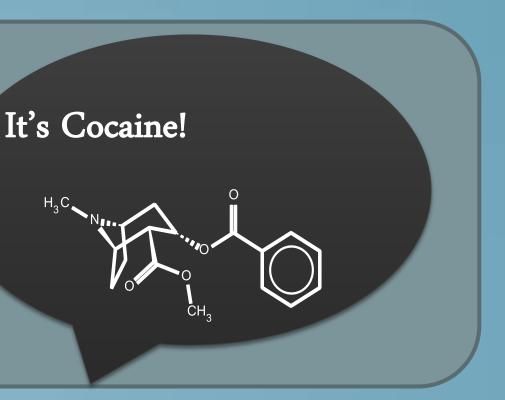
## Step 2

# Step 3

In creating a temperature program, there needs to be balance between good peak separation and a short run. The method below has both of those qualities.

| Step  | Rate (°C/min) | Temperature (°C) | Hold (min) |
|-------|---------------|------------------|------------|
| Start |               | 50               | 2.5        |
| 1     | 40            | 150              | 1          |
| 2     | 20            | 200              | 1          |
| 3     | 60            | 230              | 1          |

Bell, S. (2013). Forensic Chemistry. United States of America: Pearson Education, Inc. Smith, J. G. (2011). Organic Chemistry. New York: McGraw-Hill.



## Results

### My main problem with concentration was that pdichlorobenzene because the predominate compound once it was in the column, reducing the other two peaks to small bumps. I finally found that the correct ratio is: 0.05 g p-dichlorobenzene, 0.5 g caffeine, and 1 g hydroquinone. The final spectrum is to the left.

Viai IIIIIC – II IIIIIIuic.

## Bibliography