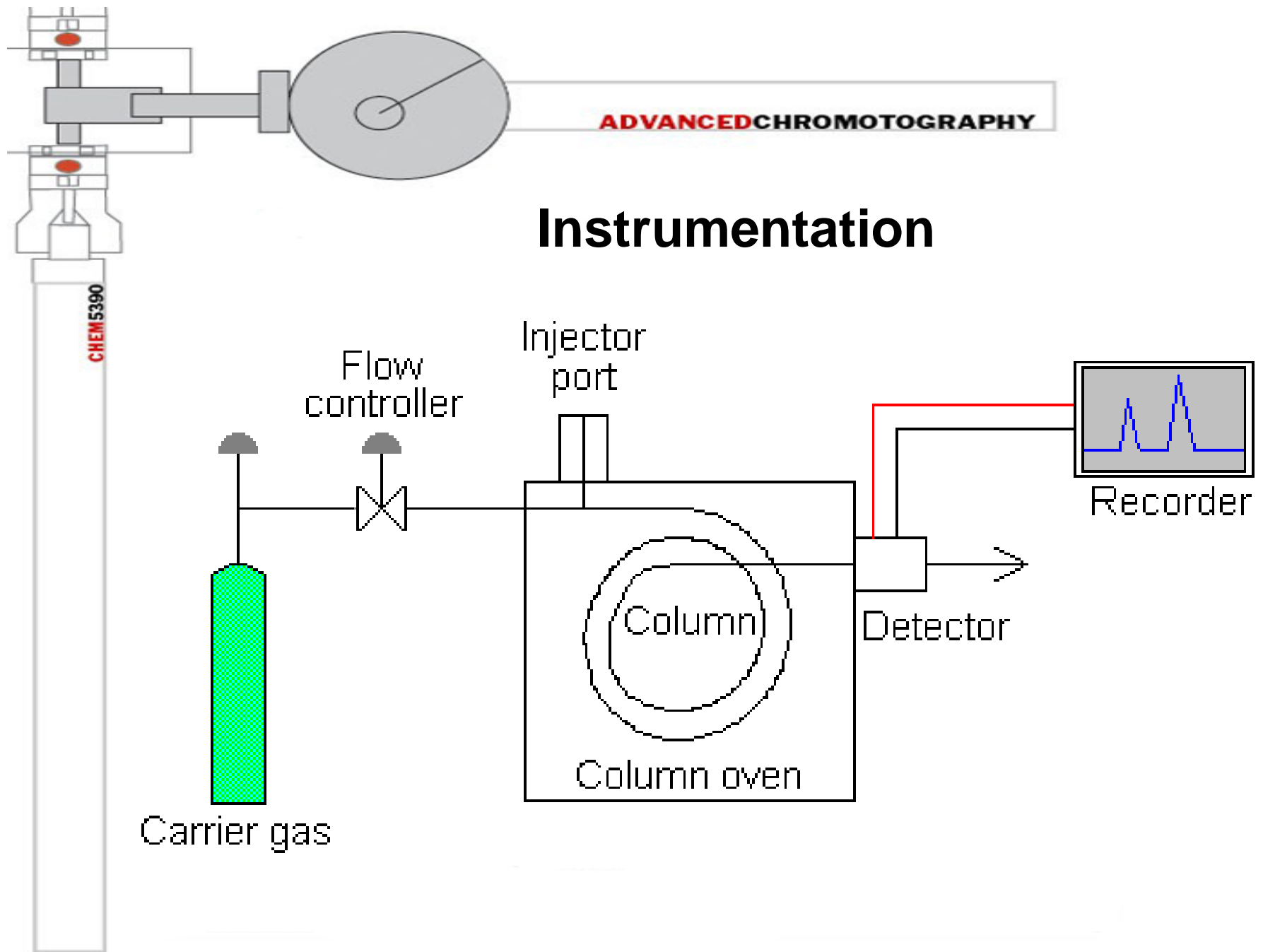
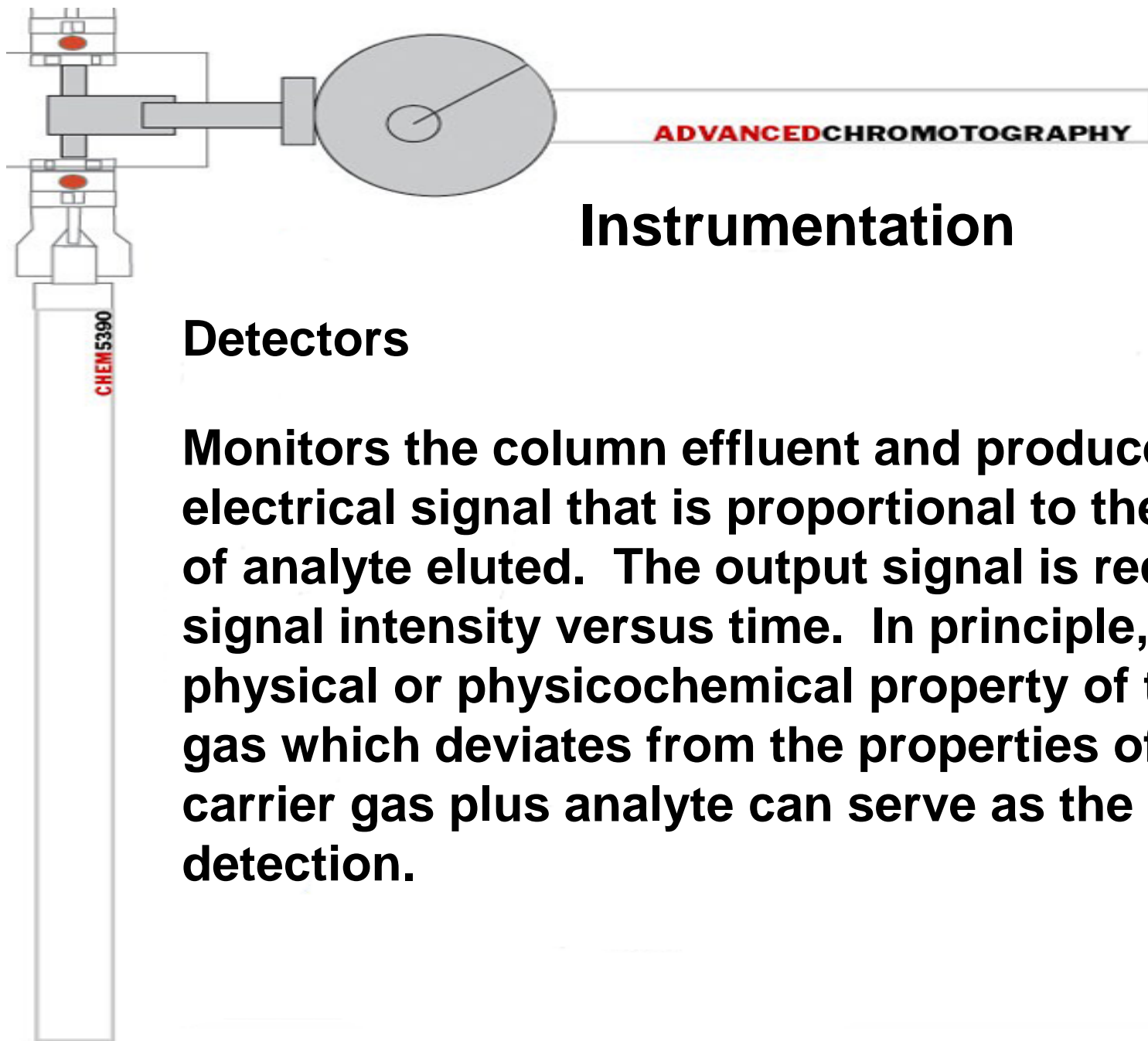


# Lecture 8: Gas Chromatography

## Instrumentation

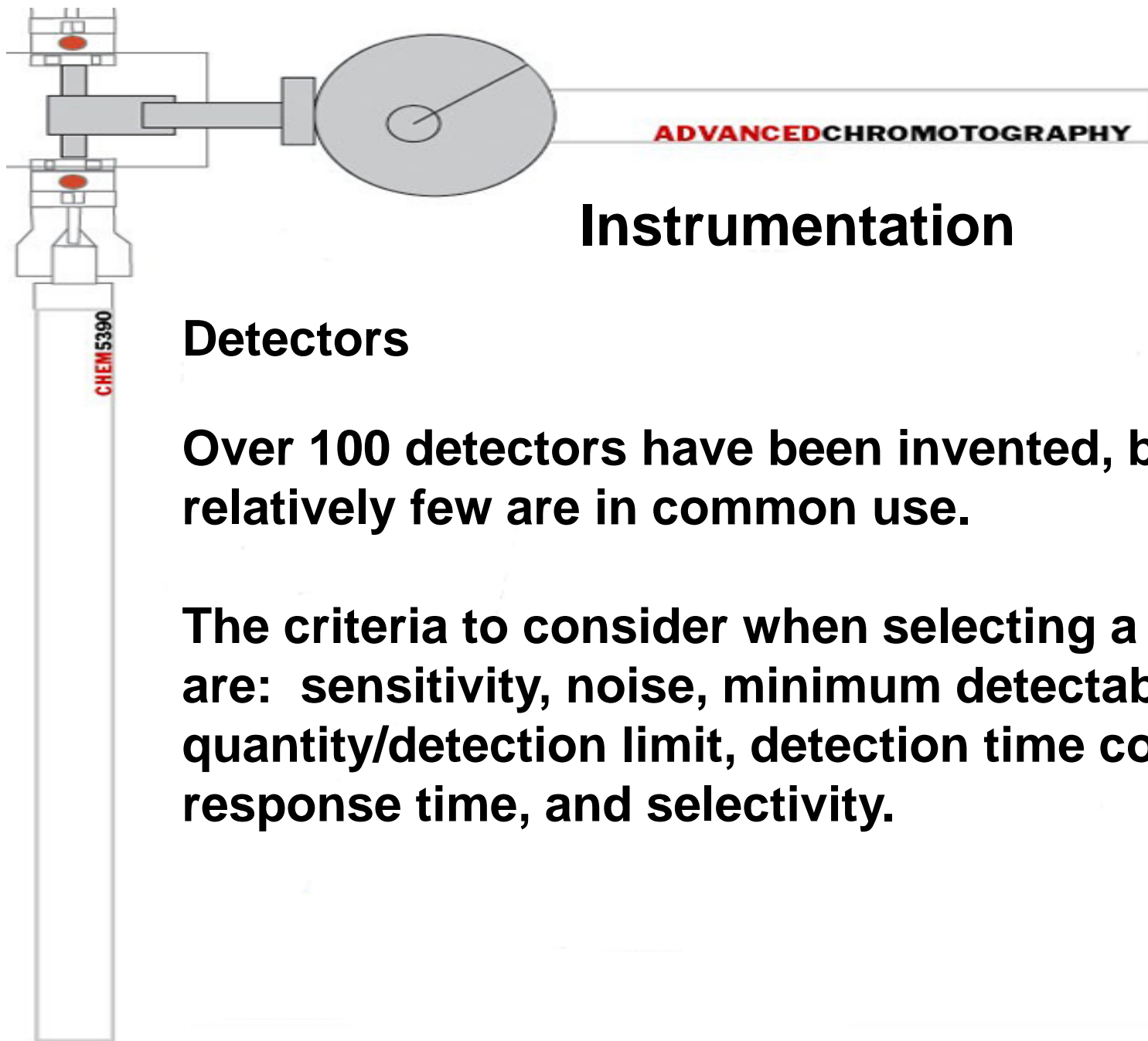




## Instrumentation

### Detectors

**Monitors the column effluent and produces an electrical signal that is proportional to the amount of analyte eluted. The output signal is recorded as signal intensity versus time. In principle, any physical or physicochemical property of the carrier gas which deviates from the properties of the carrier gas plus analyte can serve as the basis of detection.**

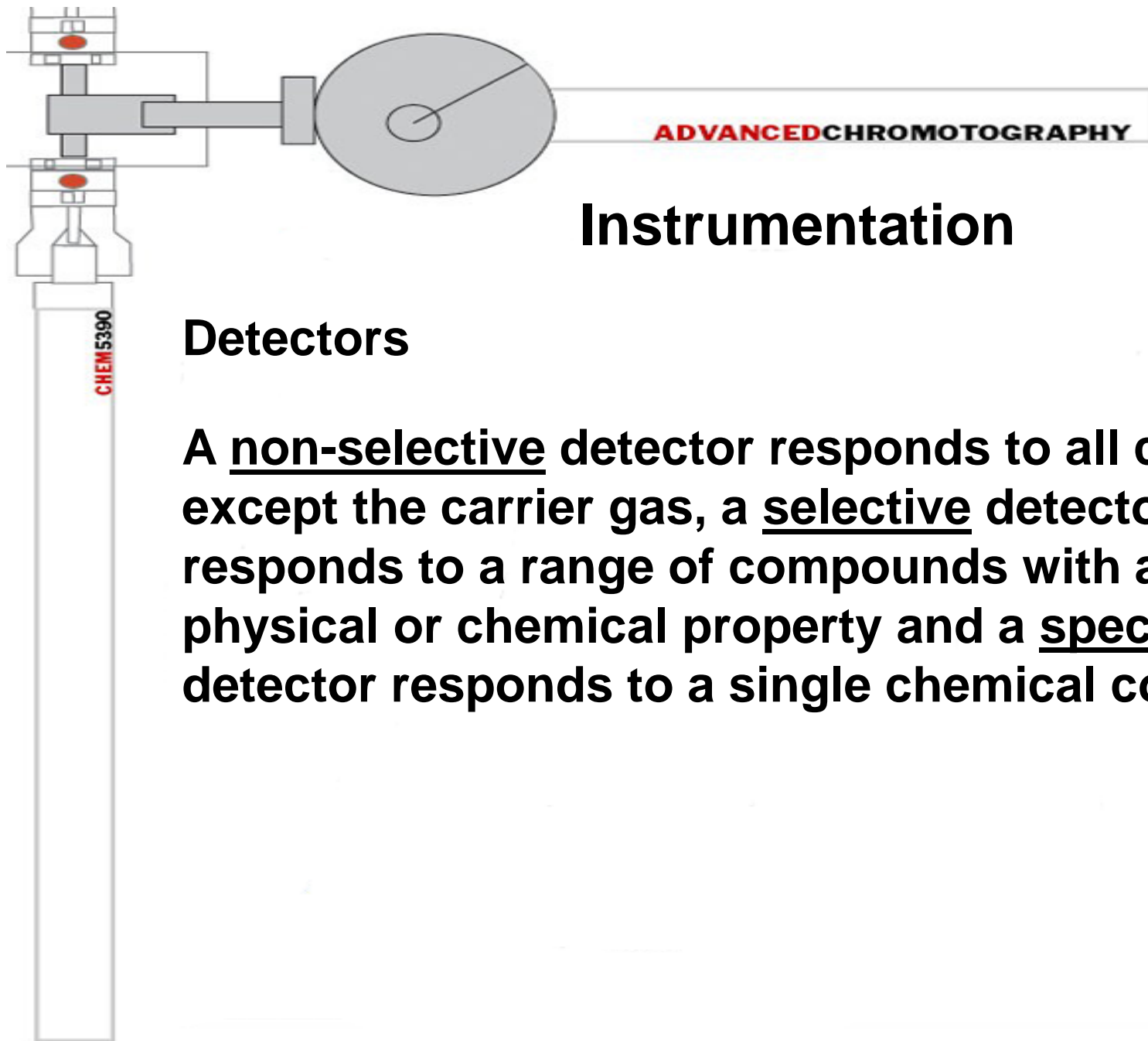


## Instrumentation

### Detectors

**Over 100 detectors have been invented, but relatively few are in common use.**

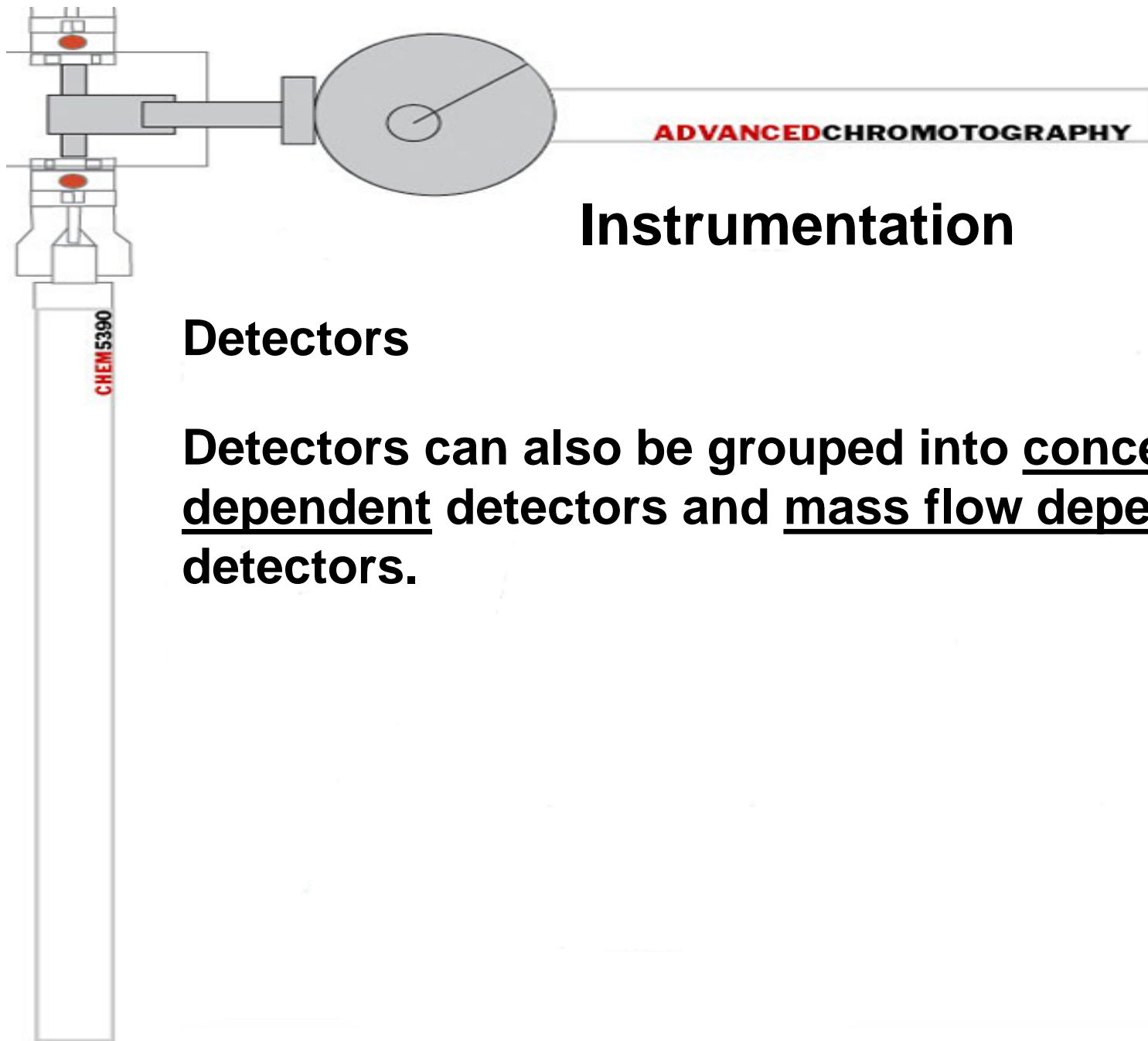
**The criteria to consider when selecting a detector are: sensitivity, noise, minimum detectable quantity/detection limit, detection time constant or response time, and selectivity.**



## Instrumentation

### Detectors

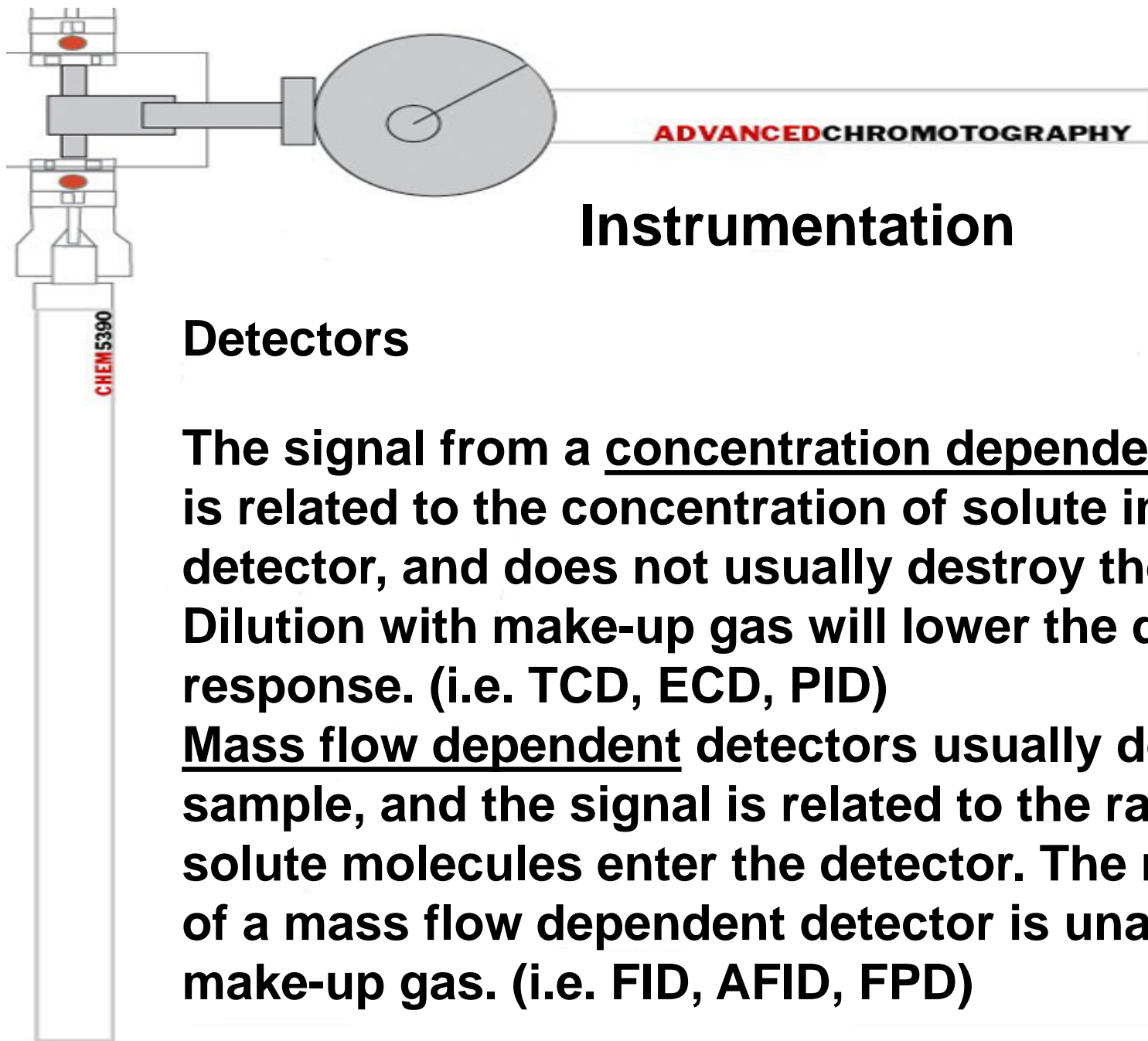
A non-selective detector responds to all compounds except the carrier gas, a selective detector responds to a range of compounds with a common physical or chemical property and a specific detector responds to a single chemical compound.



## Instrumentation

### Detectors

Detectors can also be grouped into concentration dependent detectors and mass flow dependent detectors.

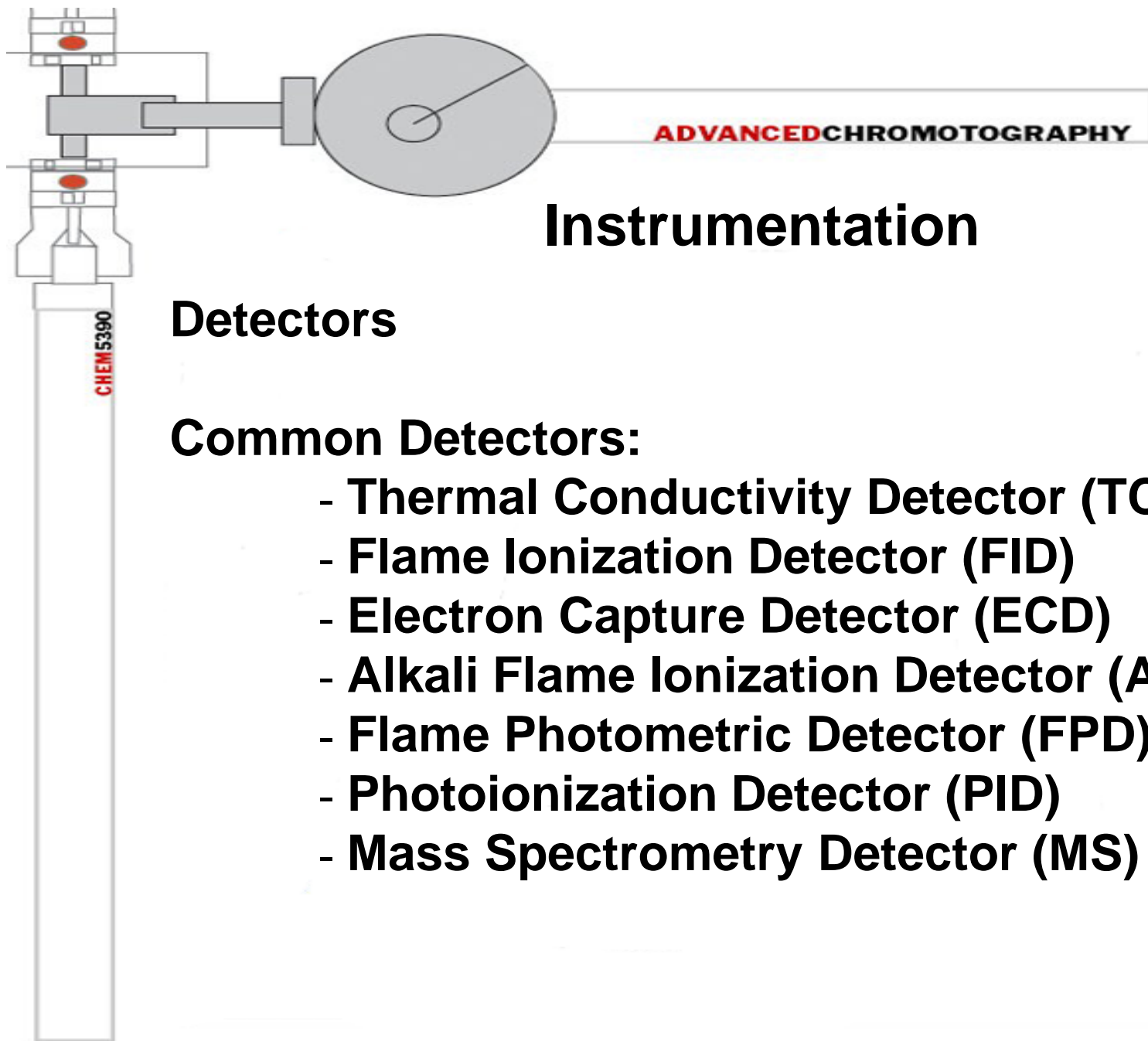


## Instrumentation

### Detectors

The signal from a concentration dependent detector is related to the concentration of solute in the detector, and does not usually destroy the sample. Dilution with make-up gas will lower the detector's response. (i.e. TCD, ECD, PID)

Mass flow dependent detectors usually destroy the sample, and the signal is related to the rate at which solute molecules enter the detector. The response of a mass flow dependent detector is unaffected by make-up gas. (i.e. FID, AFID, FPD)



# Instrumentation

## Detectors

### Common Detectors:

- Thermal Conductivity Detector (TCD)
- Flame Ionization Detector (FID)
- Electron Capture Detector (ECD)
- Alkali Flame Ionization Detector (AFID)
- Flame Photometric Detector (FPD)
- Photoionization Detector (PID)
- Mass Spectrometry Detector (MS)



# Instrumentation

## Detectors

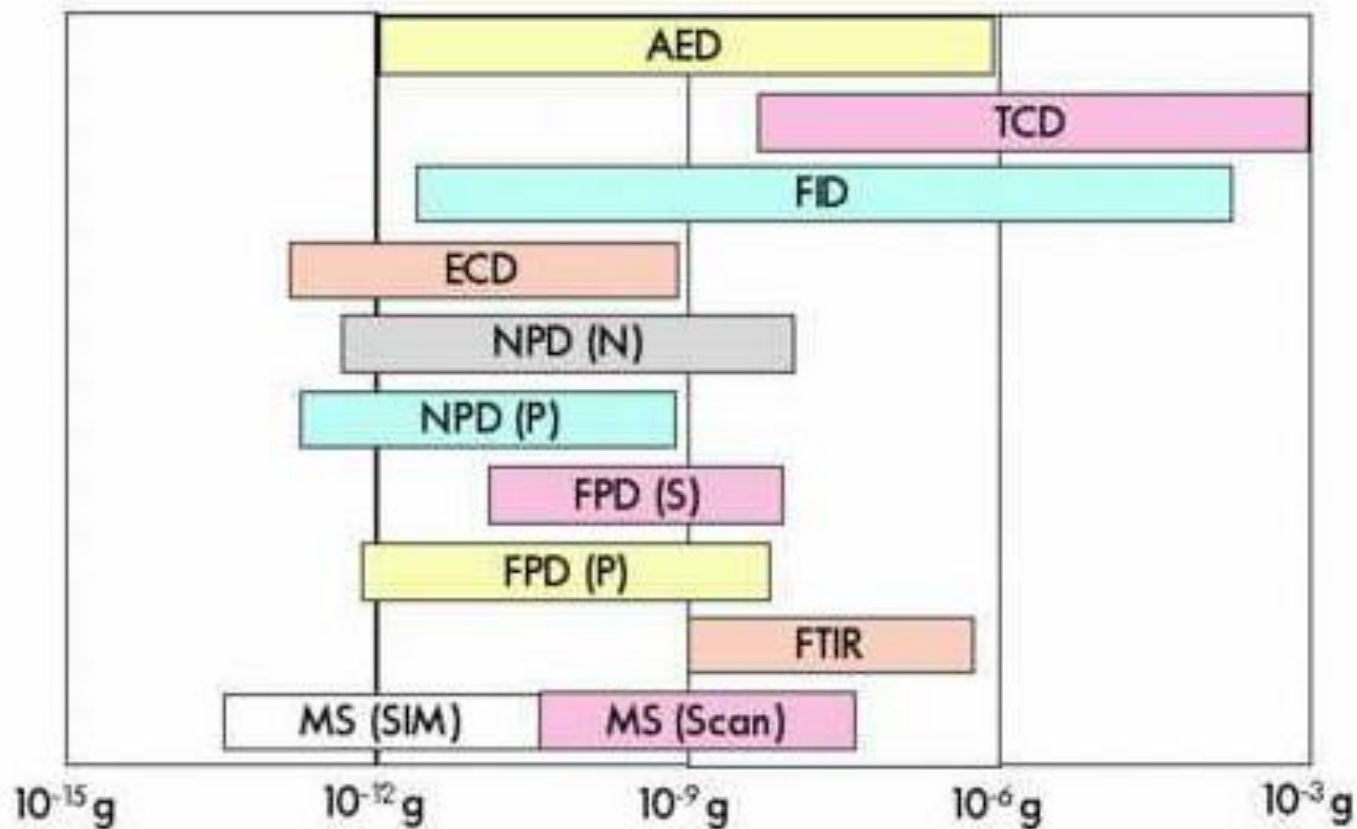
**Table 3.17** Classification of the most common gas chromatographic detectors.

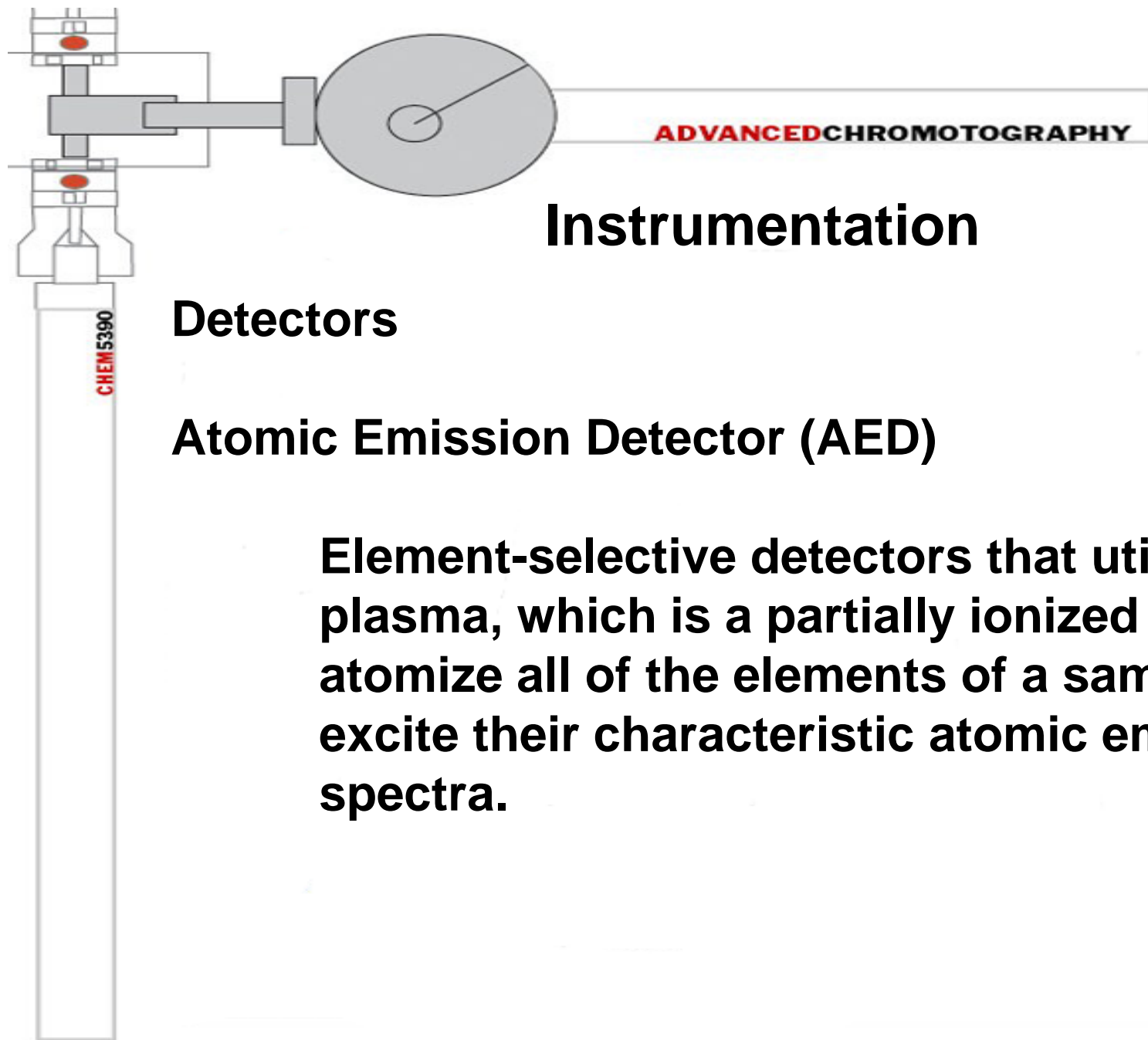
Detector	Response	Optimal detection limit	Linear range	Classification
TCD	Organic and inorganic solutes	$10^{-9} \text{ g ml}^{-1}$	$10^4$	Concentration; nondestructive
FID	All organic solutes except formic acid and formaldehyde	$10^{-12} \text{ g ml}^{-1}$	$10^7$	Mass flow-rate; destructive
ECD	Halogenated and nitro compounds	$10^{-16} \text{ mol ml}^{-1}$	$10^3$ – $10^4$ (pulsed)	Concentration; nondestructive
AFID	P- or N-containing solutes	N: $10^{-14} \text{ g s}^{-1}$ P: $10^{-13} \text{ g s}^{-1}$	$10^3$ – $10^5$	Mass flow-rate; destructive
FPD	P- or S-containing solutes	S: $10^{-10} \text{ g s}^{-1}$ P: $10^{-12} \text{ g s}^{-1}$	S: $10^3$ P: $10^5$	Mass flow-rate; destructive



# Instrumentation

## Detectors



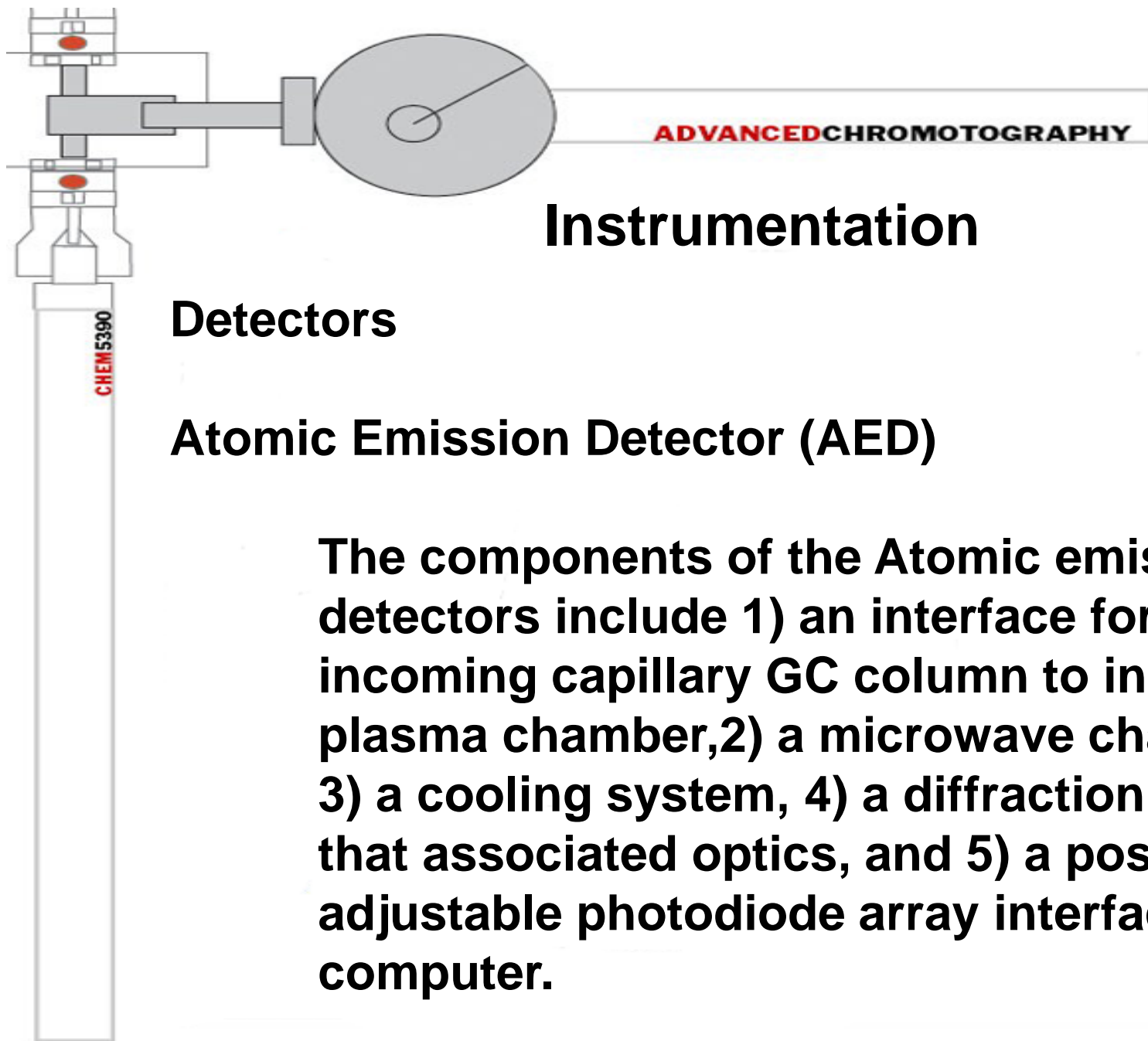


## Instrumentation

### Detectors

#### Atomic Emission Detector (AED)

Element-selective detectors that utilize plasma, which is a partially ionized gas, to atomize all of the elements of a sample and excite their characteristic atomic emission spectra.

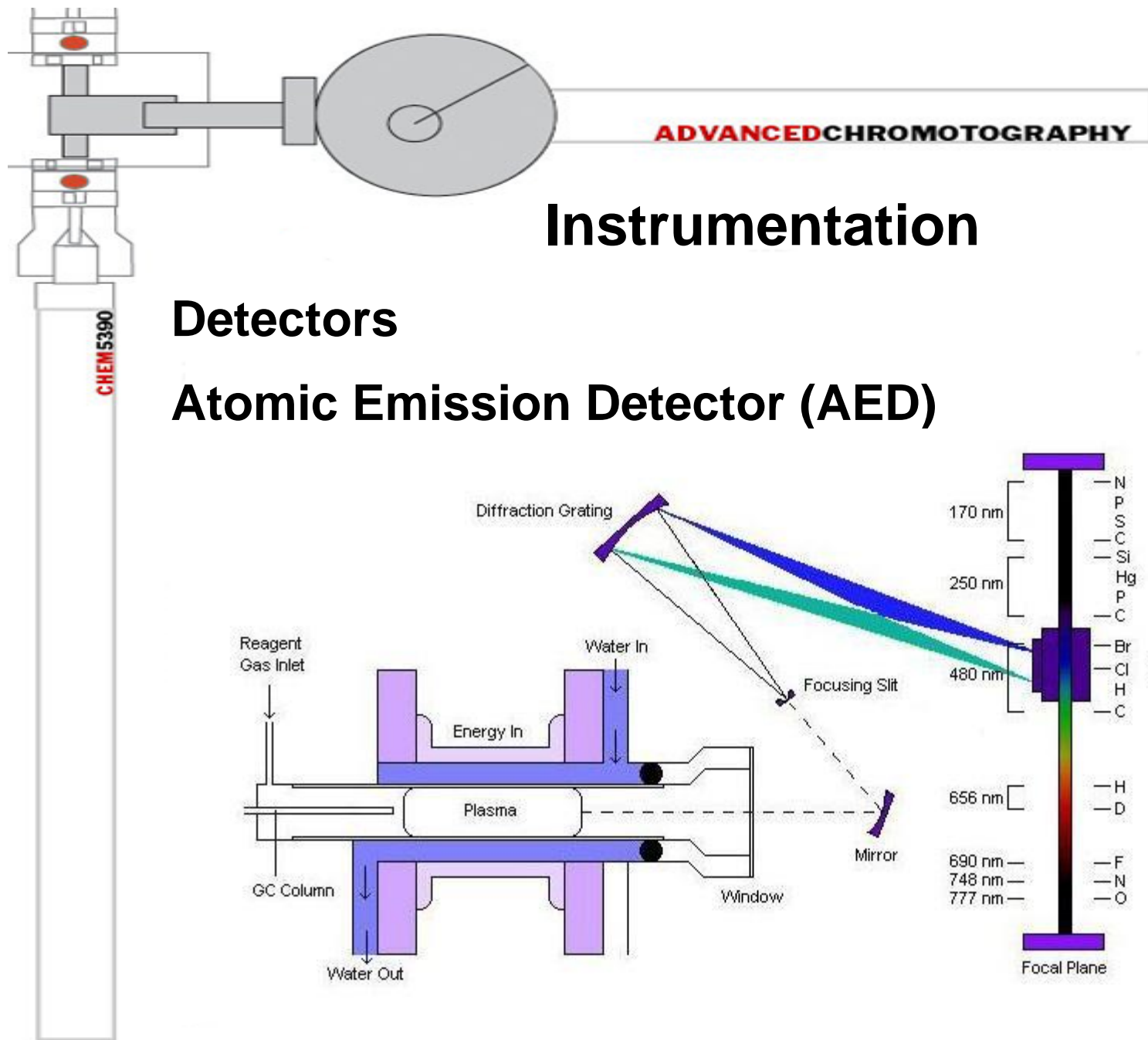


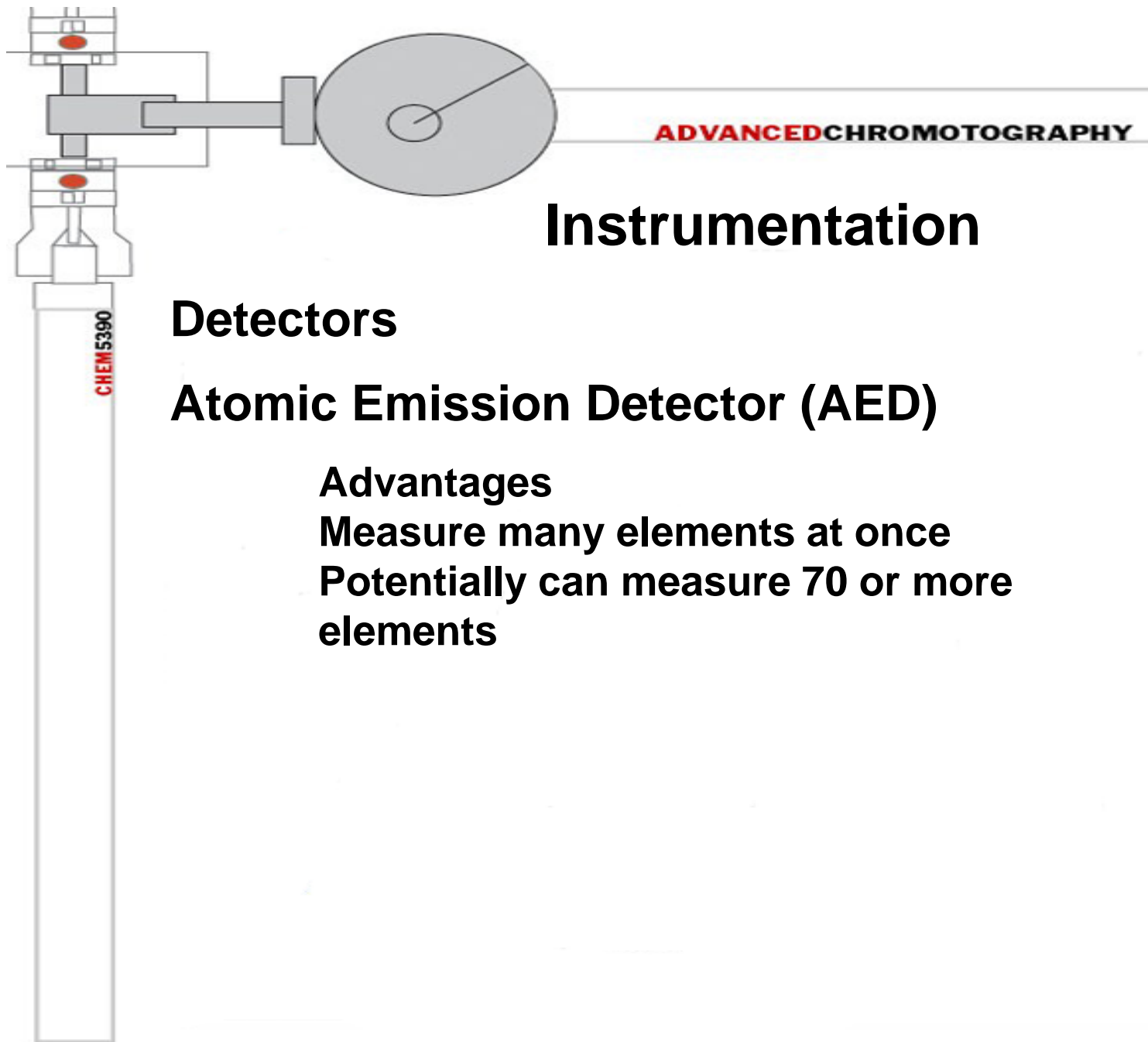
## Instrumentation

### Detectors

#### Atomic Emission Detector (AED)

The components of the Atomic emission detectors include 1) an interface for the incoming capillary GC column to induce plasma chamber, 2) a microwave chamber, 3) a cooling system, 4) a diffraction grating that associated optics, and 5) a position adjustable photodiode array interfaced to a computer.





# Instrumentation

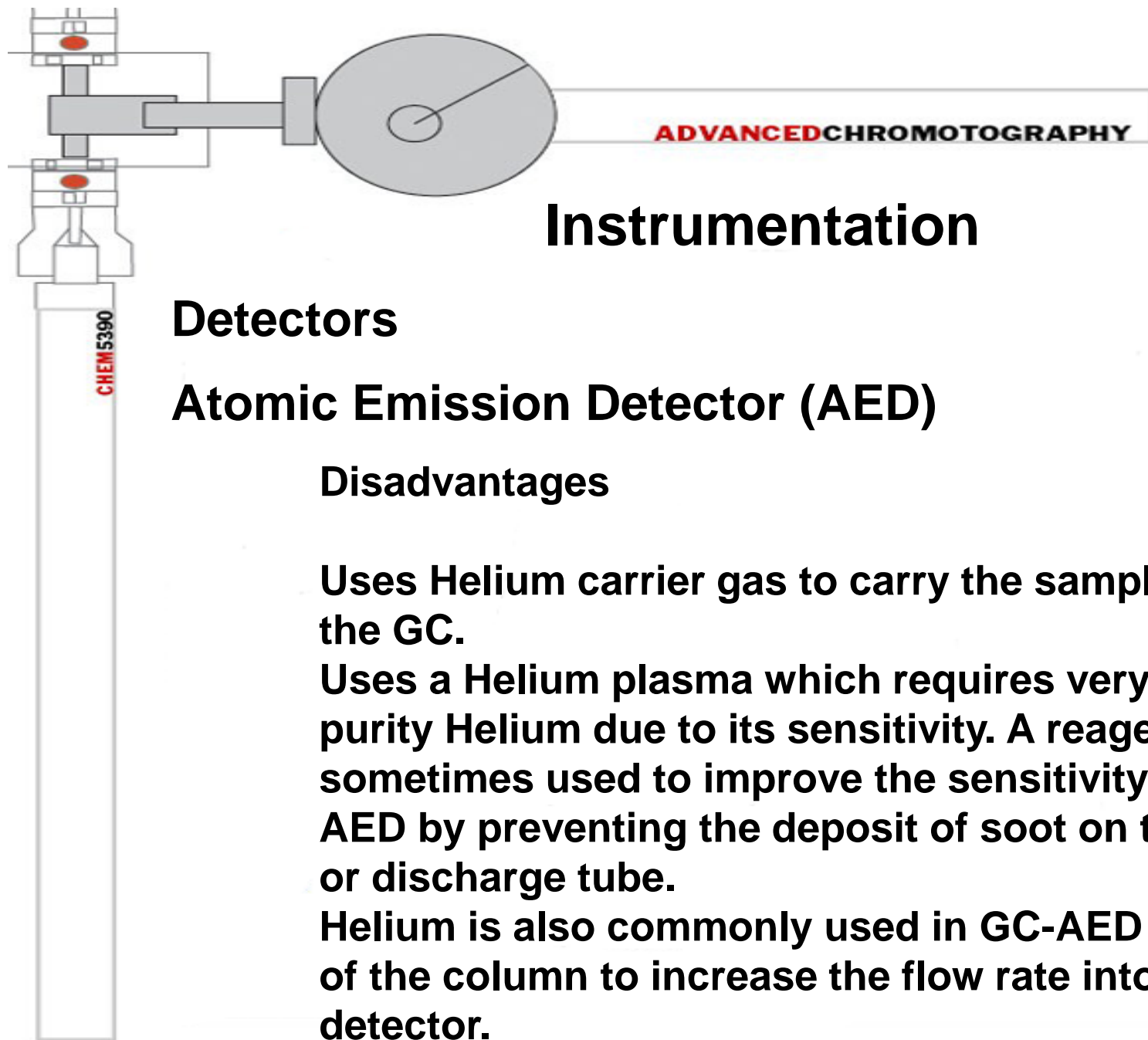
## Detectors

### Atomic Emission Detector (AED)

#### Advantages

Measure many elements at once

Potentially can measure 70 or more elements



## **Detectors**

### **Atomic Emission Detector (AED)**

#### **Disadvantages**

**Uses Helium carrier gas to carry the sample through the GC.**

**Uses a Helium plasma which requires very high purity Helium due to its sensitivity. A reagent gas is sometimes used to improve the sensitivity of the AED by preventing the deposit of soot on the lamp or discharge tube.**

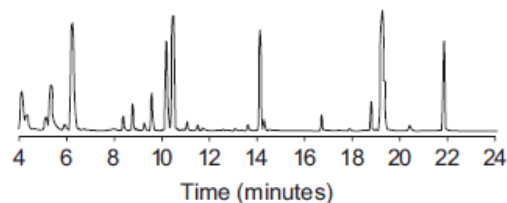
**Helium is also commonly used in GC-AED at the exit of the column to increase the flow rate into the detector.**

## Detectors

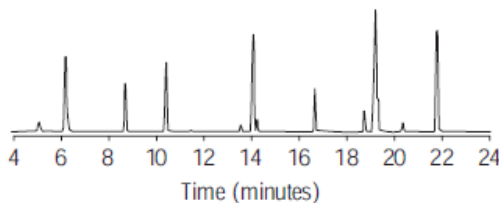
**AED**  
Can  
monitor  
different  
elements.

### Emission Spectra for Element Confirmation

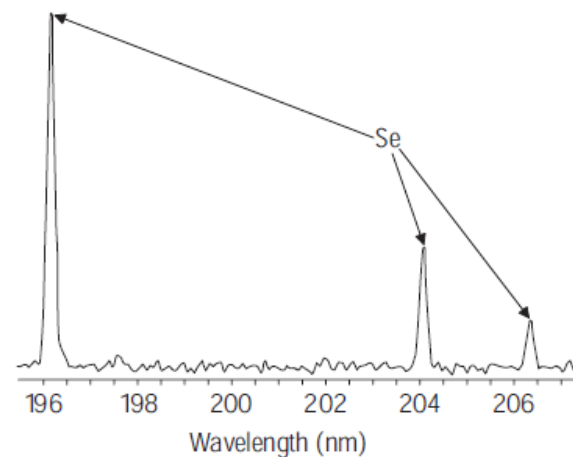
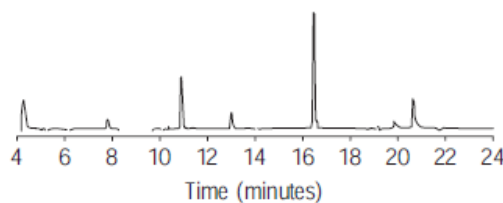
#### Carbon



#### Sulfur

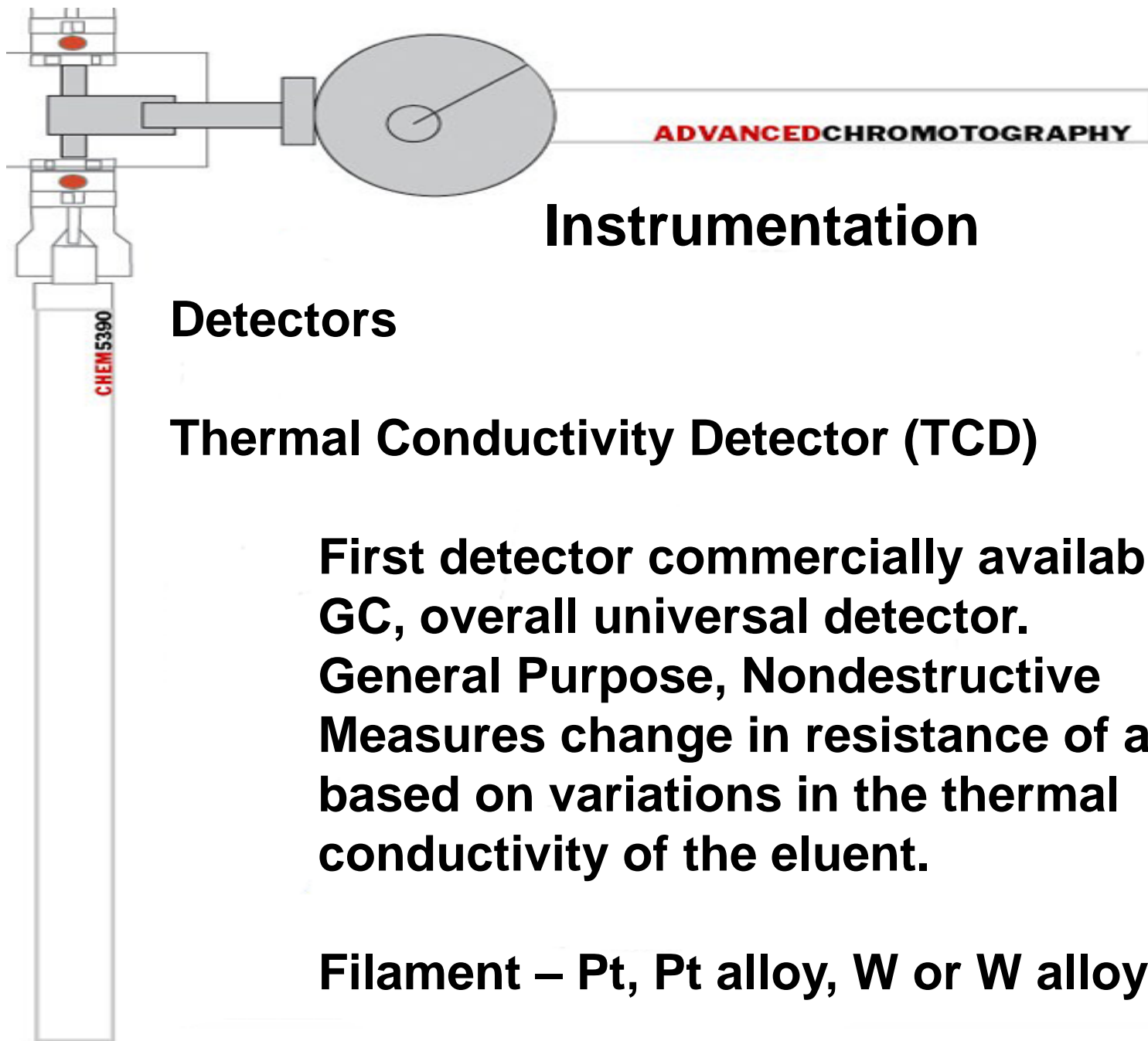


#### Selenium



An extract of garlic shows low-level sulfur and selenium compounds important in flavor and nutritional analyses. Selenium is confirmed by its three characteristic atomic lines from 196 to 206 nm.





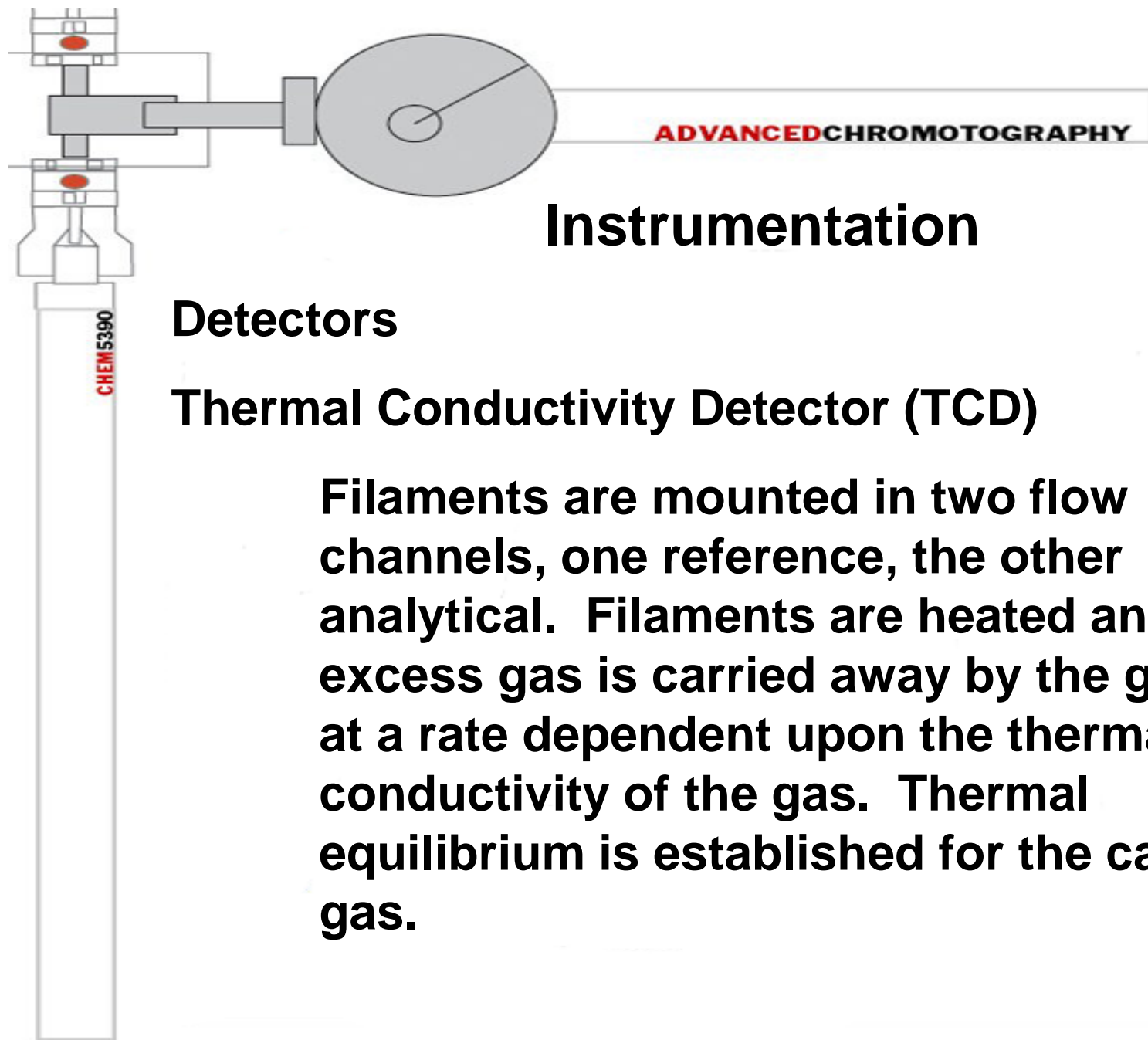
## Instrumentation

### Detectors

#### Thermal Conductivity Detector (TCD)

**First detector commercially available for GC, overall universal detector.  
General Purpose, Nondestructive  
Measures change in resistance of a wire based on variations in the thermal conductivity of the eluent.**

**Filament – Pt, Pt alloy, W or W alloys.**

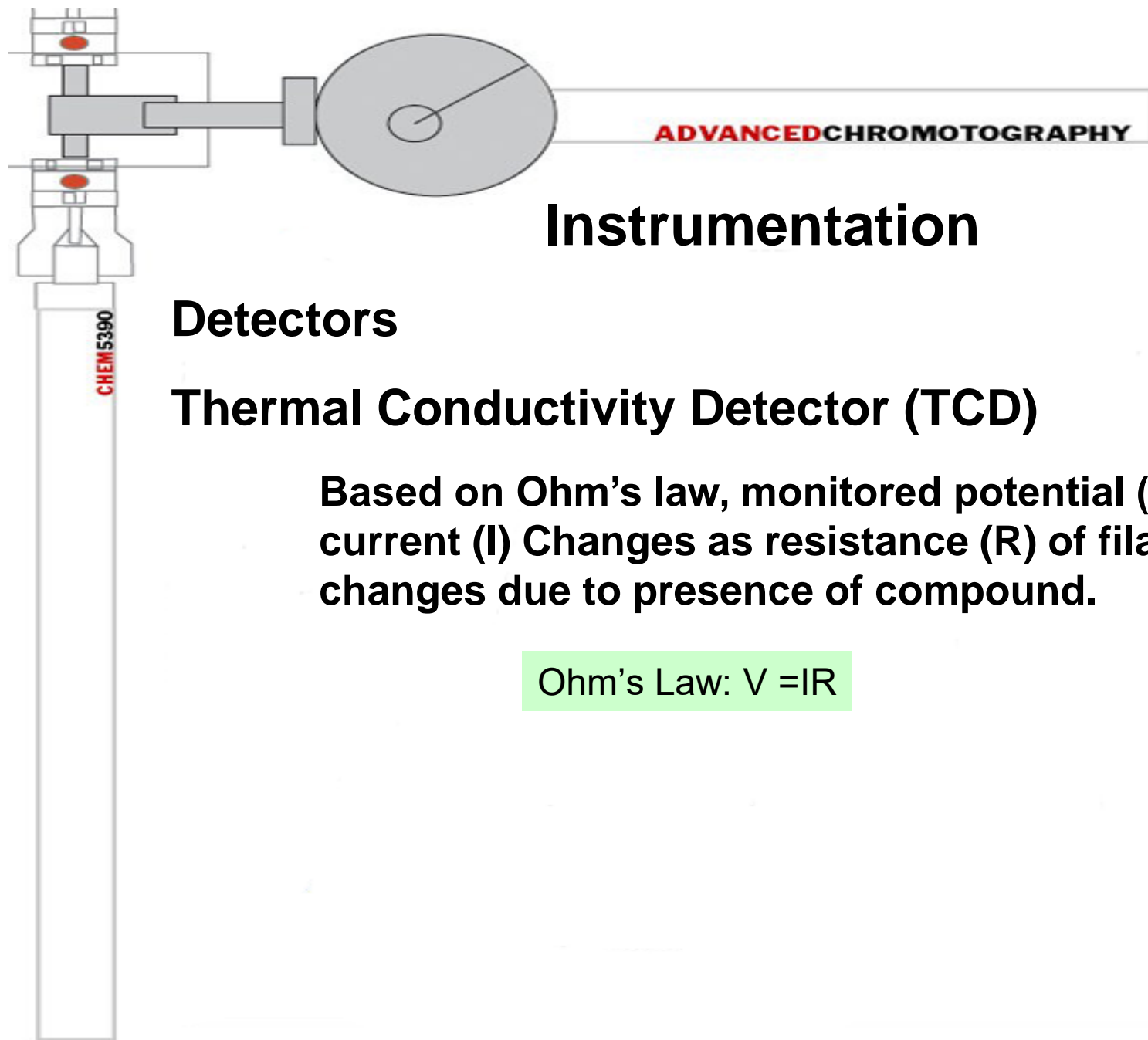


## Instrumentation

### Detectors

#### Thermal Conductivity Detector (TCD)

Filaments are mounted in two flow channels, one reference, the other analytical. Filaments are heated and excess gas is carried away by the gas flow at a rate dependent upon the thermal conductivity of the gas. Thermal equilibrium is established for the carrier gas.



# Instrumentation

## Detectors

### Thermal Conductivity Detector (TCD)

Based on Ohm's law, monitored potential (V) or current (I) Changes as resistance (R) of filament changes due to presence of compound.

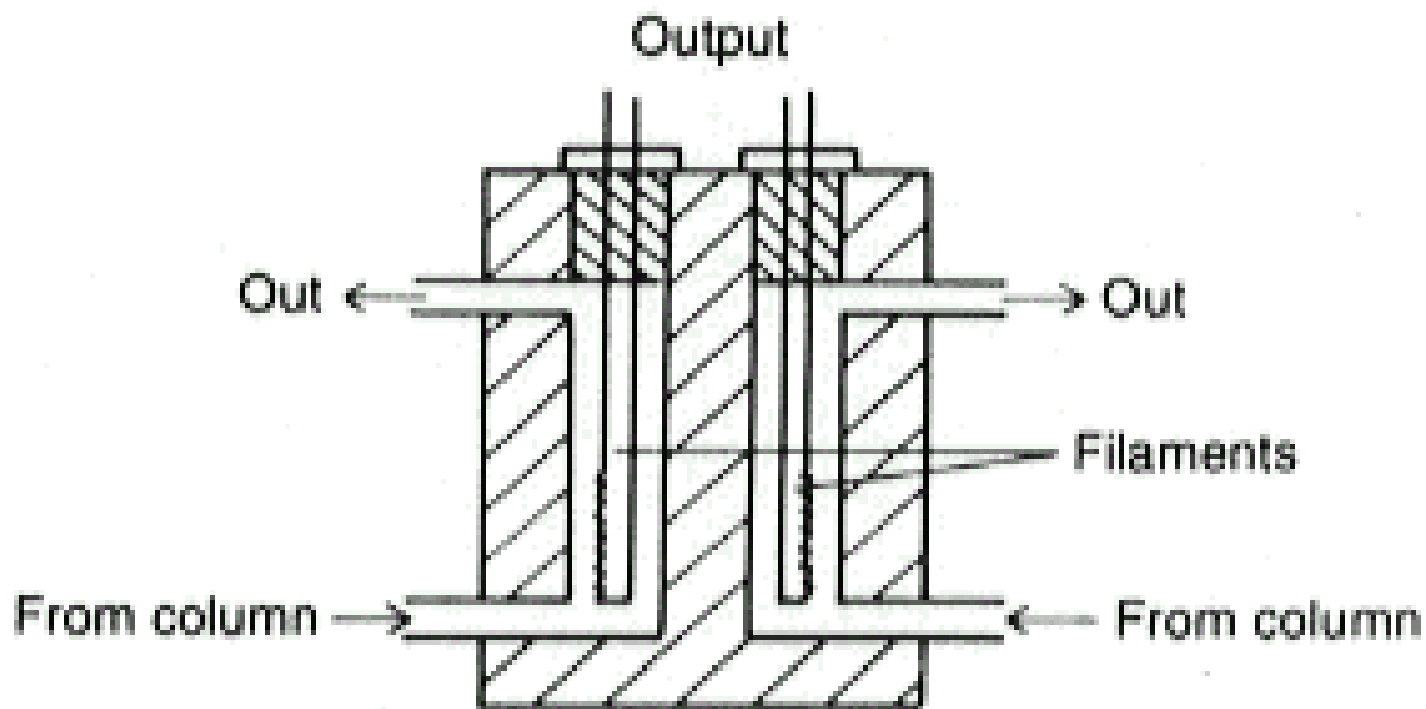
$$\text{Ohm's Law: } V = IR$$

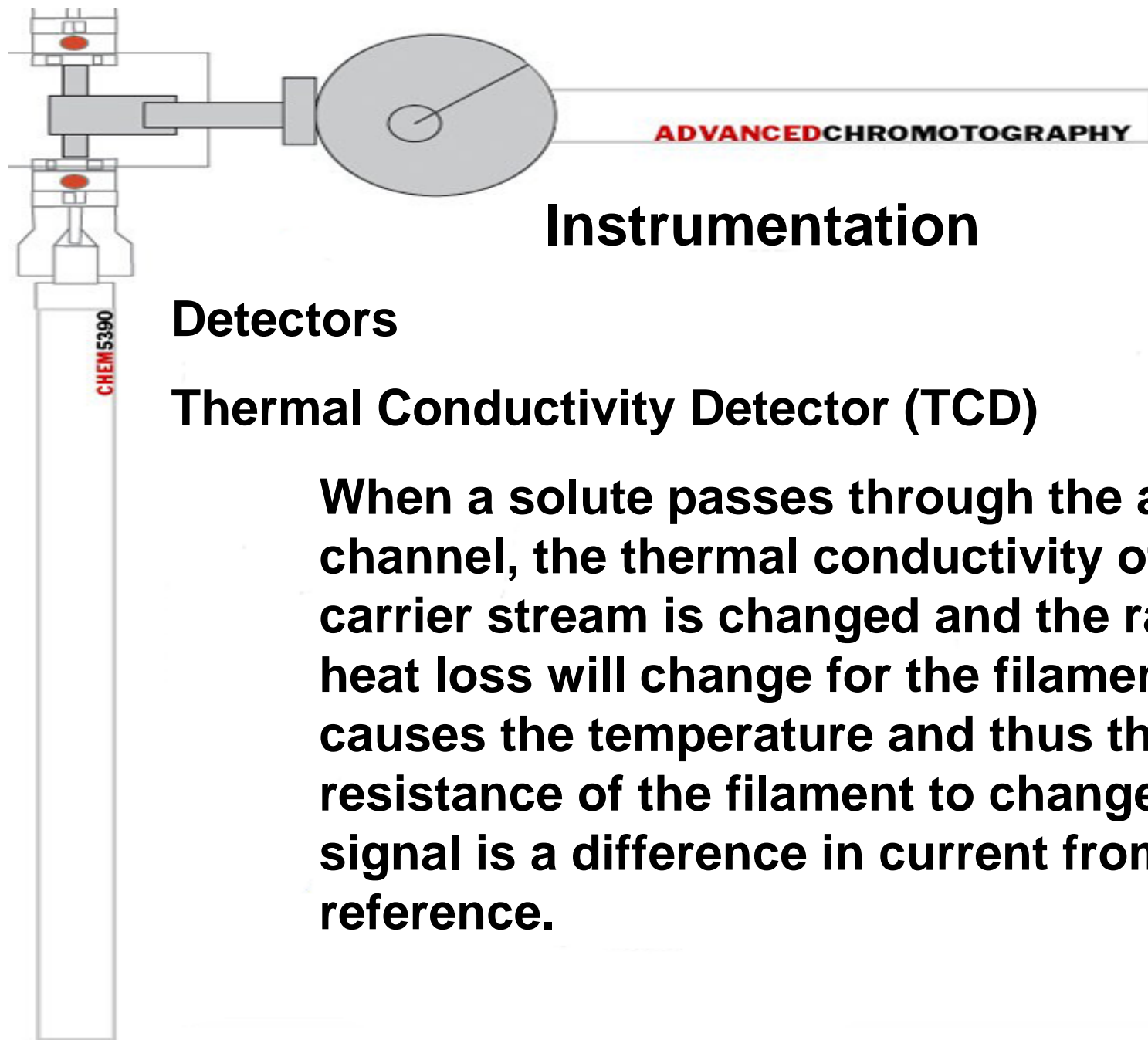


## Instrumentation

### Detectors

### Thermal Conductivity Detector (TCD)



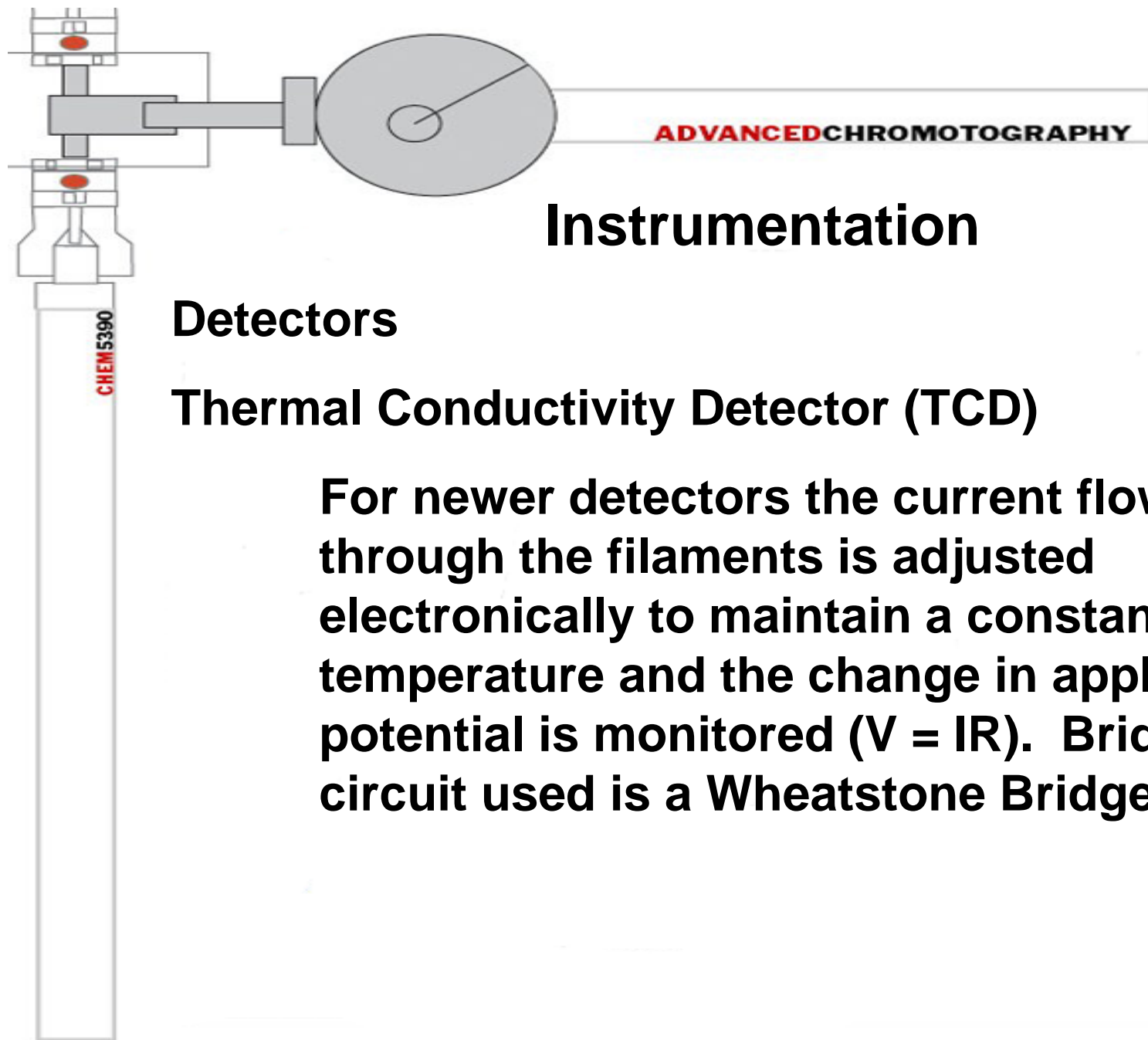


## Instrumentation

### Detectors

#### Thermal Conductivity Detector (TCD)

When a solute passes through the analyte channel, the thermal conductivity of the carrier stream is changed and the rate of heat loss will change for the filament. This causes the temperature and thus the resistance of the filament to change. The signal is a difference in current from the reference.



## Instrumentation

### Detectors

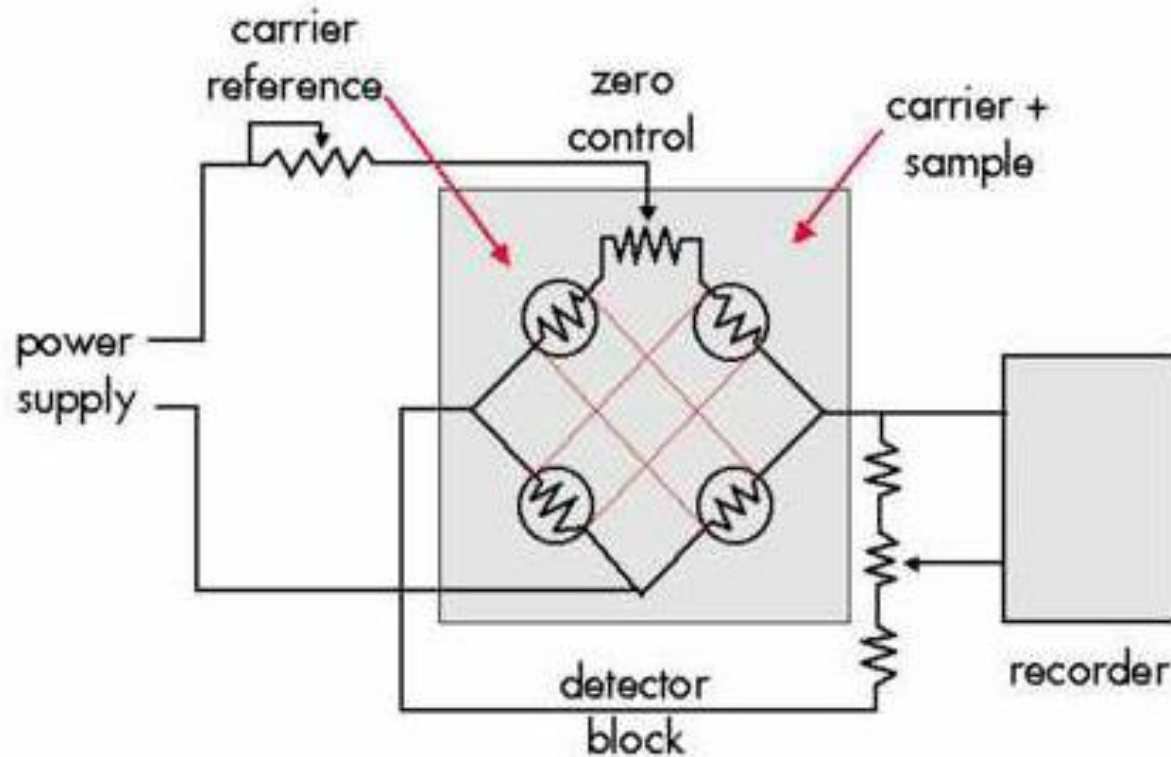
#### Thermal Conductivity Detector (TCD)

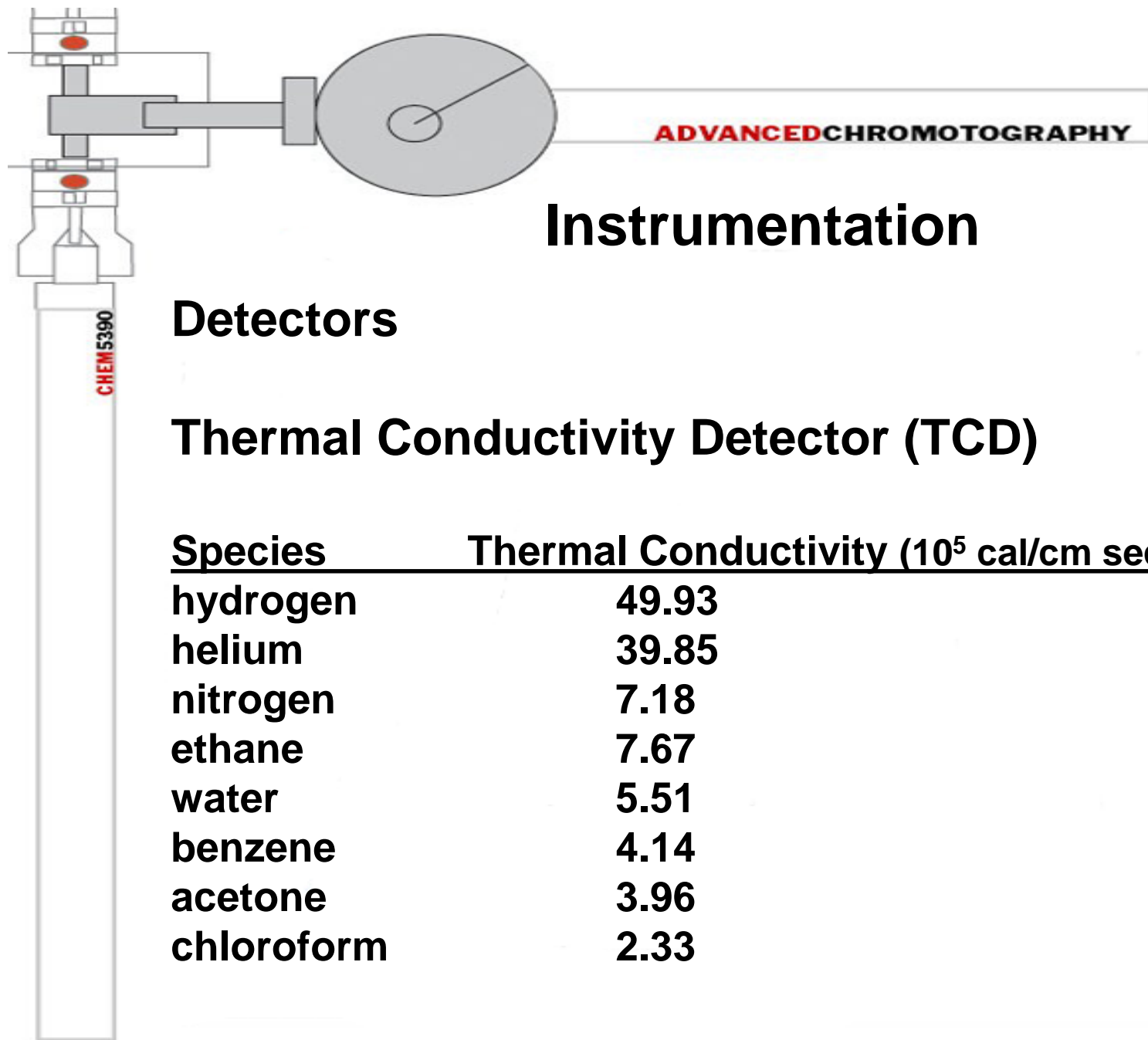
For newer detectors the current flowing through the filaments is adjusted electronically to maintain a constant temperature and the change in applied potential is monitored ( $V = IR$ ). Bridge circuit used is a Wheatstone Bridge.

# Instrumentation

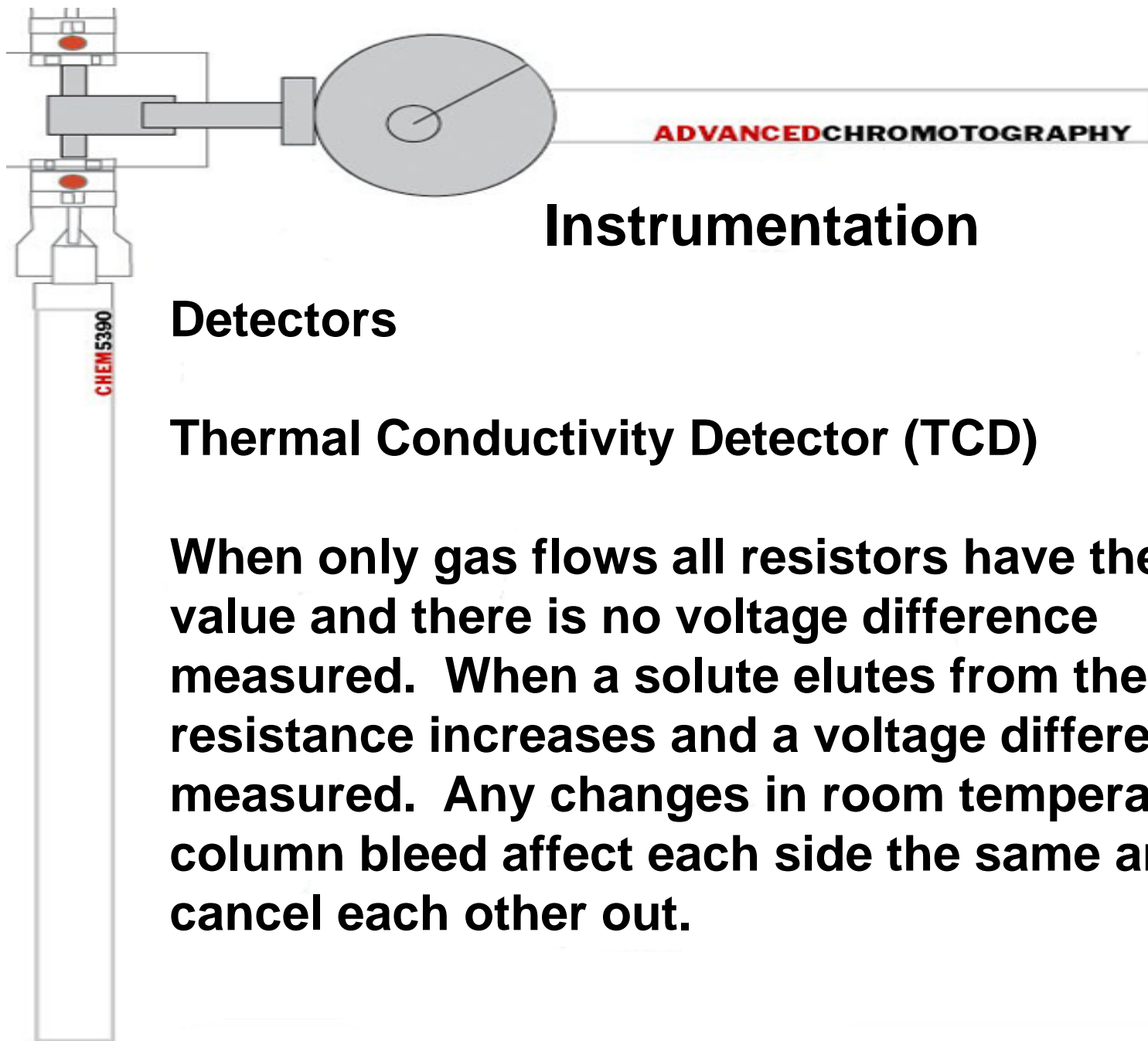
## Detectors

### Thermal Conductivity Detector (TCD)







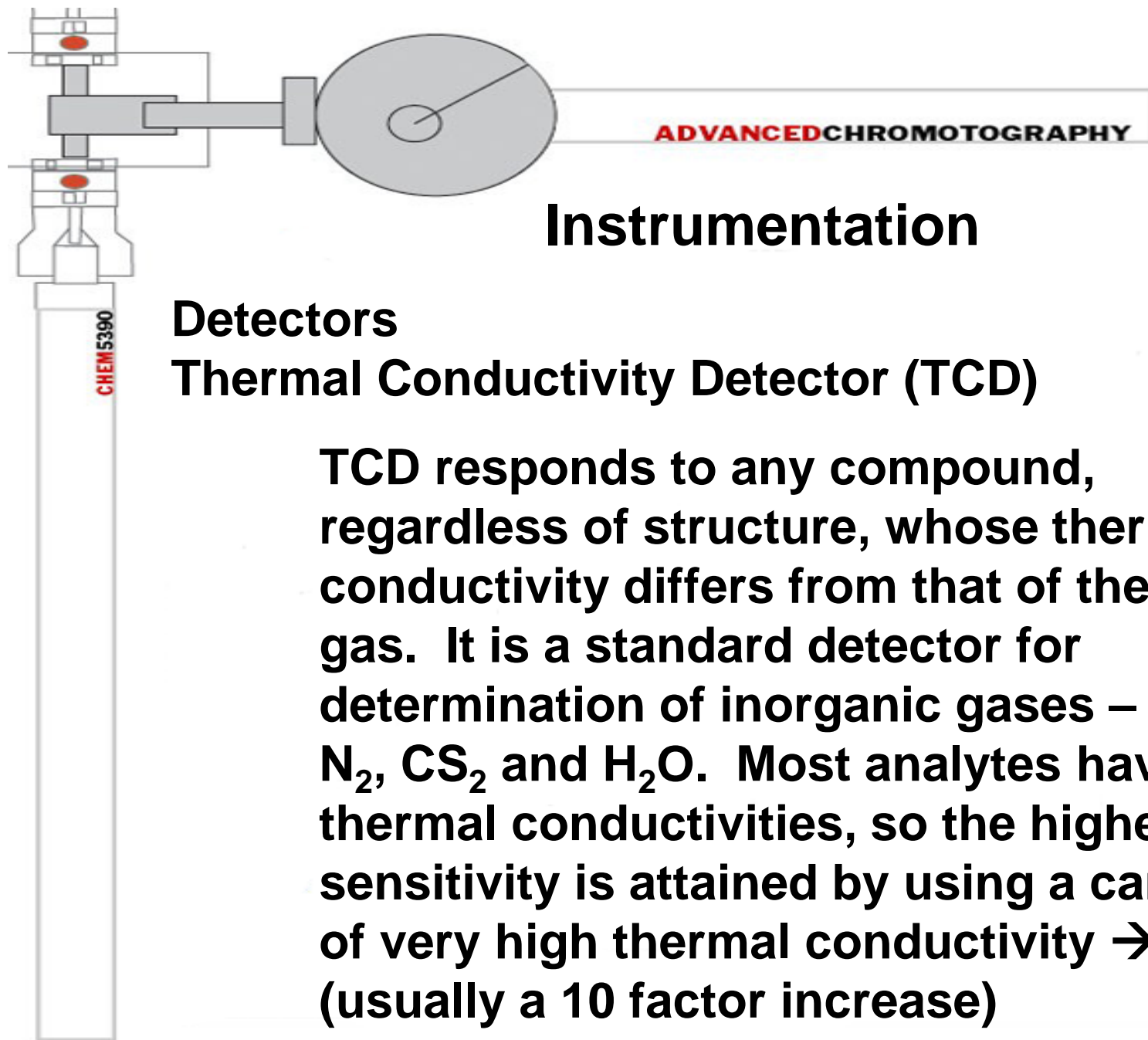


## Instrumentation

### Detectors

#### Thermal Conductivity Detector (TCD)

**When only gas flows all resistors have the same value and there is no voltage difference measured. When a solute elutes from the column, resistance increases and a voltage difference is measured. Any changes in room temperature or column bleed affect each side the same and cancel each other out.**

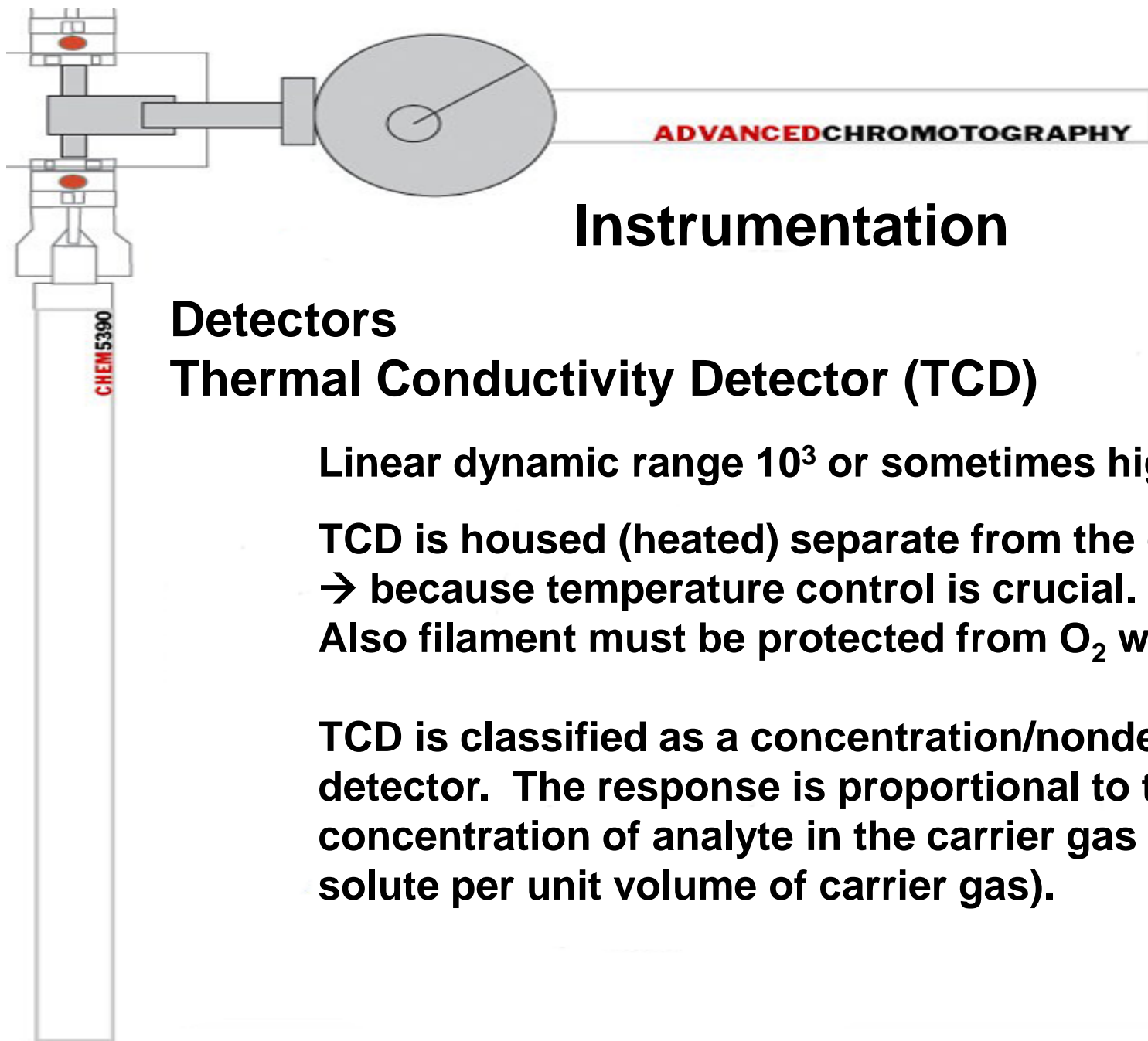


## Instrumentation

### Detectors

#### Thermal Conductivity Detector (TCD)

TCD responds to any compound, regardless of structure, whose thermal conductivity differs from that of the carrier gas. It is a standard detector for determination of inorganic gases –  $\text{H}_2$ ,  $\text{O}_2$ ,  $\text{N}_2$ ,  $\text{CS}_2$  and  $\text{H}_2\text{O}$ . Most analytes have low thermal conductivities, so the highest sensitivity is attained by using a carrier gas of very high thermal conductivity  $\rightarrow$  H or He (usually a 10 factor increase)



# Instrumentation

## Detectors

### Thermal Conductivity Detector (TCD)

Linear dynamic range  $10^3$  or sometimes higher.

TCD is housed (heated) separate from the column oven  
→ because temperature control is crucial.

Also filament must be protected from  $O_2$  while hot.

TCD is classified as a concentration/nondestructive detector. The response is proportional to the relative concentration of analyte in the carrier gas (i.e. mass of solute per unit volume of carrier gas).

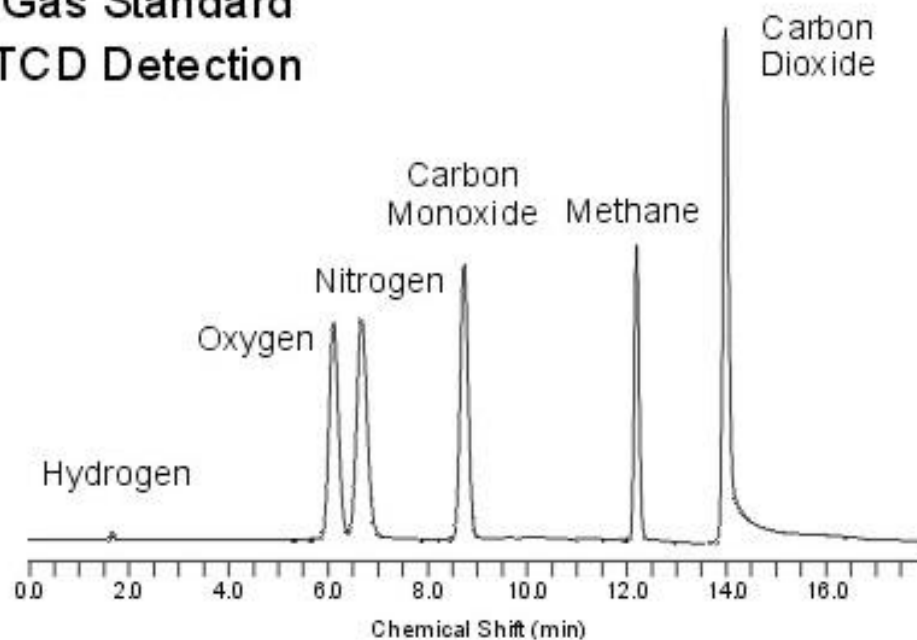


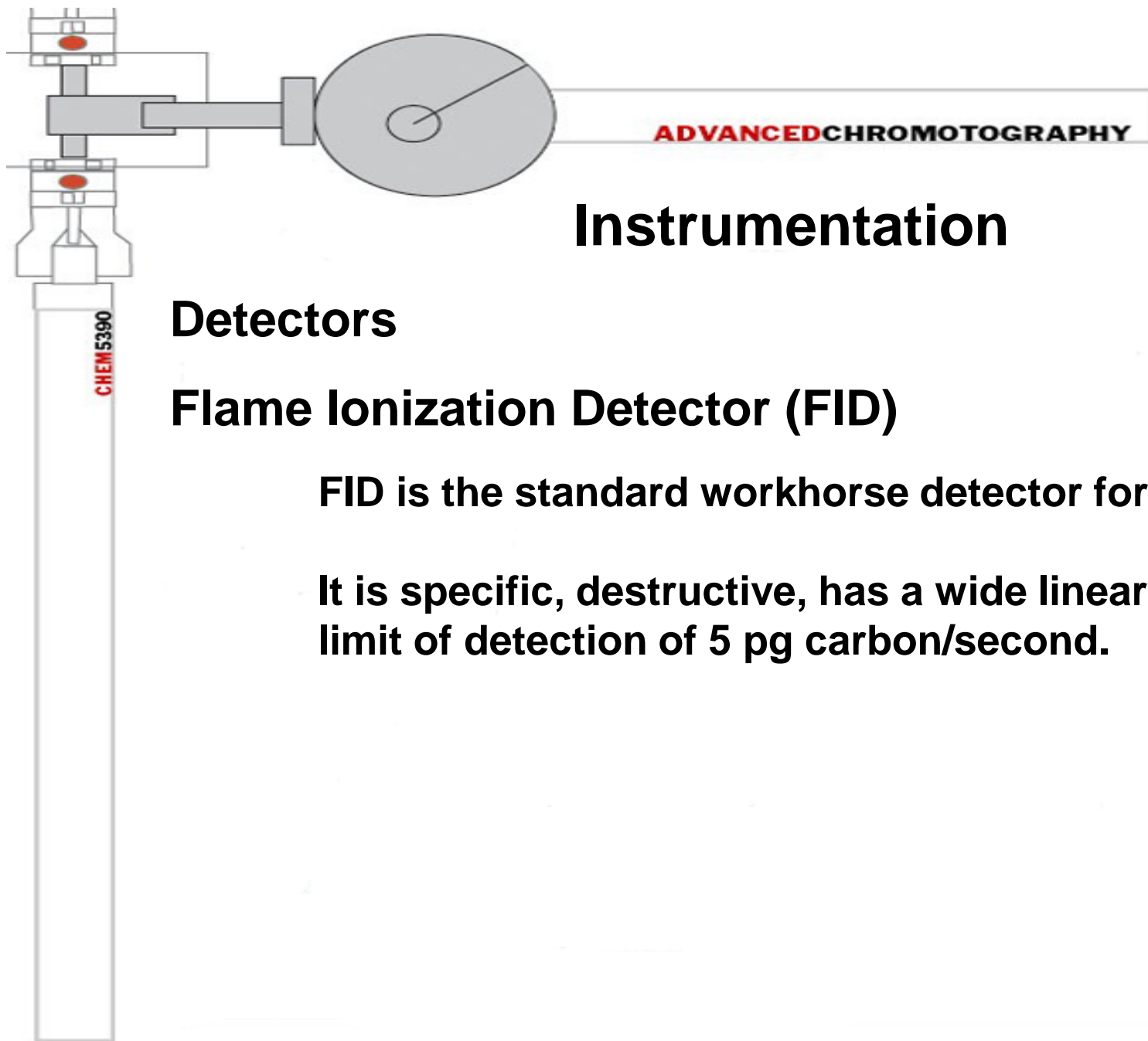
# Instrumentation

## Detectors

### Thermal Conductivity Detector (TCD)

Gas Standard  
TCD Detection





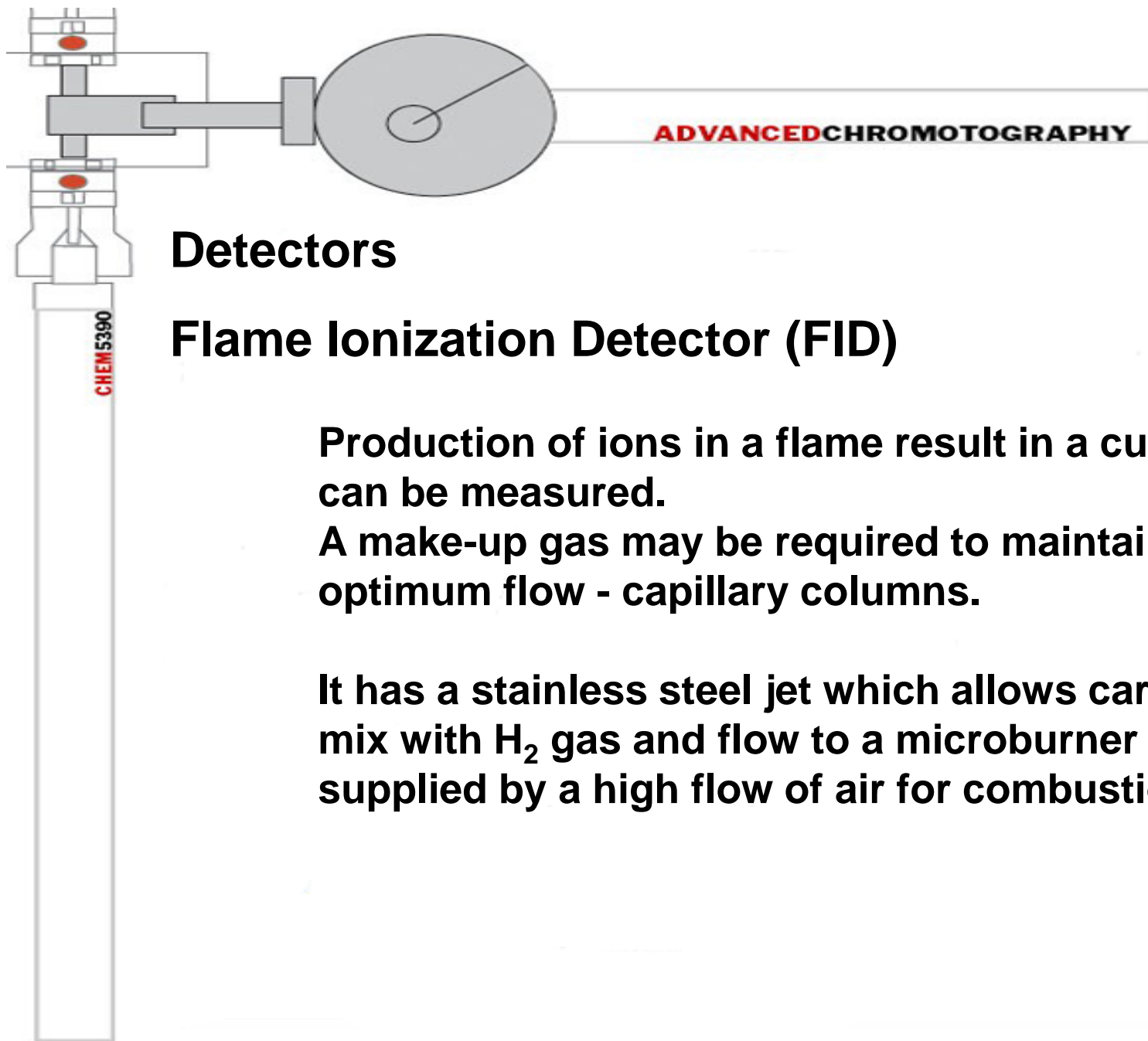
# Instrumentation

## Detectors

### Flame Ionization Detector (FID)

FID is the standard workhorse detector for the GC.

It is specific, destructive, has a wide linear range and a limit of detection of 5 pg carbon/second.



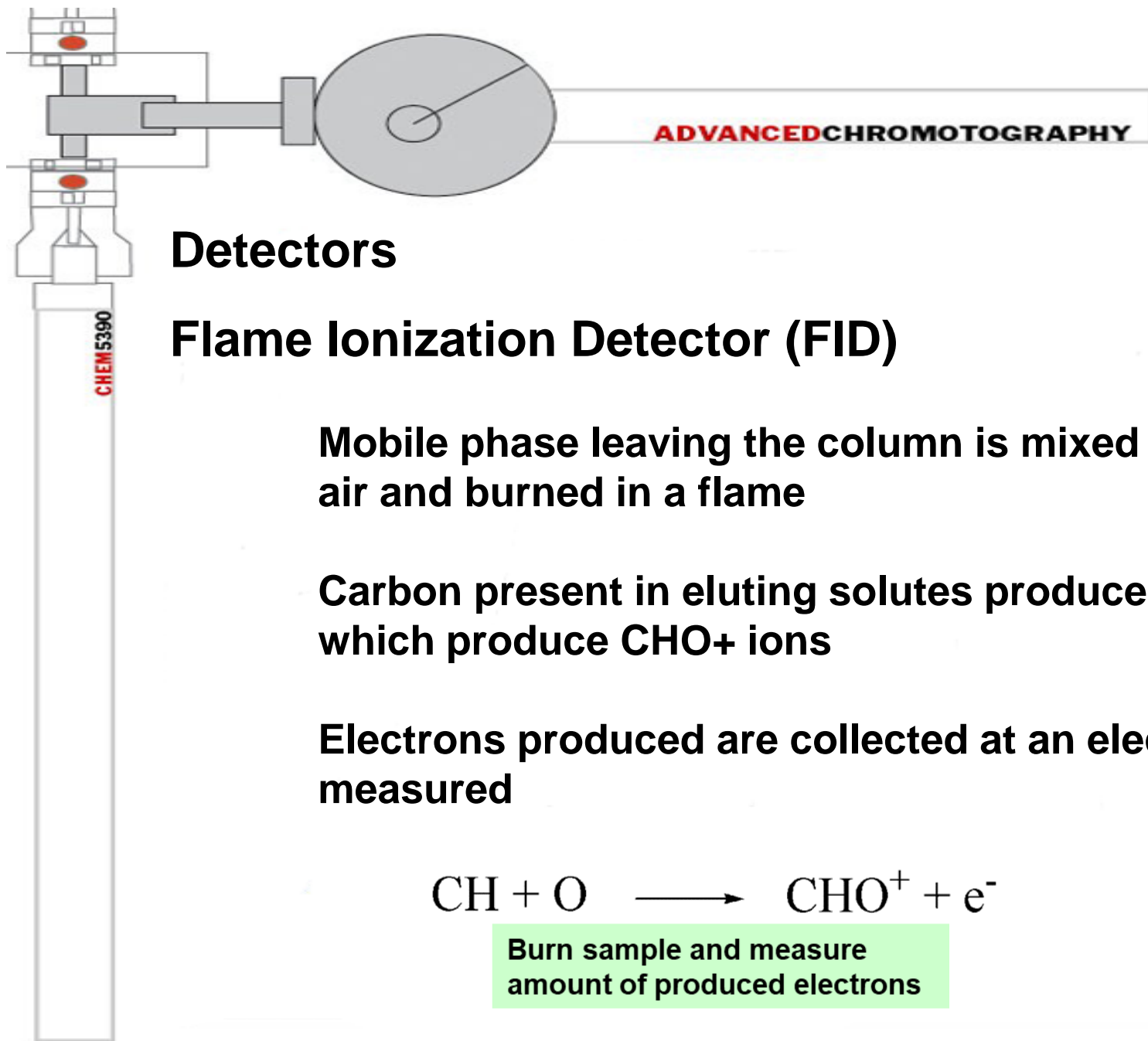
## **Detectors**

### **Flame Ionization Detector (FID)**

**Production of ions in a flame result in a current that can be measured.**

**A make-up gas may be required to maintain an optimum flow - capillary columns.**

**It has a stainless steel jet which allows carrier gas to mix with H<sub>2</sub> gas and flow to a microburner tip, which is supplied by a high flow of air for combustion.**

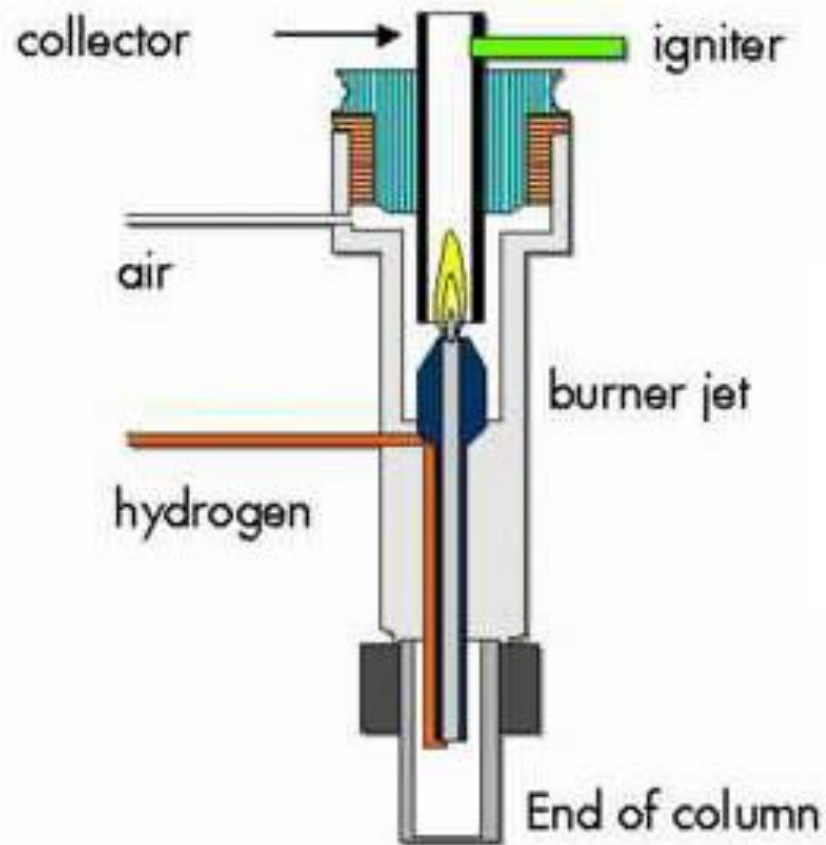
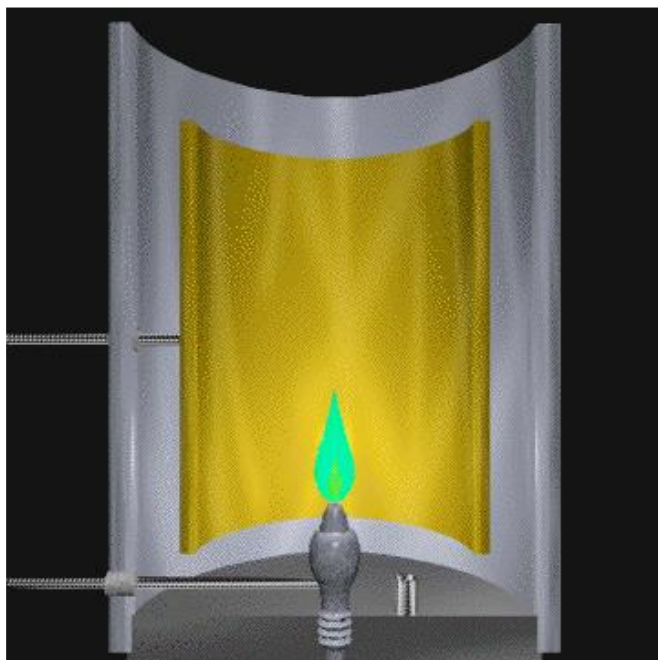




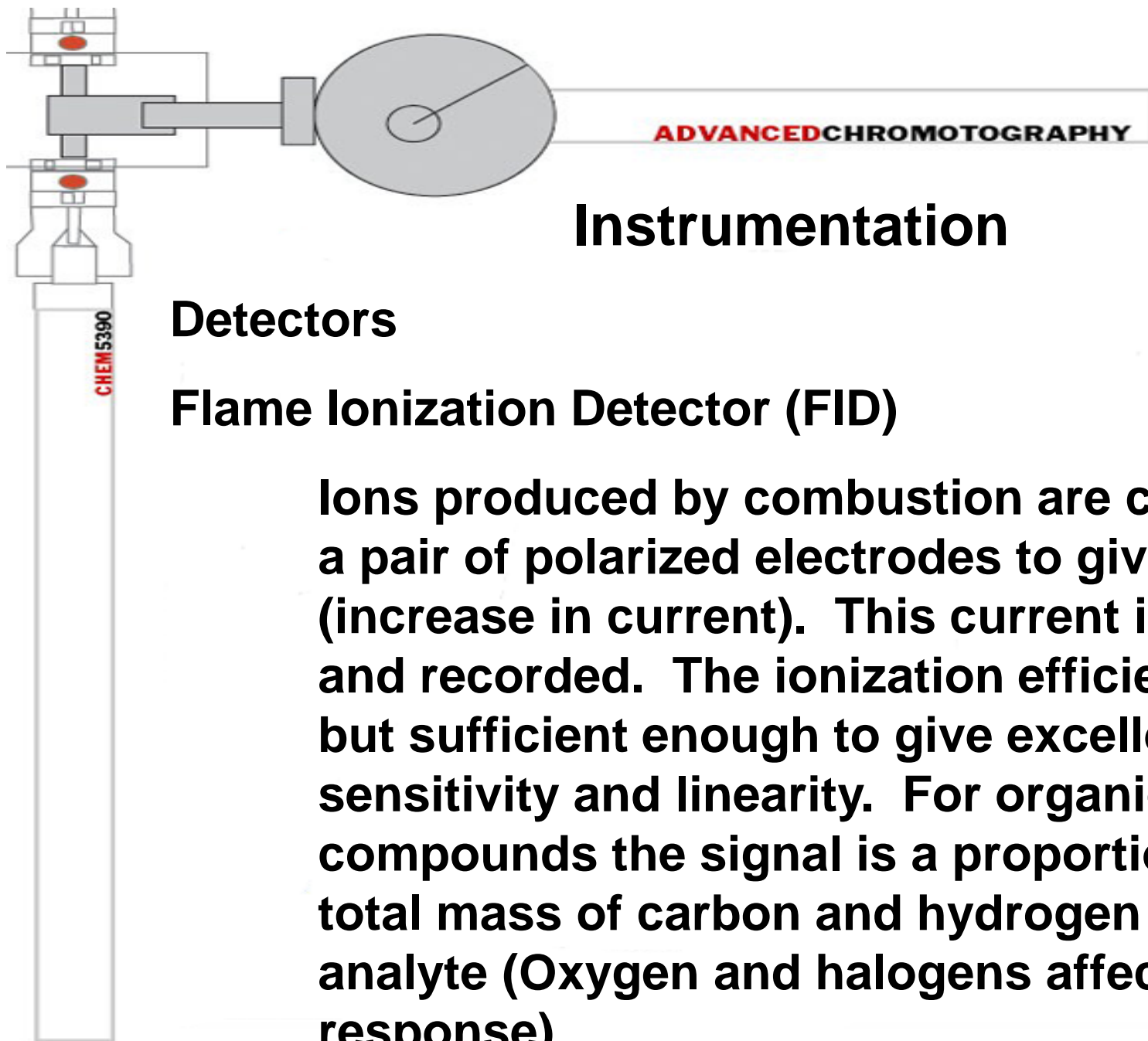
## Instrumentation

### Detectors

### Flame Ionization Detector





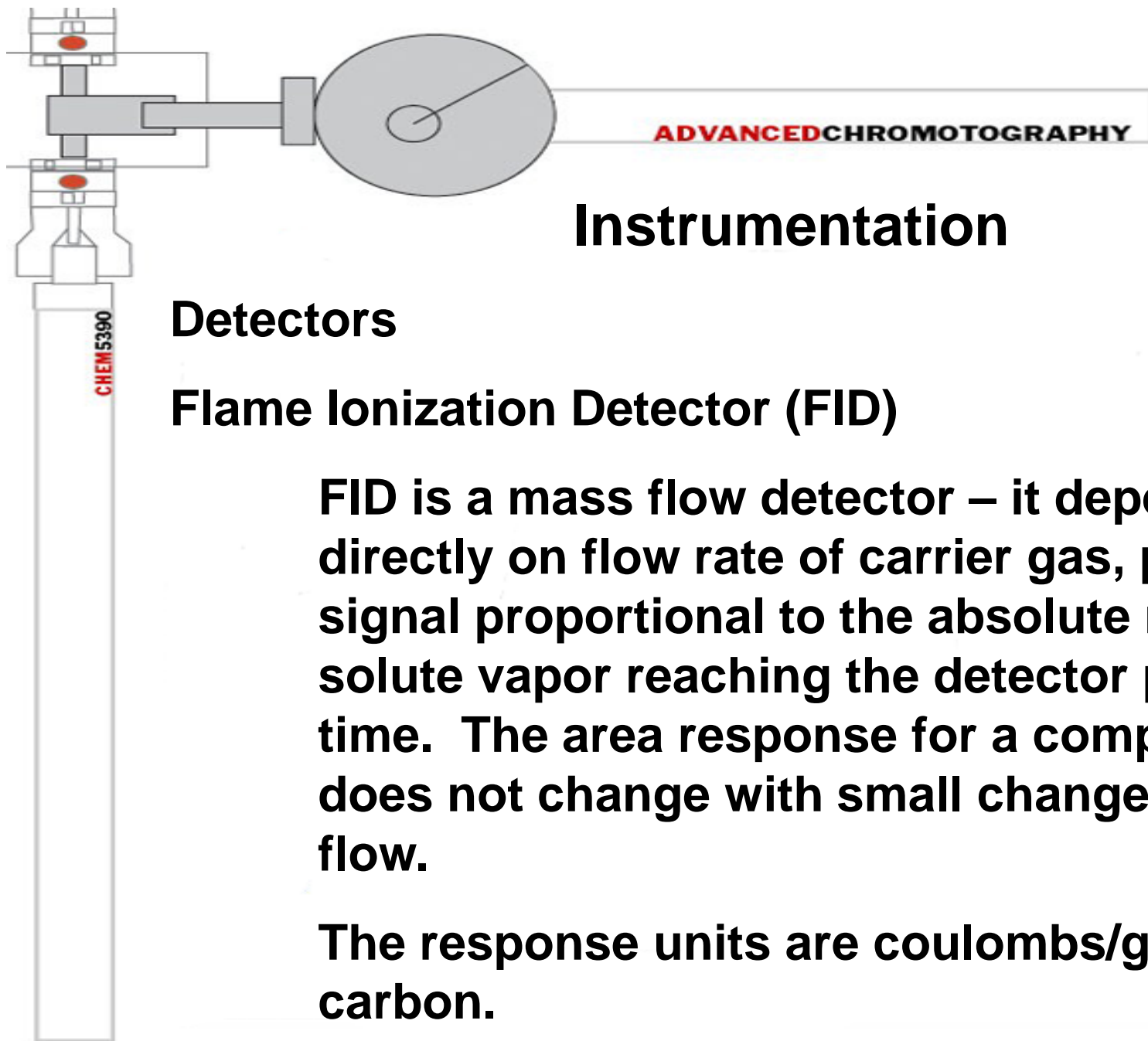


## Instrumentation

### Detectors

#### Flame Ionization Detector (FID)

Ions produced by combustion are collected at a pair of polarized electrodes to give a signal (increase in current). This current is amplified and recorded. The ionization efficiency is low but sufficient enough to give excellent sensitivity and linearity. For organic compounds the signal is proportional to the total mass of carbon and hydrogen in the analyte (Oxygen and halogens affect this response).



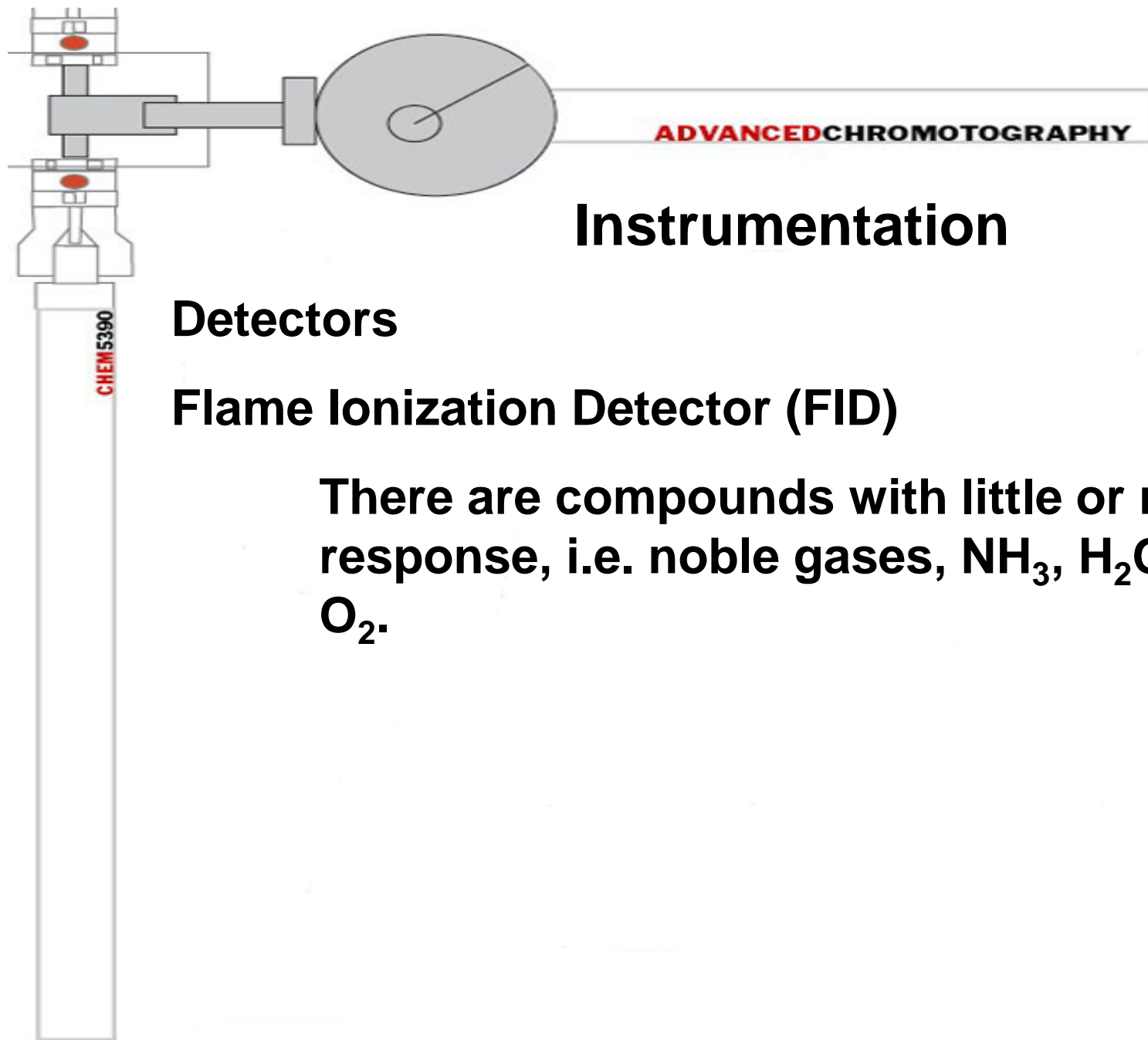
## Instrumentation

### Detectors

#### Flame Ionization Detector (FID)

**FID is a mass flow detector – it depends directly on flow rate of carrier gas, produces a signal proportional to the absolute mass of solute vapor reaching the detector per unit time. The area response for a compound does not change with small changes in carrier flow.**

**The response units are coulombs/gram of carbon.**



# Instrumentation

## Detectors

### Flame Ionization Detector (FID)

There are compounds with little or no FID response, i.e. noble gases,  $\text{NH}_3$ ,  $\text{H}_2\text{O}$ ,  $\text{CO}_2$ ,  $\text{N}_2$ ,  $\text{O}_2$ .

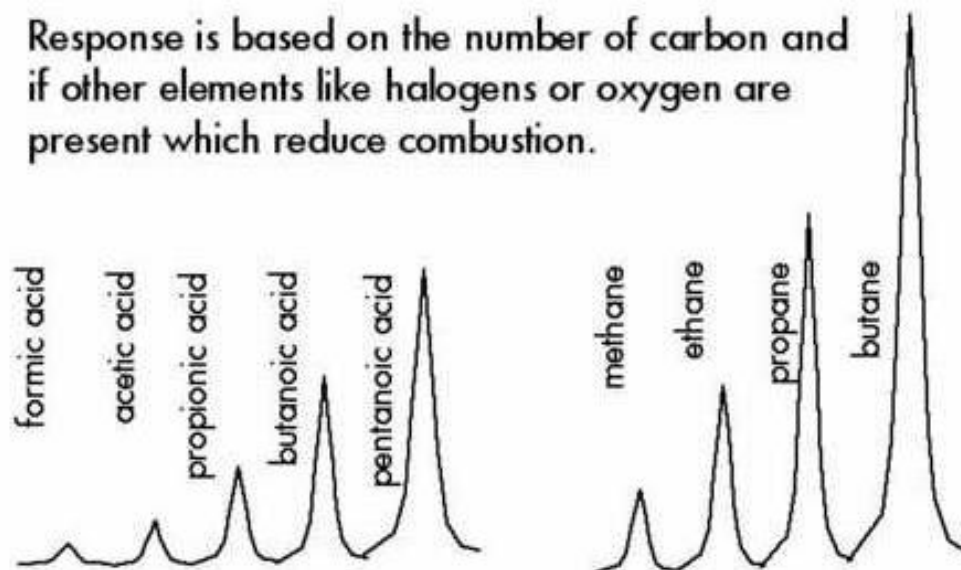
# Instrumentation

## Detectors

### Flame Ionization Detector (FID)

The effective carbon number (ECN) has been developed to estimate the relative response for any compound.

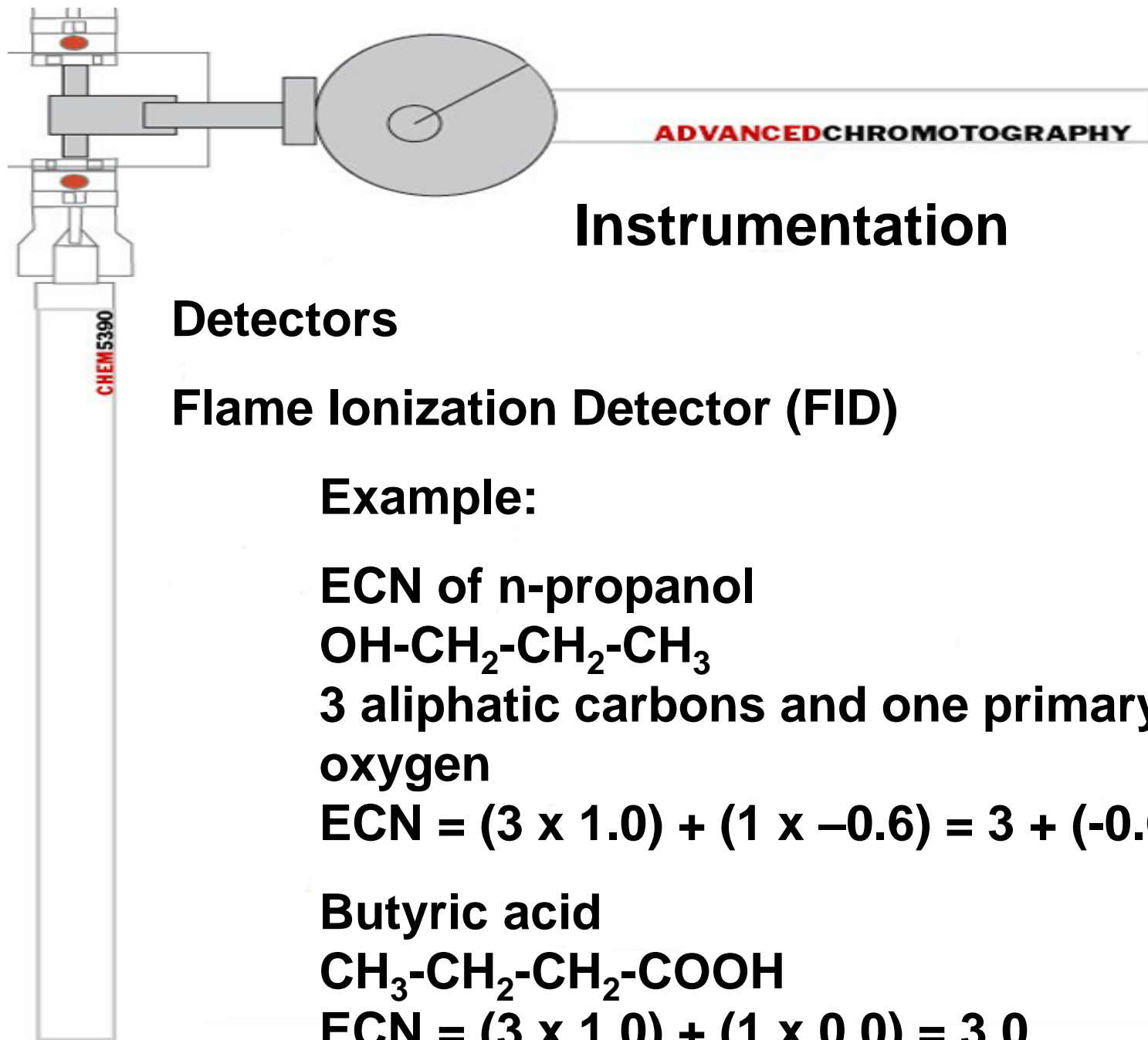
Response is based on the number of carbon and if other elements like halogens or oxygen are present which reduce combustion.



# Instrumentation

**TABLE 5.3 Contributions to Effective Carbon Number**

Atom	Type	Effective Carbon No. Contribution
C	Aliphatic	1.0
C	Aromatic	1.0
C	Olefinic	0.95
C	Acetylenic	1.30
C	Carbonyl	0.0
C	Nitrile	0.3
O	Ether	-1.0
O	Primary alcohol	-0.6
O	Secondary alcohol	-0.75
O	Tertiary alcohol, esters	-0.25
Cl	Two or more on single aliphatic C	-0.12 each
Cl	On olefinic C	+0.05
N	In amines	Similar to O in corresponding alcohols



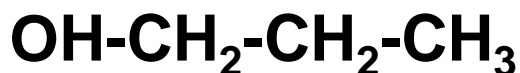
## Instrumentation

### Detectors

#### Flame Ionization Detector (FID)

Example:

ECN of n-propanol



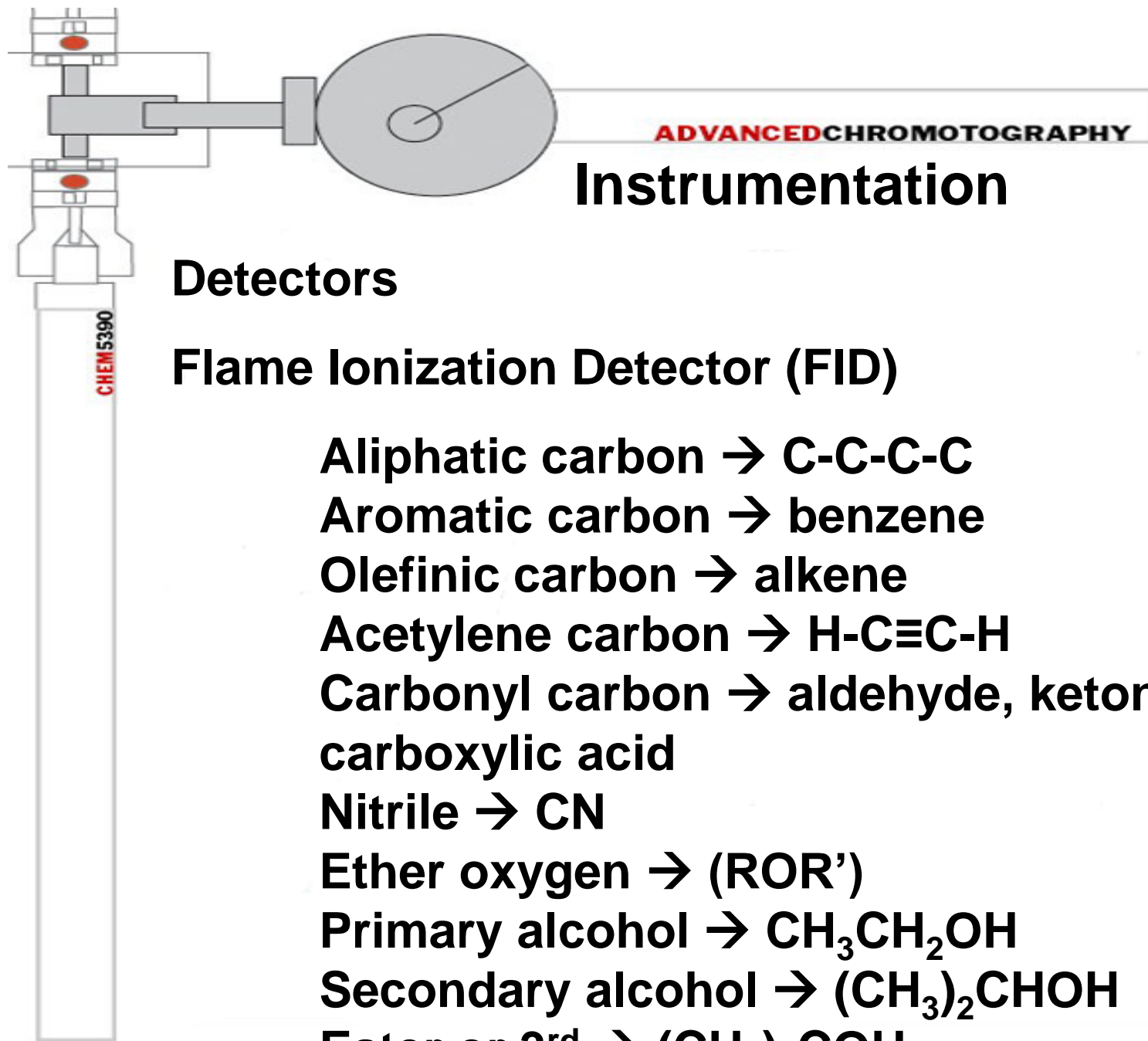
3 aliphatic carbons and one primary alcohol oxygen

$$\text{ECN} = (3 \times 1.0) + (1 \times -0.6) = 3 + (-0.6) = 2.4$$

Butyric acid



$$\text{ECN} = (3 \times 1.0) + (1 \times 0.0) = 3.0$$



## Instrumentation

### Detectors

#### Flame Ionization Detector (FID)

Aliphatic carbon  $\rightarrow$  C-C-C-C

Aromatic carbon  $\rightarrow$  benzene

Olefinic carbon  $\rightarrow$  alkene

Acetylene carbon  $\rightarrow$  H-C $\equiv$ C-H

Carbonyl carbon  $\rightarrow$  aldehyde, ketone, carboxylic acid

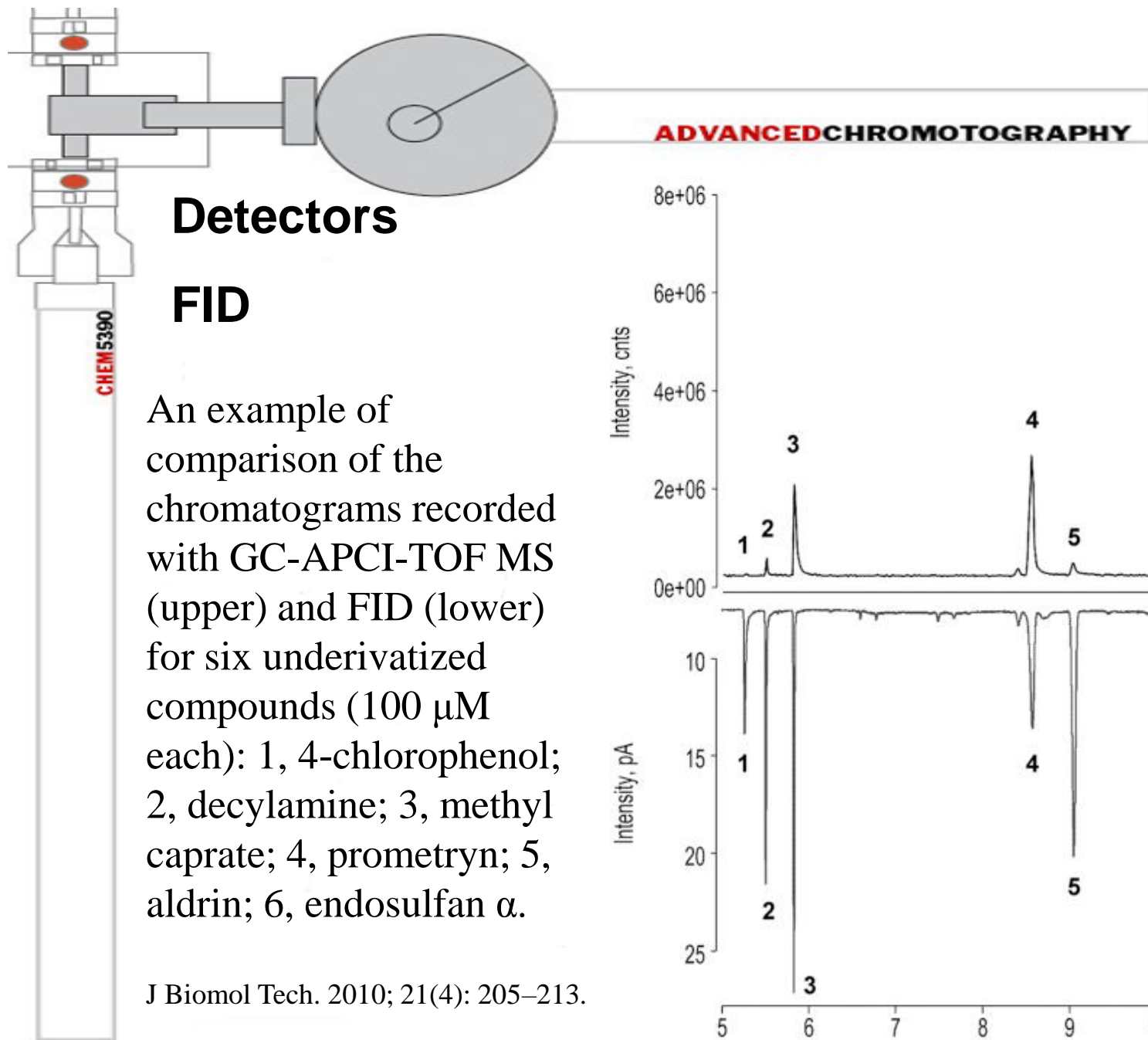
Nitrile  $\rightarrow$  CN

Ether oxygen  $\rightarrow$  (ROR')

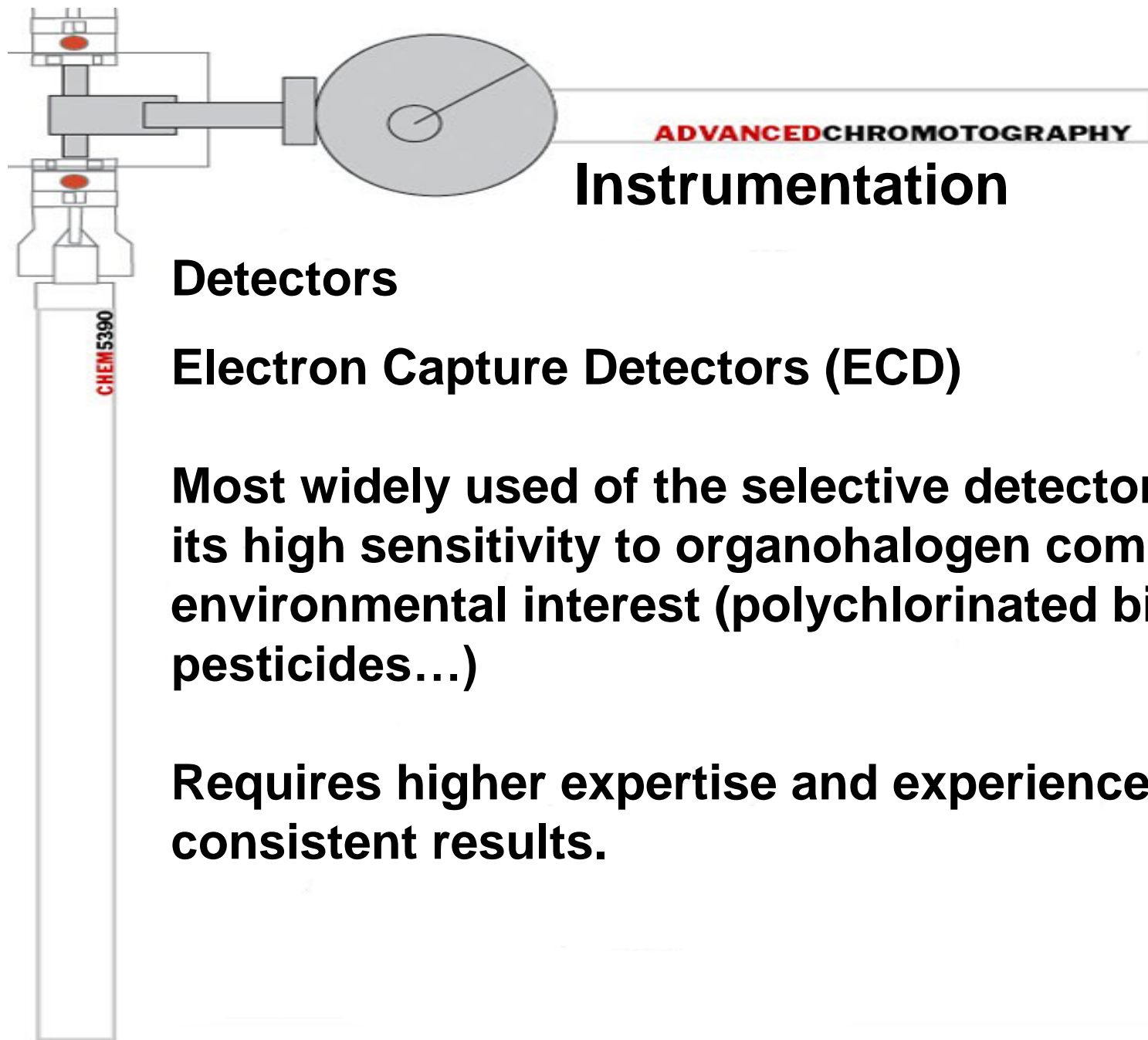
Primary alcohol  $\rightarrow$  CH<sub>3</sub>CH<sub>2</sub>OH

Secondary alcohol  $\rightarrow$  (CH<sub>3</sub>)<sub>2</sub>CHOH

Ester or 3<sup>rd</sup>  $\rightarrow$  (CH<sub>3</sub>)<sub>3</sub>COH







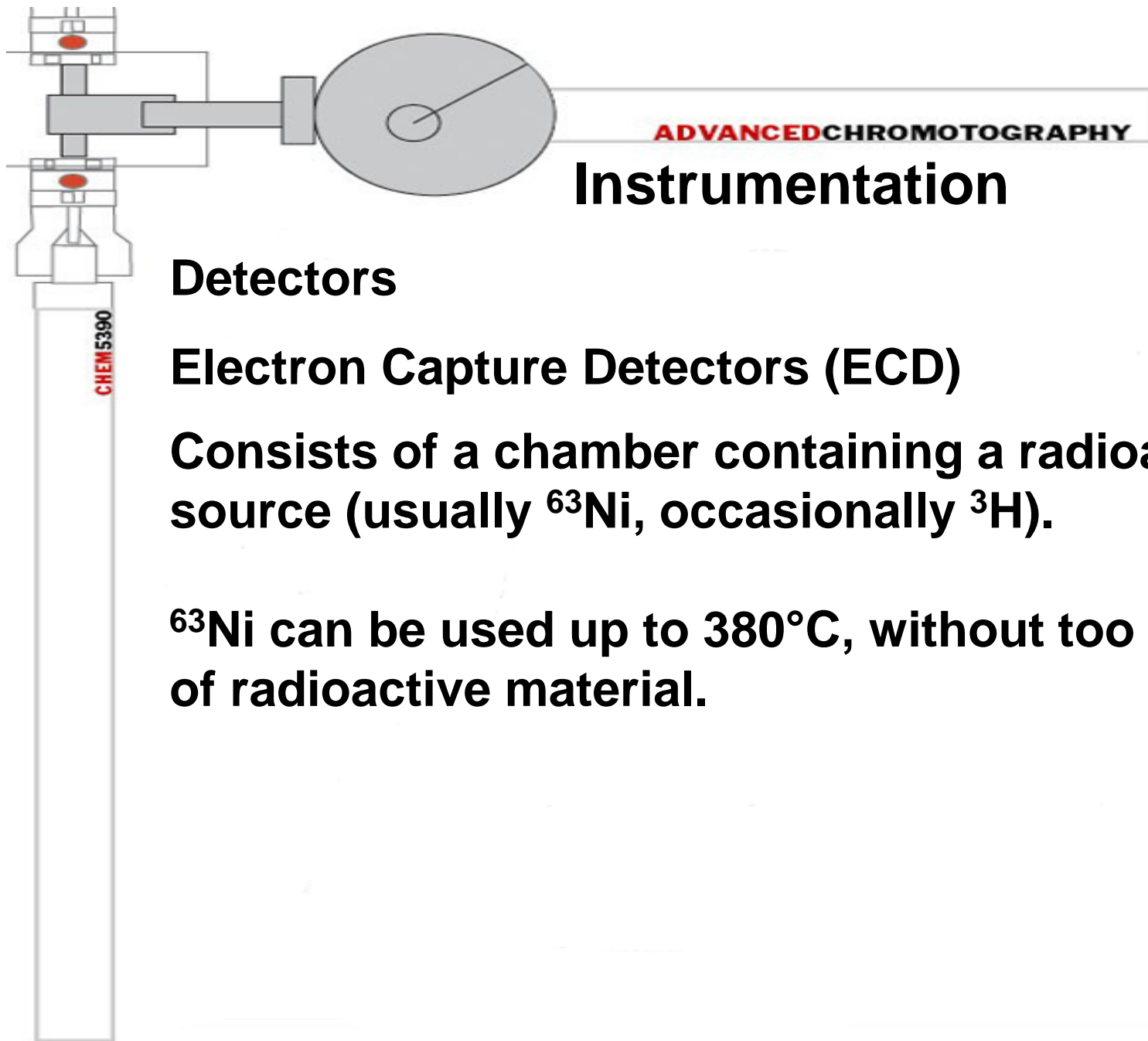
## Instrumentation

### Detectors

#### Electron Capture Detectors (ECD)

**Most widely used of the selective detectors due to its high sensitivity to organohalogen compounds of environmental interest (polychlorinated biphenyls, pesticides...)**

**Requires higher expertise and experience to achieve consistent results.**



**ADVANCED CHROMATOGRAPHY**

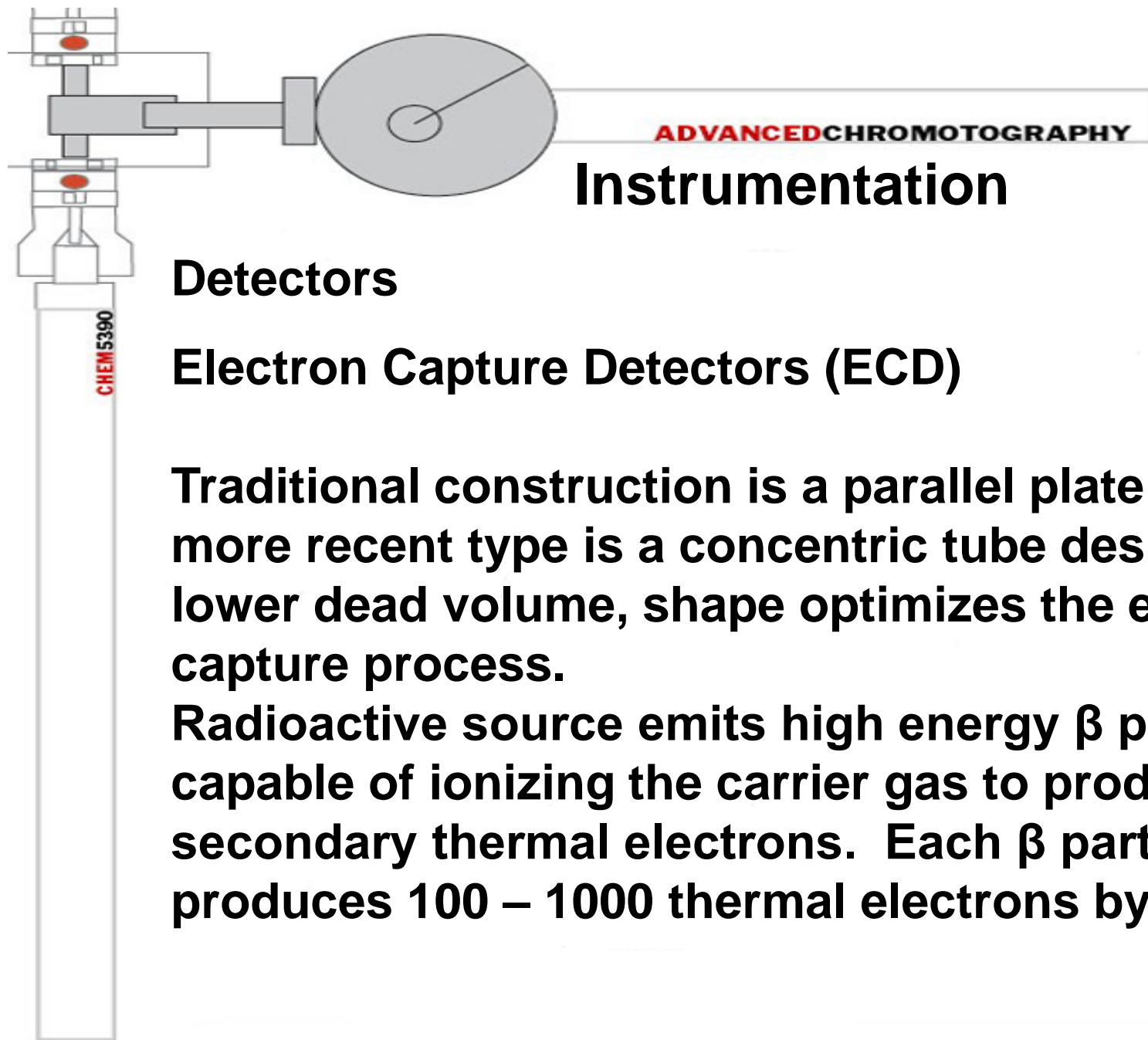
## **Instrumentation**

### **Detectors**

#### **Electron Capture Detectors (ECD)**

**Consists of a chamber containing a radioactive source (usually  $^{63}\text{Ni}$ , occasionally  $^3\text{H}$ ).**

**$^{63}\text{Ni}$  can be used up to  $380^\circ\text{C}$ , without too much loss of radioactive material.**



## Instrumentation

### Detectors

#### Electron Capture Detectors (ECD)

Traditional construction is a parallel plate design – more recent type is a concentric tube design with lower dead volume, shape optimizes the electron capture process.

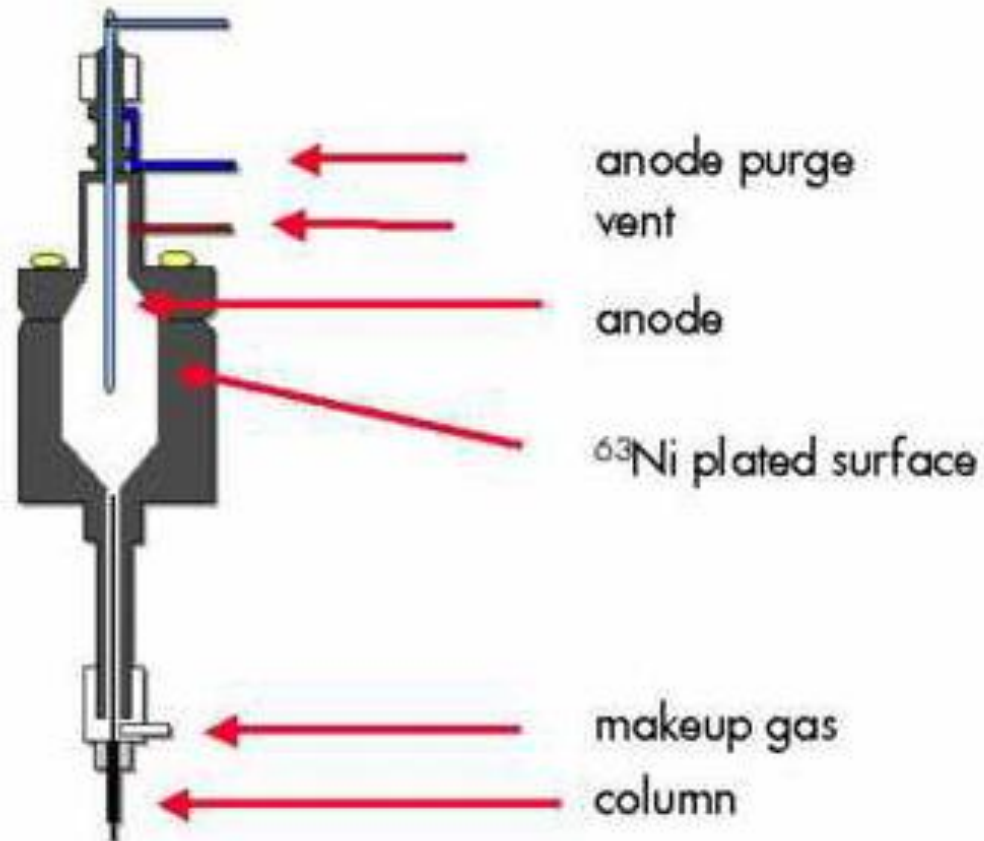
Radioactive source emits high energy  $\beta$  particles capable of ionizing the carrier gas to produce secondary thermal electrons. Each  $\beta$  particle produces 100 – 1000 thermal electrons by collision.

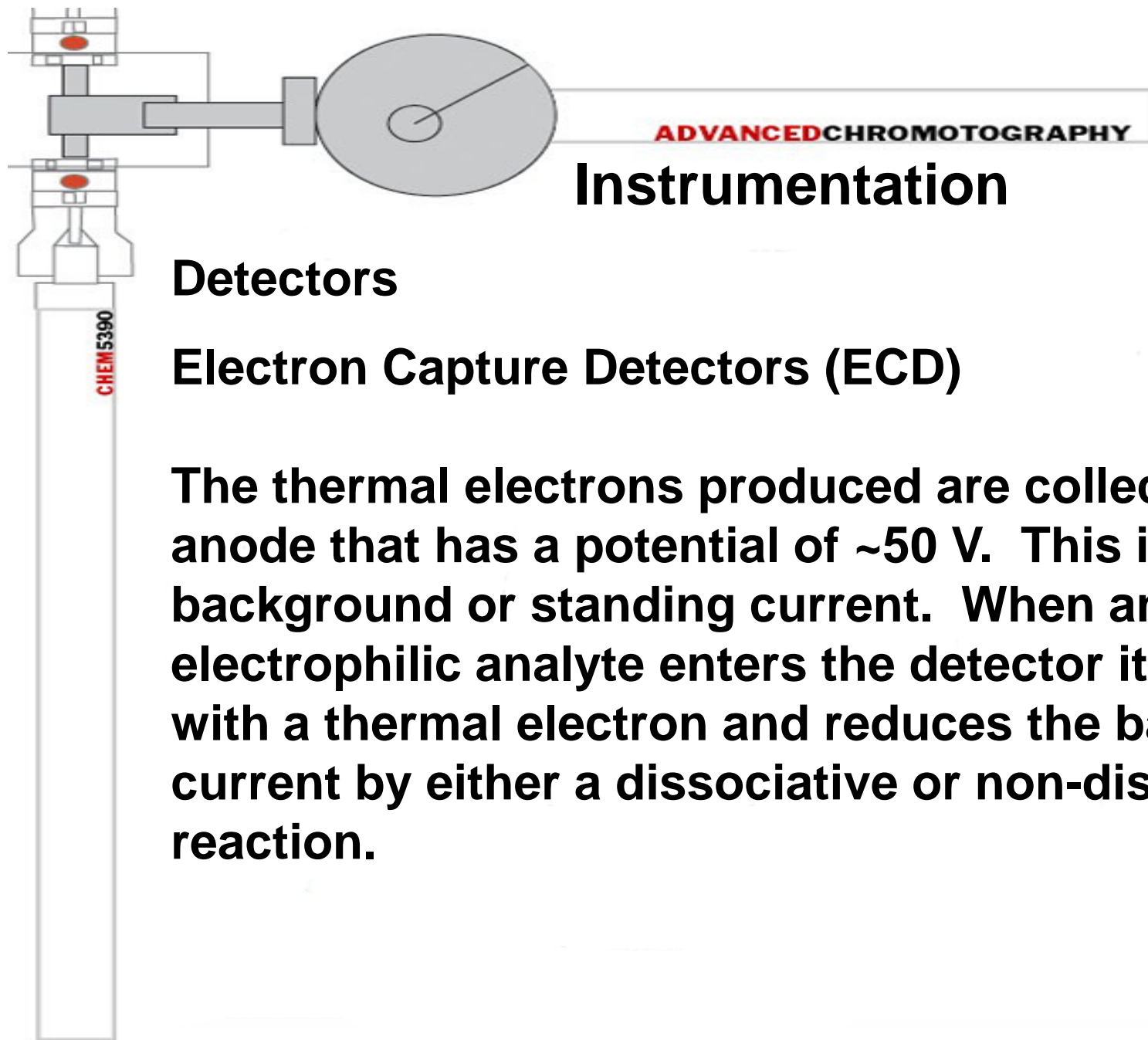
**ADVANCED**CHROMATOGRAPHY

## Instrumentation

### Detectors

### Electron Capture Detectors (ECD)



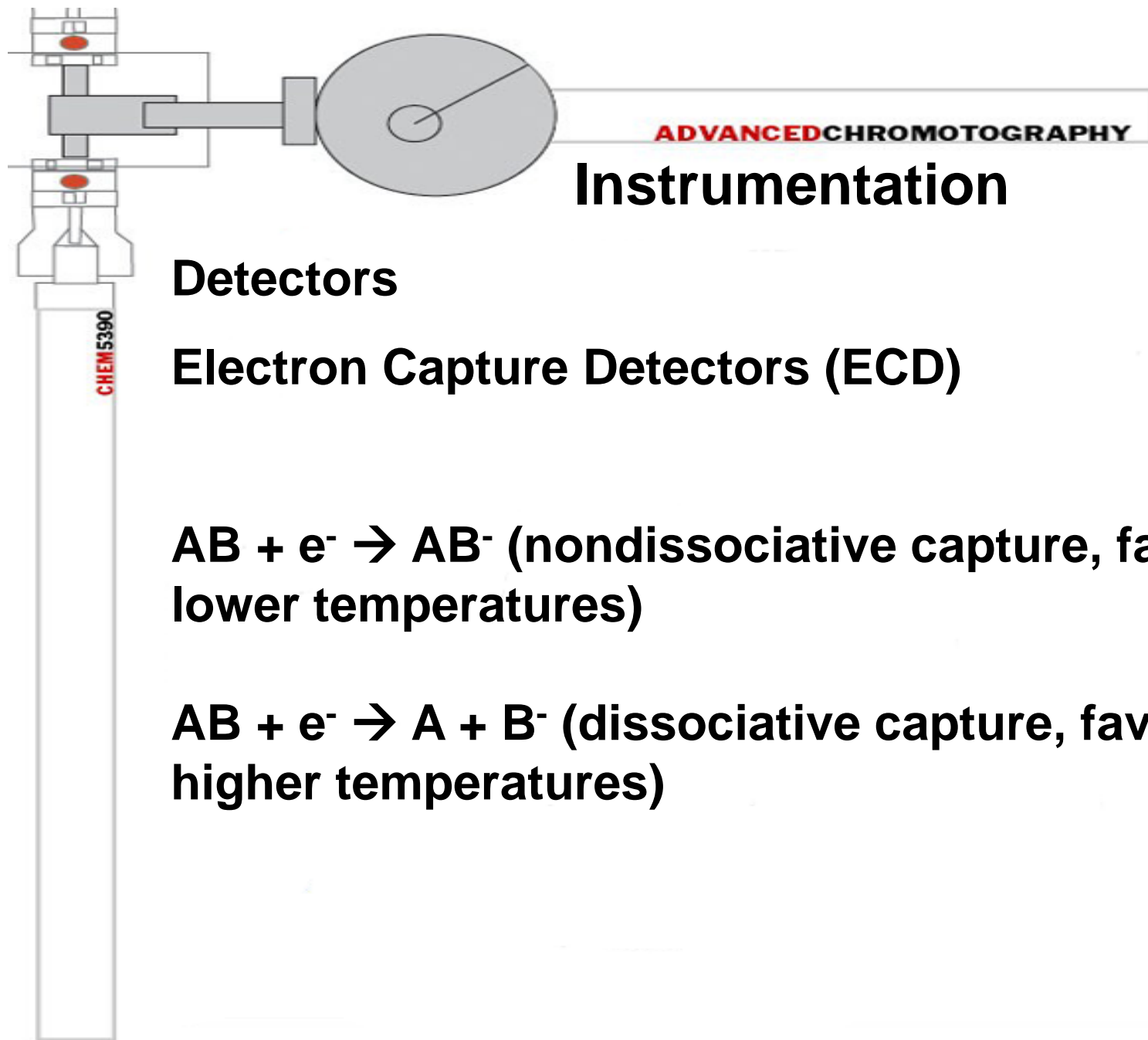


## Instrumentation

### Detectors

#### Electron Capture Detectors (ECD)

The thermal electrons produced are collected at an anode that has a potential of  $\sim 50$  V. This is the background or standing current. When an electrophilic analyte enters the detector it collides with a thermal electron and reduces the background current by either a dissociative or non-dissociative reaction.

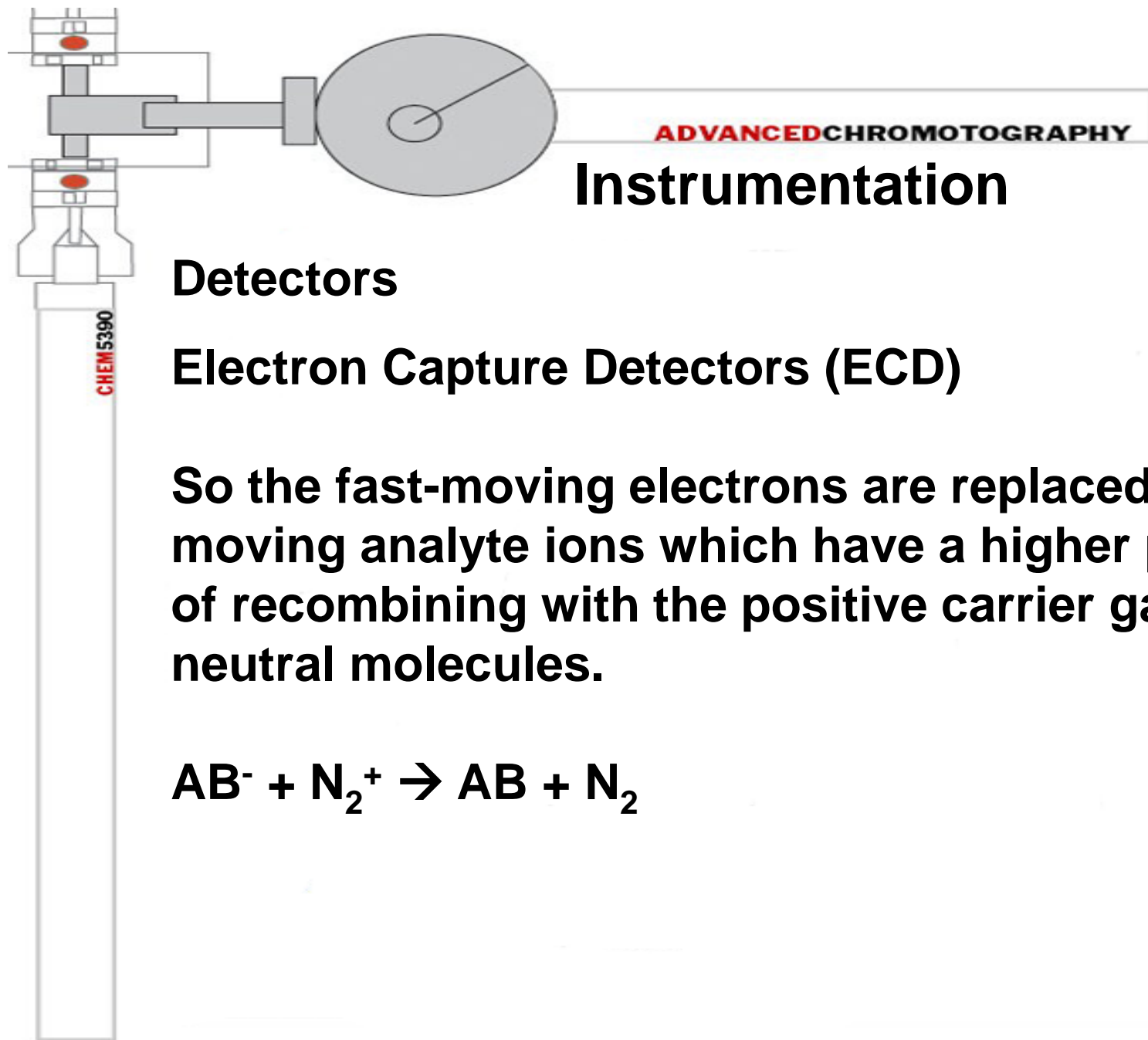


## **Detectors**

### **Electron Capture Detectors (ECD)**

**$AB + e^- \rightarrow AB^-$  (nondissociative capture, favored at lower temperatures)**

**$AB + e^- \rightarrow A + B^-$  (dissociative capture, favored at higher temperatures)**

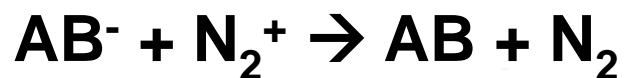


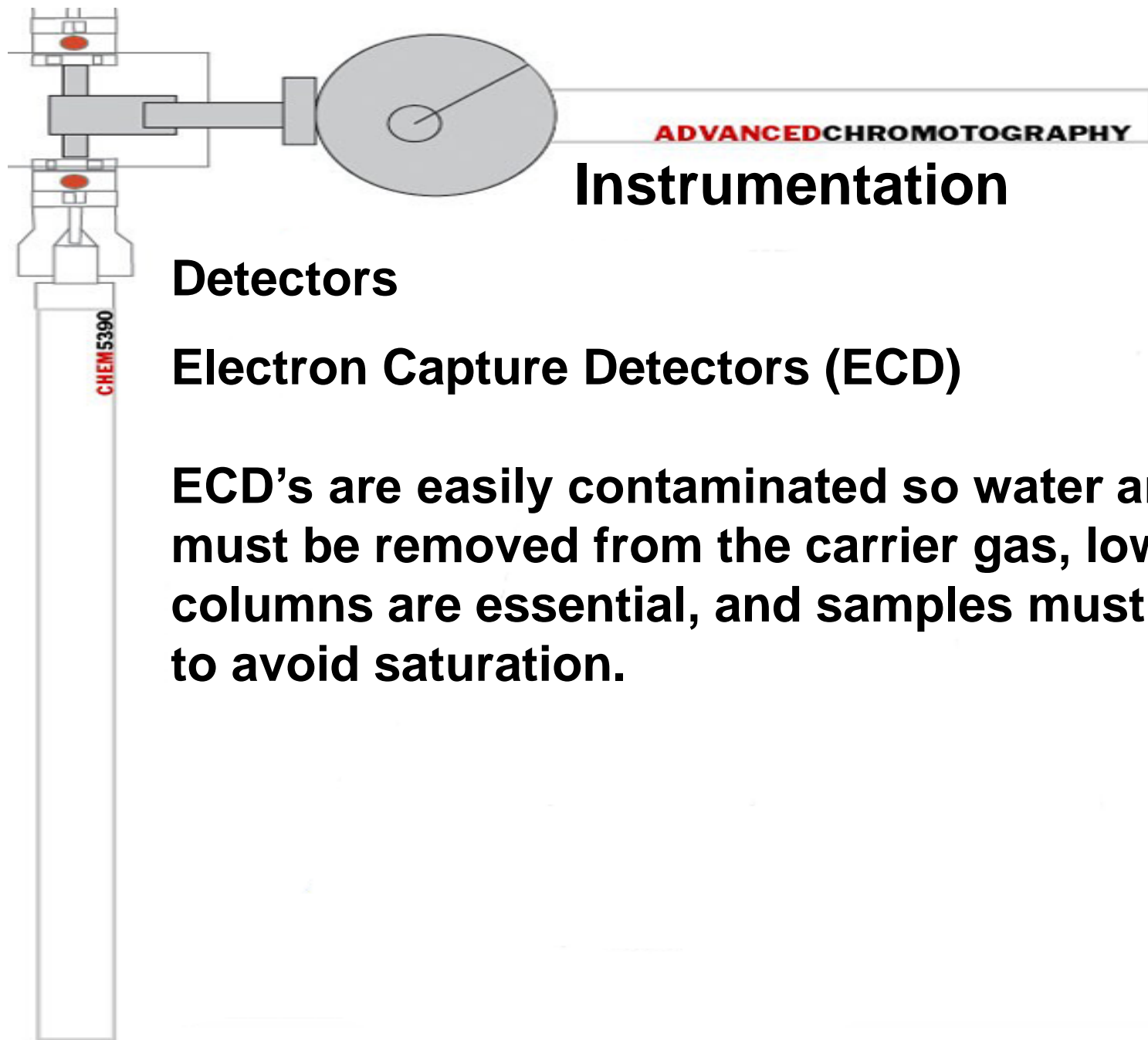
## Instrumentation

### Detectors

#### Electron Capture Detectors (ECD)

So the fast-moving electrons are replaced by slow-moving analyte ions which have a higher probability of recombining with the positive carrier gas to form neutral molecules.





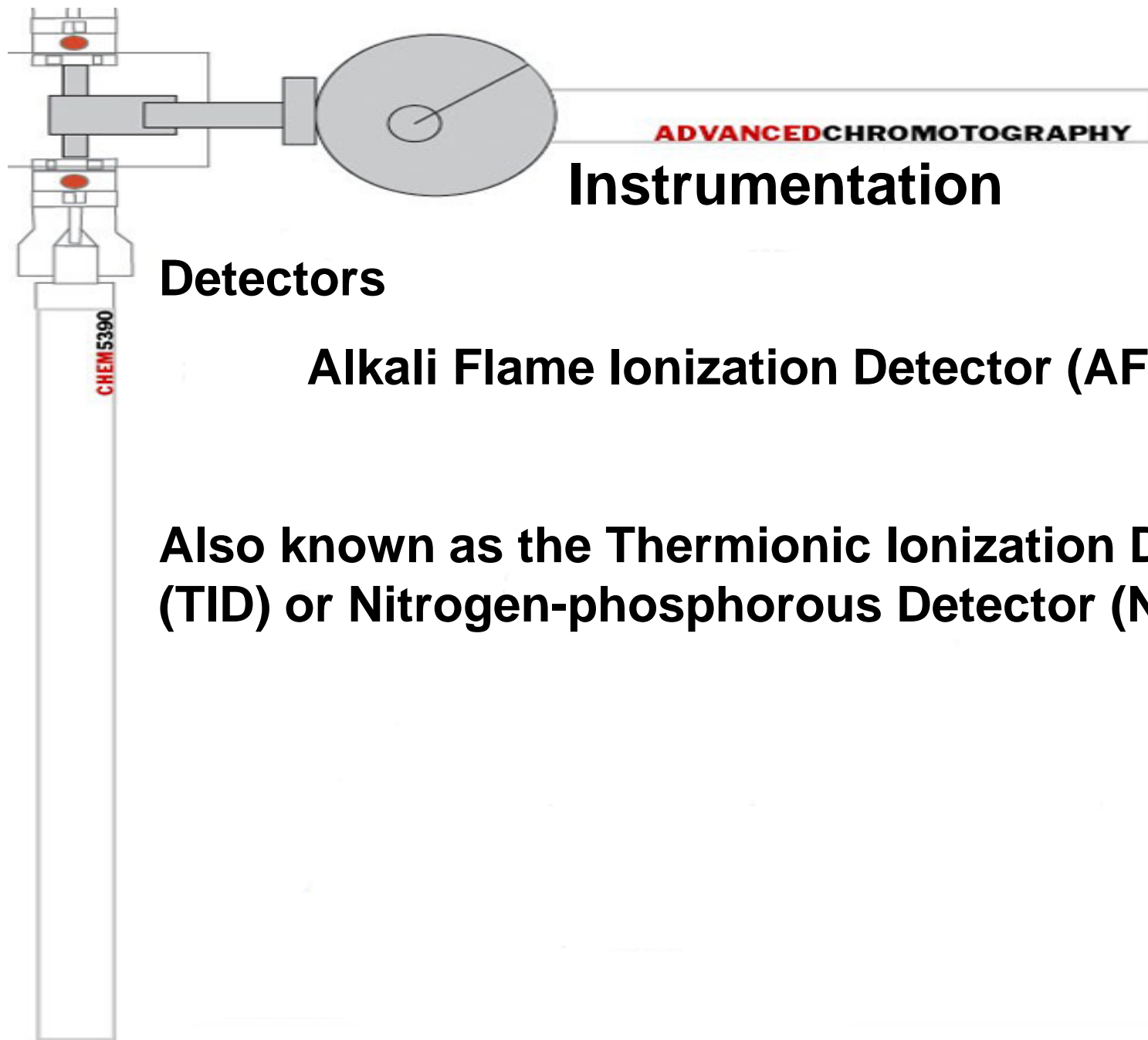
## Instrumentation

### Detectors

### Electron Capture Detectors (ECD)

ECD's are easily contaminated so water and oxygen must be removed from the carrier gas, low bleed columns are essential, and samples must be dilute to avoid saturation.



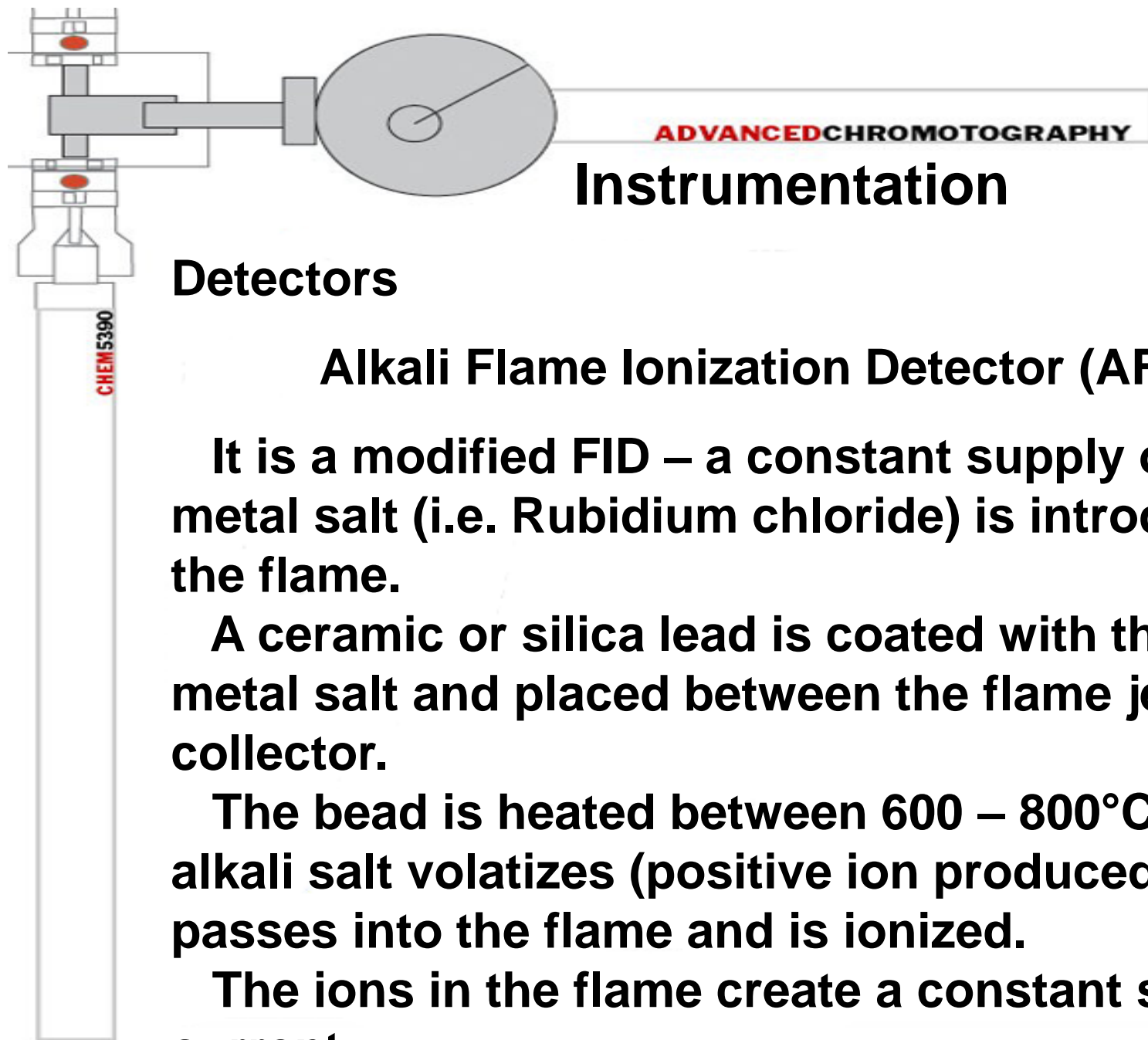


## Instrumentation

### Detectors

#### Alkali Flame Ionization Detector (AFID)

Also known as the Thermionic Ionization Detector (TID) or Nitrogen-phosphorous Detector (NPD).



## Instrumentation

### Detectors

#### Alkali Flame Ionization Detector (AFID)

It is a modified FID – a constant supply of an alkali metal salt (i.e. Rubidium chloride) is introduced into the flame.

A ceramic or silica bead is coated with the alkali metal salt and placed between the flame jet and ion collector.

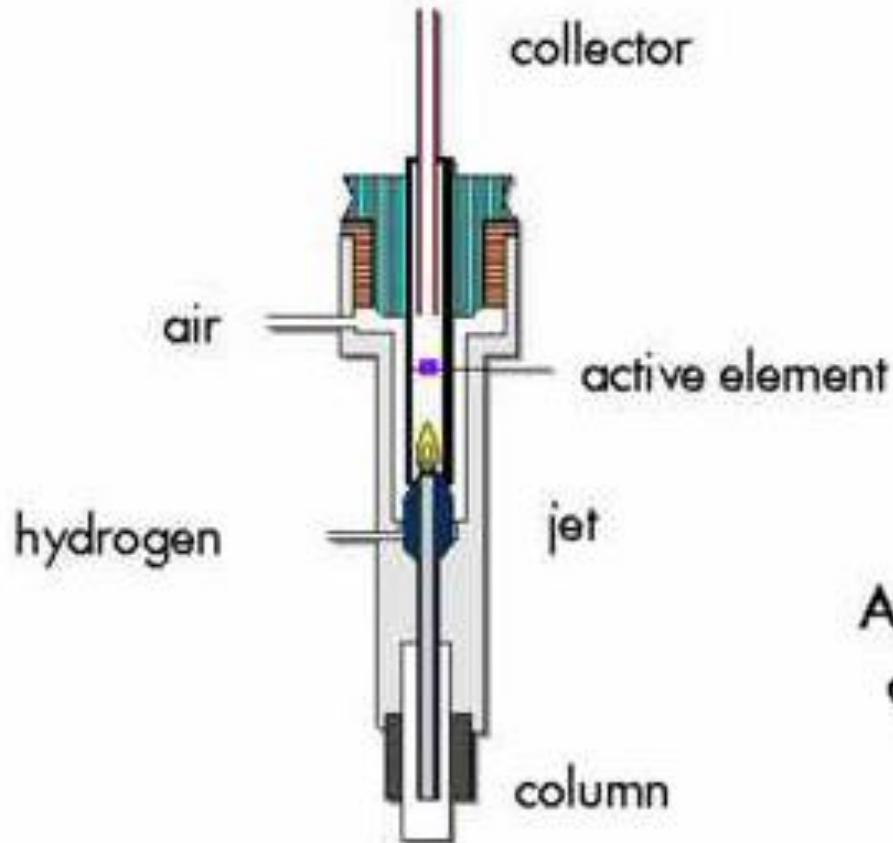
The bead is heated between 600 – 800°C. The alkali salt volatilizes (positive ion produced) and passes into the flame and is ionized.

The ions in the flame create a constant standing current.

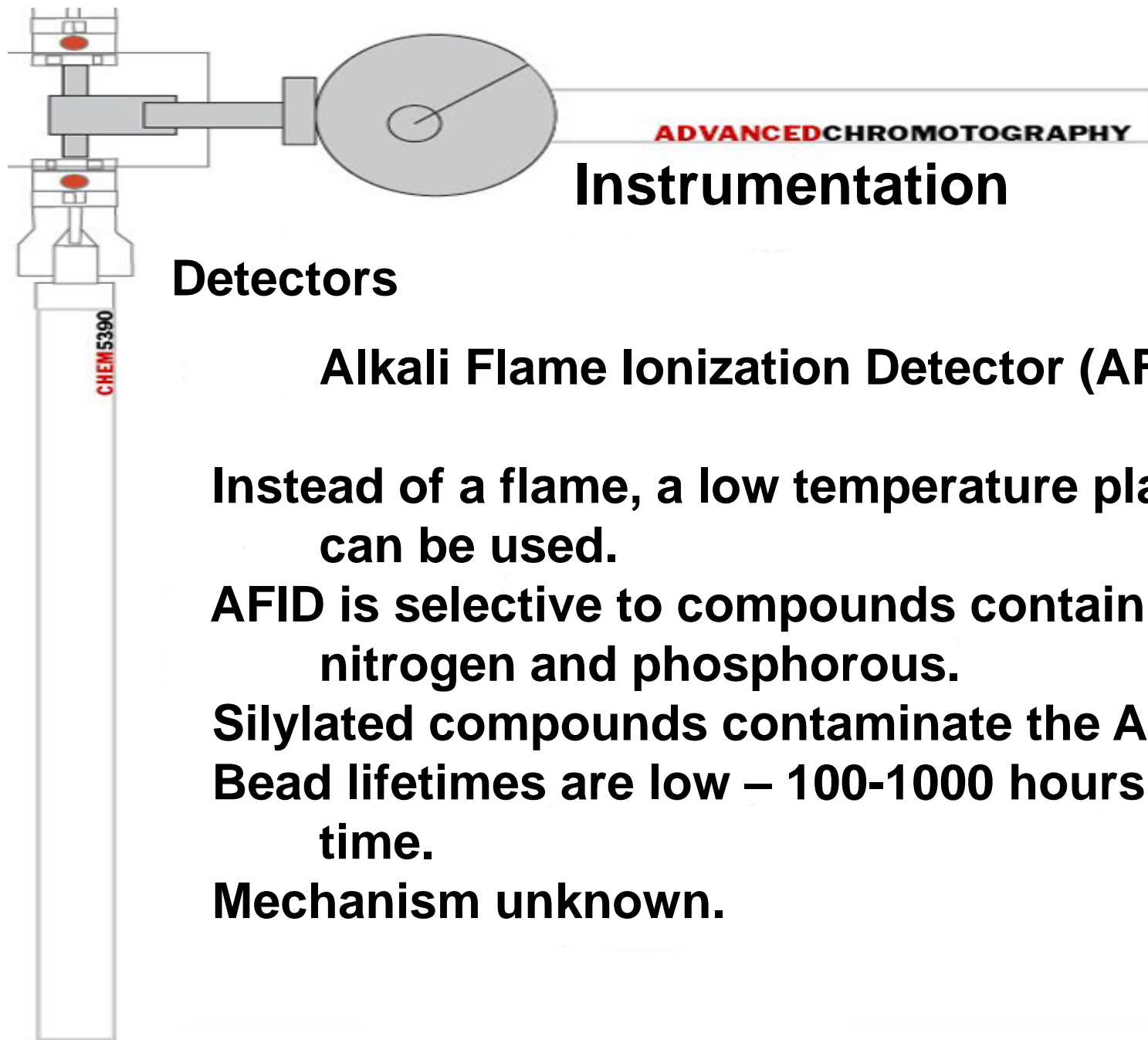
# Instrumentation

## Detectors

### Alkali Flame Ionization Detector (AFID)



Active element is  
a K or Rb salt.



## Instrumentation

### Detectors

#### Alkali Flame Ionization Detector (AFID)

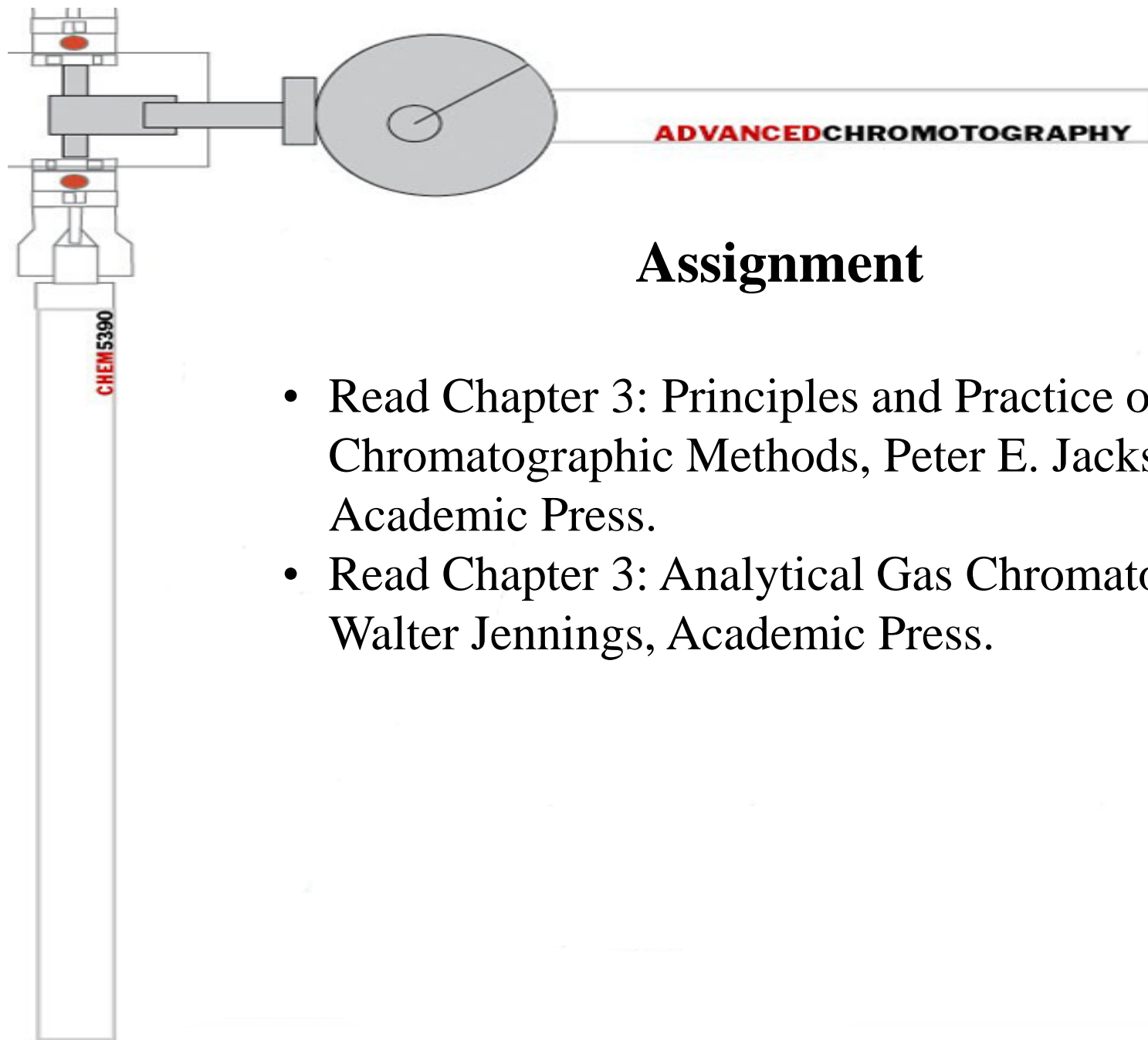
Instead of a flame, a low temperature plasma of  $H_2$  can be used.

AFID is selective to compounds containing nitrogen and phosphorous.

Silylated compounds contaminate the AFID.

Bead lifetimes are low – 100-1000 hours operating time.

Mechanism unknown.



## Assignment

- Read Chapter 3: Principles and Practice of Modern Chromatographic Methods, Peter E. Jackson, Academic Press.
- Read Chapter 3: Analytical Gas Chromatography, Walter Jennings, Academic Press.