FORENSIC TOXICOLOGY

Forensic Toxicology deals with the study of the adverse effects of drugs and chemicals on biological systems, and the interpretation of those results for legal purposes.

Forensic Toxicologist :

◦ Detect and identify drugs and toxins in body fluids, tissues and organs
◦ Interpret the potential biological consequences of the detected drugs or toxins

Forensic Toxicology has close ties to Analytical Toxicology and Analytical Chemistry.

Drugs and Toxins

A drug is any substance, other than a normal constituent of the body, that when applied to or introduced into a living organisms the effect of altering body functions.

Paracelus (1493-1541) – paraphrased ➔Physician-alchemist

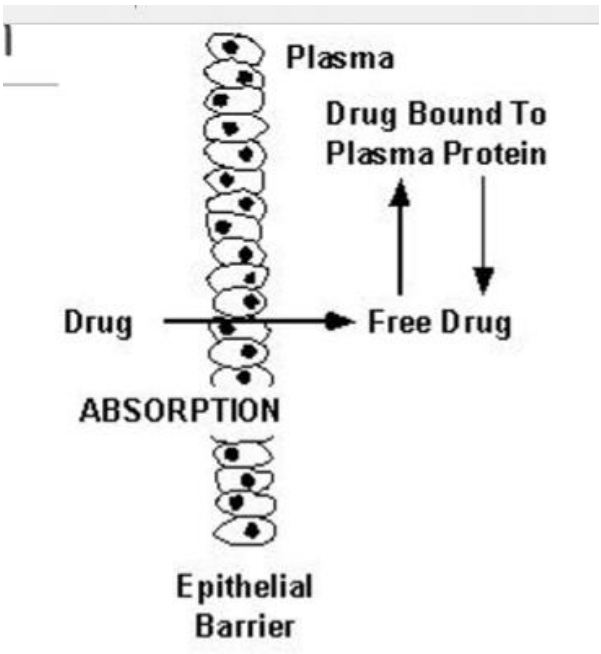
'All substances are poisons; there is none which is not a poison. The right dose differentiates a poison from a remedy.'

How are drugs and toxins handled by the body?

➔need to understand how drugs are handled by the body

The description of how drugs are handled by the body is often described as occurring in four phases:
Absorption, Distribution, Metabolism and Excretion : "ADME"

Absorption: Drugs must pass through an epithelial cell layer to enter the body



Routes of absorption :

➔gastrointestinal tract (stomach, small intestine, large intestine)

➔Sub-lingual

➔Lungs(inhalation)

➔Skin(dermal)

➔Injection – subcutaneous, intramuscular, intravenous

Implications :

➔The optimal site for maximum drug absorption is dependent on the physicochemical properties of the drug

➔Route of administration can influence how quickly the drug reaches the blood stream and the brain

➔After death, high local concentrations of drugs in the GI tract or lungs can be a clue as to the site of administration

Distribution :

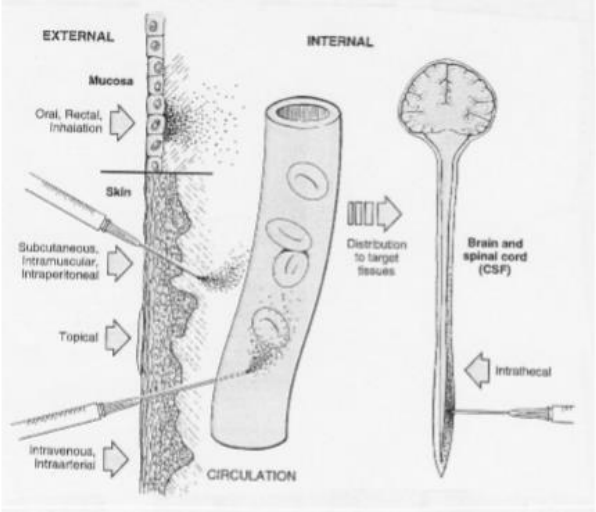
◦ Once drugs pass through the cellular membranes at the site of administration, most end up circulating in the blood

◦ The blood is responsible for distributing the drug to different tissues including the ultimate site of action (e.g. the brain)

◦ Blood (plasma or serum) concentrations are used as a surrogate measure of the levels of drug at the site of action

◦ Blood concentration versus effect relationships

Distribution with different administration methods



Metabolism (Biotransformation) :

➔Many drugs undergo biotransformation to metabolites before they are excreted from the body

➔enzymes located in body tissues catalyze this biotransformation : oxidation, reduction, conjugation reactions

➔The liver is the most common site for metabolic conversion : other tissues do play a role, dependant on the drug

➔Each drug has characteristic metabolic pathways

Implication :

◦The Forensic toxicologist needs to know the identity of metabolites
◦ Some drugs are metabolized so quickly that the only way to detect their presence is through the measurement of metabolites
◦ Some metabolites are pharmacologically active and can increase the length of time a subject is under the drug's influence
◦ Metabolites can remain in the body for long periods of time, influencing the time frame for detection

Example : Cocaine

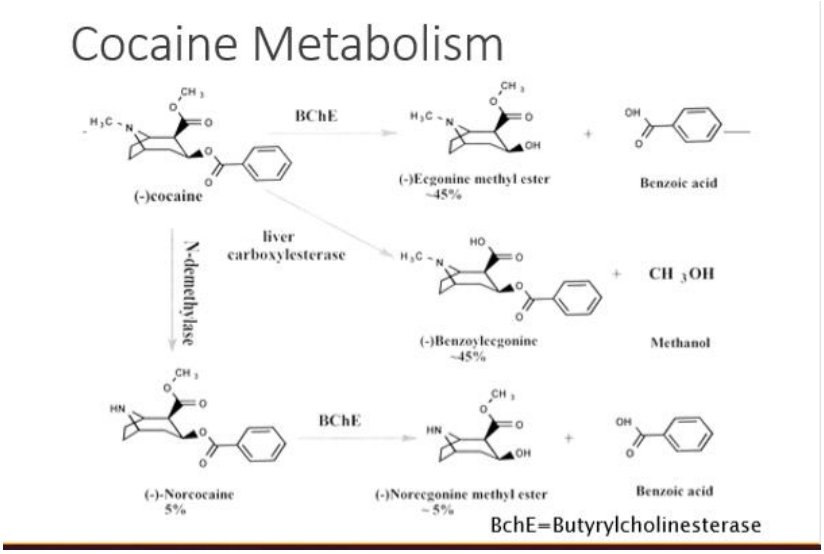
◦ Extensively metabolized

◦ Only 1% excreted unchanged in the urine

◦ Major metabolites – benzoylecgonine, ecgonine methyl ester

◦ Other metabolites - ecgonine

◦ Cocaine combines with ethanol in the liver to form norcocaine, cocaethylene, and norcocaethylene, pharmacologically active metabolites



Excretion :

➔Most drugs are eliminated either through the urine or through the feces

➔Some limited number of small volatile drugs can be eliminated via the breadth

➔To be excreted into the urine, drugs must be water soluble :
Drugs that are fat soluble are metabolized to water-soluble forms by the body prior to excretion in the urine

➔The relative importance of the routes of elimination vary depending on the drug

Implications :

◦ The rate at which a drug is metabolized and excreted will influence the half-life of the drug in the body
◦ Elimination rate is important in determining the length of time a person can be influenced by the drug:
◦ Cocaine has a half-life of 60 to 90 minutes ◦ THC has a half-life of 1.3 – 13 days

ADME Conclusions :

➔Understanding the time frame between administration and effect
➔Interpreting what tissue concentrations mean
➔Knowing what metabolites to look for
➔ Knowing what tissues samples to look at

Detecting and Measuring Drugs

Collecting sample for drug analysis

Sampling living individuals

1) Blood
2) Urine
3) Breath : Ethanol
4) Hair : Cocaine

Sampling deceased individuals : everything can be used, except the breath.

Extraction Techniques :

➔from biological tissues (aqueous solutions) into organic solvents before they can be analysed.

➔using pH we can extract acidic and basic drugs from the biological media

pH < 7 : acidic drugs
pH > 7 : basic drugs

The separate fractions are the concentrated and analyzed

<u>Measurement techniques :</u>

In most cases the identity of the drug is unknown. Screening assays are used first to get an initial assessment of what might be present.

1) Thin layer chromatography
2) Gas chromatography
3) Immunoassay

The screening assays are followed with confirmatory tests that are more accurate and specific to given dug

Gas chromatography/mass spectrometry (GS-MS) is the “Gold Standard”

Interpretation : what was the biological effect of the detected drug or toxin?

➔Direct cause of death
➔Indirect cause of death
➔ Influence on subject behavior
➔Placement at the scene of a crime
➔Example, carbon monoxide exposure in fire deaths
➔Absence of a therapeutic drug that may have influenced behavior Example, seizure or heart attack due to a missed pharmaceutical drug treatment

Challenges : help from other crime scene information to optimize analytical testing and to interpret toxicological findings

➔victim’s symptoms
➔post-mortem pathological findings
➔ examination of the victim’s personal effects including presence of drug containers or household chemicals

Ethanol

Widely used and abused drugs in Western countries

Jurisdictions have special laws regulating ethanol

For this reason, legal cases involving ethanol typically represent a significant percent of the Forensic Toxicologist’s workload

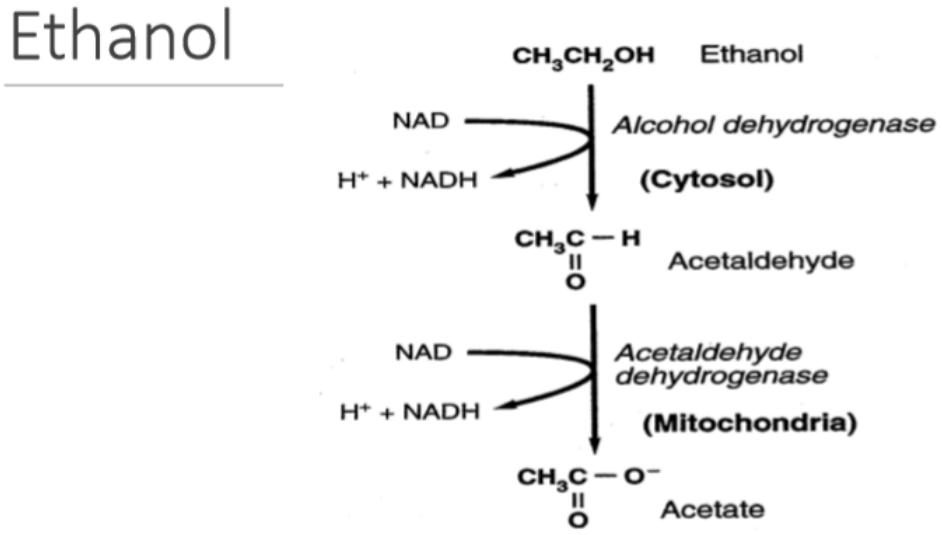
Ethanol ADME

Simple molecule, very water soluble, volatile
Readily absorbed in the gut with oral ingestion
Appears in the blood within minutes of ingestion

Once in blood stream, distributes evenly in the total body water

Metabolized largely in the liver (>90%)

Ultimately most – 95-98% - of the ingested ethanol is metabolized to carbon dioxide and water (generating energy in the process);
<u>a small percent is excreted intact in the breath</u>



Measurement of Ethanol Exposure

➔intact ethanol can be measured in the blood

➔excretion of intact ethanol in urine is small

➔excretion of intact ethanol in breath is small but measurable

➔metabolites are not helpful in detection or quantification

Blood Alcohol Concentration (BAC)

Metric of alcohol intoxication for legal or medical purposes

Blood alcohol content is usually expressed as a percentage of alcohol per volume in the blood

BAC of 0.10 means 0.10% (one tenth of one percent) of a person's blood, by volume, is alcohol

1% is equivalent to 1g/100 mL – 1g/dL
0.01% is equivalent to 10mg/100 mL

There is a very good relationship between concentration of alcohol in the blood and pharmacological effects

Supported by years of experimental data There is also a good relationship between ethanol concentrations in the blood and concentrations in the breath

The ratio of ethanol in the blood to ethanol in alveolar air is approximately 2,100 to 1 at 34 degree Celsius.

	Progressive effects of ethanol	
BAC (% by vol.)	Behavior	Impairment
0.010–0.029	• Average individual appears normal	• Subtle effects that can be detected with special tests
0.030–0.059	• Mild euphoria, relaxation, decreased inhibition	• Concentration
0.06–0.09	• Blunted feelings, disinhibition, extraversion	• Reasoning, depth perception, peripheral vision, glare recovery
0.10–0.19	• Over-expression, emotional swings	• Reflexes, reaction time, gross motor control, staggering, slurred speech
0.20–0.29	• Stupor, loss of understanding, impaired sensations	• Severe motor impairment, memory blackout, loss of consciousness
0.30–0.39	• Severe CNS depression, unconsciousness • Death is possible	• Bladder function, breathing, heart rate

Blood Alcohol Concentration

BAC is tied to the number of drinks and the time frame over which they were ingested ◦Also Influenced by:

Body weight; the higher the body weight, the lower the BAC per drink

Body fat, the higher the percent body fat, the higher the BAC per drink

Gender, Females have higher BAC for the same number of drinks as males of the same body weight

Food, the presence of food will slow the rate of absorption

Rate of elimination :

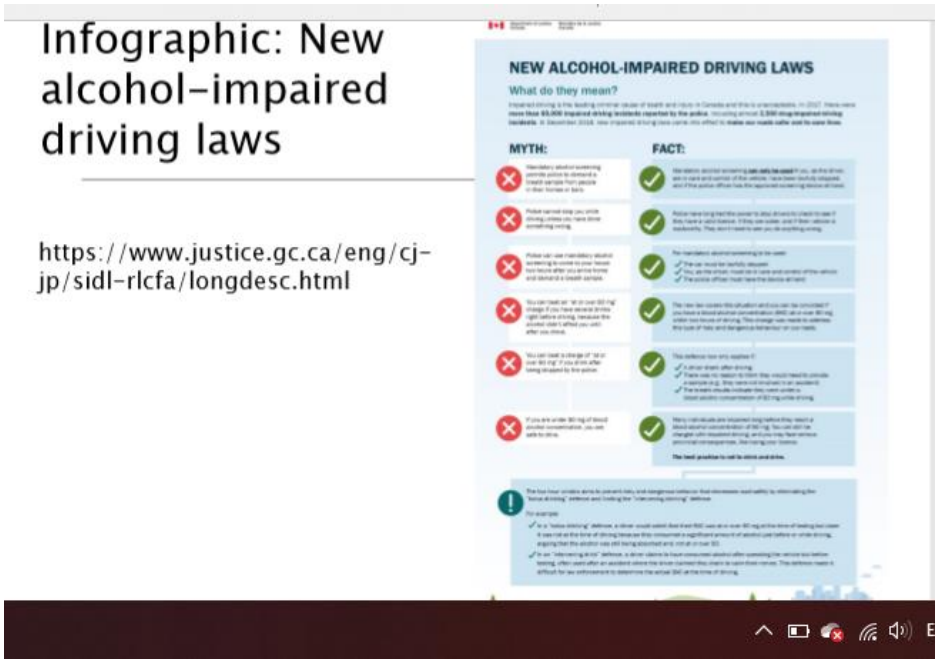
Ethanol is cleared from the system at an average rate of 0.015% (blood volume percent) per hour)

<u>Ethanol and the law</u>

For the purposes of law enforcement, BAC is used to define intoxication and provides a measure of impairment

In Ontario and the rest of Canada, the maximum legal BAC for fully licensed drivers is 80 milligrams of alcohol in 100 millilitres of blood (0.08%)

Driving with BAC in excess of 0.08 is a criminal offence



Breath testing for Ethanol

The amount of ethanol exhaled in the breath is in direct proportion to the concentration in the blood

A unique property of ethanol due to its volatility and distribution in total body water

Breath testers operate on the fact that at 34 C, the ratio of alcohol in the blood to alcohol in alveolar breath is approximately 2,100 to 1

Many types of breath testers are designed to analyze a set volume of breath

The captured breath is exposed to infrared light; the degree of interaction of the light with alcohol in the captured breath sample allows the instrument to measure ethanol

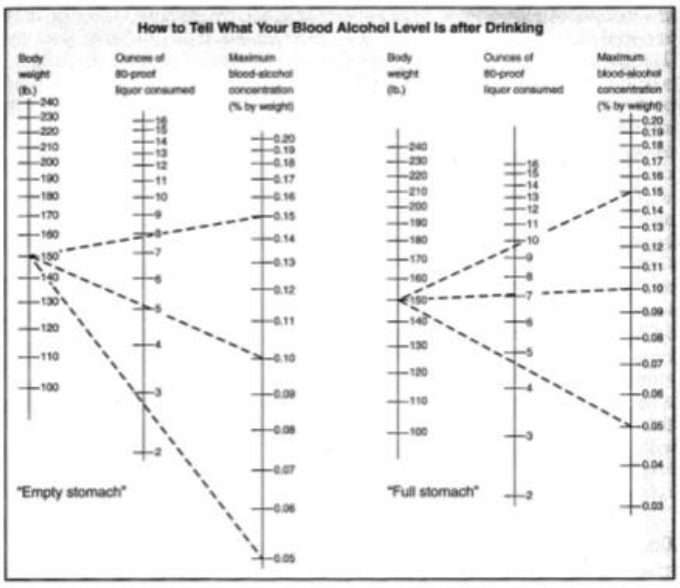
Measurement of Blood Alcohol Concentration

Gas chromatography offers the toxicologist the most widely used approach for determining alcohol levels in blood

Blood must always be drawn under medically accepted conditions by a qualified individual using a non-alcoholic disinfectant

Once blood is removed from an individual, its preservation is best ensured when it is sealed in an airtight container after an anticoagulant and a preservative have been added and stored in a refrigerator
Avoidance of loss of ethanol from samples due to volatility

Ethanol

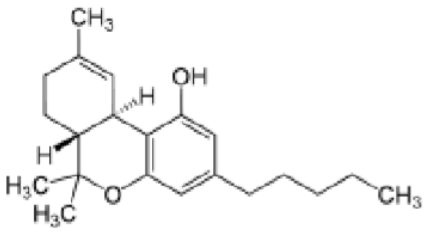


THC

Tetrahydrocannabinol (THC)

Tetrahydrocannabinol (THC)

- The principal psychoactive substance from the Cannabis plant



THC is very different from ethanol in its AADME characteristics

◦ THC is very lipid soluble

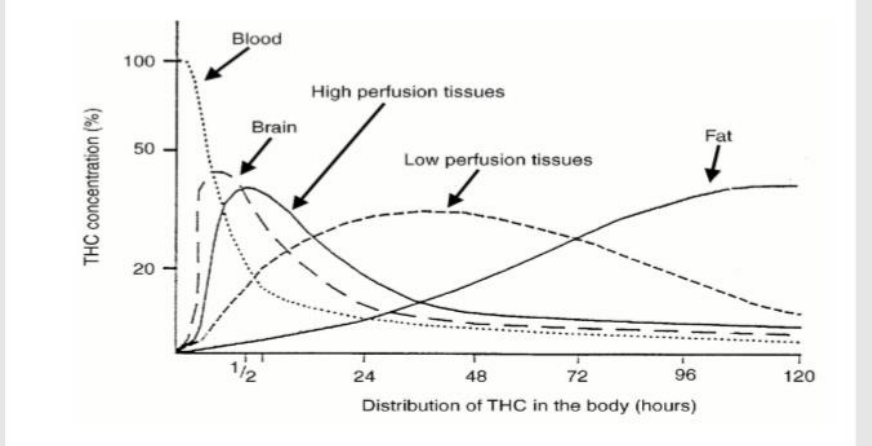
◦ It is not absorbed well from the GI tract - oral bioavailability 5-10%, and after oral ingestion there is a lag before pharmacological effects are seen

◦ Absorption from the lungs is very rapid, effects seen within minutes (serum levels peak within 10 minutes)

◦Highly concentrated in tissues over the blood, concentrates in fat
◦Drug can then be re-released from fat tissues over time

◦THC is metabolized to 11-hydroxy-THC which is also pharmacologically active

◦These combined characteristics result in a very long biological half-life
◦Tissue elimination is about 7 days, and complete elimination of a single dose may take up to 30 days



Detection of THC or 11-hydroxy-THC in urine is indicative of exposure but not necessarily of impairment

Actual psychological effects may dissipate within 3-5 hours despite the long half life in the system

The relationship between blood levels of THC and pharmacological effect is not well established

Debate within the scientific and legal community on relevance

Impact of THC levels on driving ability also debated

➔interpretation of forensic toxicology results for THC are not as straightforward as those for ethanol

Opioid Metabolism

Has a genetic component, ie allelic variation in cytochrome P450s that metabolize opioids

For some background:

A Z DePriest, B L Puet, A C Holt, A Roberts, E J Cone. Metabolism and Disposition of Prescription Opioids: A Review

Forensic Sci Rev 27 (2), 115-45 Jul 2015

Fentanyl

Synthetic opiate

Variable excretion rates

Data suggests this may be the result of differential distribution about the body

10% is cleared from the body through renal excretion after being acted on by a P450 the rest is liver-based

Mather Clin. Pharmacokinet. 1993 5:422

Updated review :

Kuip et al (2017) A Review of Factors Explaining Variability in Fentanyl Pharmacokinetics; Focus on Implications for Cancer Patients. Br J Clin Pharmacol, 83 (2), 294-313