

# Hallucinogens flashcard



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the word hallucinogen comes from the Latin word *\_\_alucinere*, meaning to wander in mind, talk idly, or prate

DSM5 defines a hallucinogen as a sensory perception that has the compelling sense of reality of a true perception but that occurs without external stimulation of the relevant sensory organ-

distortions of sensory information, some drugs cause mental confusion or delirium

other terms for hallucinogens: psychedelics, illusionogens, and deliriant

show many plants capable of producing hallucinogenic or deliriant effects

600 hallucinogens comprise a very chemically heterogeneous class of drugs

fourfold categorization comprised of what 3 categories based on neurotransmitter characterizations and the fourth being in a miscellaneous group

4 categories are:

1. cholinergic hallucinogens
2. serotonergic hallucinogens (eg. LSD)
3. methylated amphetamines (eg. MDMA, ecstasy)
4. other hallucinogens

another word for methylated amphetamines: noradrenergic hallucinogens

humans were quick to discover that ingestion of these plants could produce what-what acting in the cholinergic nervous system produces psychoactive effects: a delirium like state, or be used as poisons

drugs acting as agonists AND antagonists

cholinergic agonist hallucinogens

cholinergic nervous system importantly involved in

2 types of cholinergic agonist hallucinogens

these drugs stimulate primarily muscarinic cholinergic receptors, producing greater than normal neural activity in the cholinergic nervous system

many life sustaining physiological functions and learning & memory

amanita muscaria, ibogaine

amanita muscaria-fly agaric mushroom-mushrooms have a bright red cap, speckled with white dots, similar to the ones in the Disney film Fantasia-not a typical drug choice today Vikings called “ beserkers” because ingestion of the mushroom produces a state of “ agitated raving” and feelings of invincibility, evidence that Vikings would consume this before raiding a village amanita contains what psychoactive substance (2)

these substances are excreted where

ibotenic acid\* (which is metabolized to muscimol) and muscimol

excreted in urine largely unchanged and an active dose may be had by ingesting the urine of someone who ingested the mushroom

symptoms of cholinergic activation from amanita include-initial period of good humour and light euphoria-feelings of detachment and unreality-feelings of increased power-agitated raving-twitching and trembling of limbs-visions of the supernatural and illusions of grandeur-marked lacrimation (tearing)-salivation and sweating-pinpoint pupils (shrink up)-severe stomach pain, and painful diarrhea, coma, convulsions, and potentially death ibogaine-plant found in Gabon in central Africa, Gabonese hunters chewed the yellowish root of the plant-mainly acts as a stimulant allowing the hunters to endure long treks and searches for food main psychoactive compound in ibogaine ibotenic acid ibogaine was sold when/where/why

is ibogaine legal in Canada

in the 1960s in France as an over the counter medication for fatigue

yes!

some people found that if enough pills were taken they could experience a type of mystical or meditative effect during which what they identified as therapeutic, repressed childhood memories were unlocked, users felt that these experiences gave them great insight in themselves. Ibogaine gained a reputation among heroin users as useful if they were trying to quit. There are drug treatment centers in Canada that do what, ex. incorporate ibogaine in a treatment plan. Toronto Ibogaine Treatment Center. Cholinergic antagonist hallucinogens

4 types

some substances block muscarinic cholinergic receptors and in doing so produce a set of effects that justify their classification as deliriants or hallucinogens. Common in pesticides

-atropa belladonna-datura stramonium-henbane-mandragora officinarum

atropa belladonna

main ingredient

atropine is an alkaloid of atropa belladonna, known as a deadly nightshade or love apples, b/c it was thought to be an aphrodisiac (drug that stimulates sexual desire)

main ingredient is atropine

atropa plant is found mainly where, and is a member of what family, produces what kind of flowersmainly Europe, North Africa, Asia member of the tomato/potato family produces bell shaped purplish flowers tinged with green and has soft blue/black berriesname atropa belladonna was given to reflect 2 common uses of the plant-atropos eldest of the 3 fates in greek mythology, and it was her duty to cut the thread of life for a person's death (thus atropa reflects the use of atropine as a poison)

-belladonna comes from when women would instill the juice of nightshade berries in their eyes to dilate them, which was considered to be more beautiful

physiological effects of atropinedilated pupilsfeeling of suffocatinghusky voiceat low dosesdrug produces an initial increase in general arousalat slightly higher dosesedative effects begin to appear, dream like state where user experiences pleasing hallucinations, vivid and erotic dreams, and a sense of weightlessness (sensation of flying)upon awakeninguser may exhibit amnesia (has difficult time realizing dreams and hallucinations were not real)at even higher dosesphase of pleasantness is replaced with more powerful and frightening images, potentially lethal-very fine line btwn pleasant experience and lethal one, so repeated recreational use very uncommondatura stramonium3 other common namesatropine and other cholinergic blockers, scopolamine and hyoscyamine, are found in Datura stramonium, known as jimsonweed, jamestown weed, thorn apple datura history of usein colonial america women who claimed to witches would rub ointments made from Datura on sticks and then straddle them for feeling of weightlessness and a sensation of flying-drug would be absorbed via the

vaginal membranes producing a claim that they were flying datura has been and continues to be used for what to fortify marijuana in the far east and India today in these regions (Far East and India), what happened cigarettes are made from the leaves of Datura and Atropa and smoked to produce an intoxicating effect until 1970s, these cigarettes were sold in pharmacies in both Canada and the US for the treatment of asthma b/c of the bronchodilating effects of atropine ingestion of datura produces same effects as those of atropa Jimsonweed is cultivated today for what

how many seeds for a low to moderate dose, how many for hallucinations

decoration-has purple trumpet like flowers, large, jagged leaves, and thorny fruit pods that produce seeds each fall

10-20 for low to moderate dose 50 for hallucinations/death

henbane-contains scopolamine and hyoscyamine-called hyoscyamus niger-yellowish flowers and hairy leaves mandragora officinarum mandrake (meaning “potent male”) plant contains atropine, scopolamine, and hyoscyamine history, myth a hanged man would have an erection and ejaculation caused by the sudden snapping of the neck sending an impulse down the spinal cord, plant grew wherever a hanged man’s semen fell to the ground-myth also has it that the plant shrieks when uprooted, driving mad anyone who hears it at low doses at higher doses it is used as a sedative hypnotic type of drug at higher doses, it produces hallucinations and delirium the drug most identified with the term “hallucinogen” is the prime example of a serotonergic hallucinogen serotonergic hallucinogens often used

in what migraine medications  
LSD, chemical name and trade name  
first synthesized when and by who

important symptom in LSD

Lysergic acid diethylamide, trade name Delysid  
first synthesized in 1938 by the Swiss chemist Albert Hoffman who was studying derivatives of the ergot fungus for their vasoconstrictive action and ability to give muscle tone to the uterus

-LSD one of the most potent drugs around-  
important symptom is synesthesia

lysergic acid amide is an alkaloid obtained from ergot, a parasitic fungus found on a variety of grains (eg. rye and wheat)  
Hoffman was not attempting to produce an hallucinogenic drug, but it turned out that what

his research also produced

the 25th derivative/modification that he produced had strong hallucinogenic properties (hence the name LSD-25)

a number of compounds with therapeutic usefulness (eg. Methergine), drug used to treat migraines

did this happen on purpose or by accident, what year

next day what happened

1943 Hoffman accidentally dosed himself with LSD by getting a sample on his hands, recorded a detailed picture when he started experiencing effects

he intentionally ingested 250 micrograms to study the drug's effect

250 micrograms 5 times larger than the minimal psychoactive dose-minimal dose is about the size of a grain of salt Dr. Humphrey Osmond coined the term "psychedelic" gov't agencies in the US were interested if LSD would be useful as an aid in interrogations, but concluded that it was too unreliable to use Dr. Timothy Leary-godfather of LSD-started freely distributing the drug and told people to "turn on" to acid LSD used in psychotherapy when and why during 1950s and 1960s, rationale being that while under the influence of the drug the user would more openly communicate about inner feeling and also be more open to therapeutic suggestions LSD legal in the US until when 1966 in 1966 what happened to LSD it was made a schedule I drug in the US (no medical use and high abuse potential) use of LSD peaked when late 1960's to early 1970's from 1967-1971 the % of college students reporting at least one lifetime use of LSD rose from 1%-20%-however use has declined since that time Canadians aged 15 or older reported lifetime use for hallucinogens of about what % and past year use at what % lifetime use at 12%, past year use at 1% among university Canadian students, past year use at what % 6% since 1995, steady decline since the peak in 1995 LSD characteristics odorless, tasteless, colorless, and fairly easily crosses both the blood brain barrier and the placental barrier how is LSD typically taken orally as a tablet (eg. orange barrel, purple haze), gelatin capsule (commonly called window pane), or on absorbent paper (commonly called blotter acid) oral ingestion peak concentration how long, duration of effects 90 minutes peak concentrations 5-12 hrs duration of effects first symptoms of LSD activation of the sympathetic nervous system: dilated pupils (good index



of the hallucinogenic effect), increased body temperature and blood pressure, analgesia (inability to feel pain) after 1-2 hours of ingestion

major symptoms (8)

-hallucinogenic effects –primary visual and often experienced with the eyes closed

-one of the first effects is wave like and rhythmic movement in objects (that are not actually moving)-object trails-when eyes are closed there appear images of latticework, spirals, funnels, cobweb figures (called form constants, occurs frequently)-synesthesia: perception of one sense in another sense modality (eg. users seeing sounds, can see vividly coloured musical notes while listening to music)-distortions in perception, small periods of time seem to be very long-ego disintegration (difficult to distinguish himself from the surroundings), effect may be partially due to the anesthetic effects of the drug. One user described the effects as if their whole body had been injected with novocaine, and it was like touching your jaw at the dentist-bad trips –potential adverse effect, common in novice users, fear that effect is permanent (3%)-flashbacks: unexpected psychedelic experiences long after the most recent use of LSD, typically short lived, self terminating, and not distressing

HPPD, hallucinogen persisting perception disorder more long lasting, distressing, recurrent, and only slowly reversible psychedelic experiences occurring well after the last drug use (4% of users)-recognized diagnosable condition in DSM 5 LSD fatality rate LD50 is 14 000 micrograms which is approx 300 times the minimal psychoactive dose estimated at 50

micrograms, very difficult to OD does tolerance occur tolerance occurs to most of the hallucinogenic effects of LSD in 3-4 days-tolerance also occurs to other serotonergic hallucinogens (cross tolerance across these hallucinogens)-if tolerance has developed to LSD there will be tolerance to psilocybin or mescaline what also contributes to the development of tolerance to LSD Pavlovian conditioning is there evidence of physical dependence no does LSD cause harmful effects in babies LSD does not cause chromosomal damage (brain damage) has research supported an LSD model of schizophrenia no in schizophrenia there are true hallucinations experienced with eyes opened, in LSD produces perceptual distortions with eyes closed early research suggested that LSD suppressed firing of (Reichs and Rosecrans) serotonin neurons in the raphe nuclei by acting as an agonist at presynaptic 5-HT<sub>1</sub> receptors which serve a negative feedback regulatory function on serotonergic activity the raphe nuclei are part of what is called the ascending reticular activating system (ARAS) which is involved in the filtering of sensory information, and thus the hypothesis was that LSD interfered with this filtering leading to sensory distortions some research contradicted this explanation and a consensus grew that LSD produced its effects how mainly via an agonistic action at postsynaptic 5-HT<sub>2</sub> receptors blocking 5-HT<sub>2</sub> receptors does what antagonizes the hallucinogenic effects of LSD although it is believed that agonism at 5-HT<sub>2</sub> receptors is critically involved in LSD's hallucinogenic effects there is new evidence that the original mechanism of agonism at 5-HT<sub>1</sub> receptors suppressing raphe activity is relevant lysergic acid amide less potent, naturally occurring form of hallucinogen related to LSD-found in ergot, a fungus present on morning glory seeds-looks like rooster leg these naturally occurring sources

of lysergic acid played a role in the Salem witch trials—witch trials took place in fall of 1691 following a warm, rainy spring and summer that would have been favourable to an ergot infestation. “Holyfire” or “St. Anthony’s fire” may have resulted from the strong vasoconstrictive action of lysergic amide that had been baked into bread made from infected grains—the restriction of blood flow to the limbs caused by the vasoconstriction leads to a sensation of warmth (hence the fire), tingling sensation also a symptom of ergotism, when and where afflicted behaviour caused by ingestion of ergot—outbreak in a small French town in 1951. Other hallucinogens that resemble the neurotransmitter serotonin (3) psilocybin, dimethyltryptamine, bufotenine, psilocybin, common street name chemical name what is it Aztec and Mayan people called psilocybe mushrooms what how is it typically taken how much needed for psychoactive effect, how much for hallucinogenic effect onset of effects is how long and duration is how long “shrooms” common street name 4-phosphoryl-dimethyltryptamine—naturally occurring substance in a variety of mushrooms—teonanactl, meaning “flesh of the gods”—typically taken by eating mushrooms or drinking a brew containing them 4–8 mg, > 15 mg for hallucinogenic effect onset of effects is 30 minutes with a duration of 2–6 hrs in the body, psilocybin is converted to does tolerance occur psilocybin effects psilocin which is more lipid soluble and is thought to be the actual active agent—tolerance occurs to the effects of psilocybin and cross tolerance is displayed to other hallucinogens compared to LSD, more strongly visual, less emotionally intense, more euphoric, and less likely to produce a panic reaction dimethyltryptamine (DMT)—what does it look like, effects, onset, duration—reddish bark used as snuff—effects are rapid and short—onset as little as 10 seconds, duration of 60 min at most because of

short duration, it became known as the businessman's psychedelic (would take over lunch hour), ppl on Wall street used this to trade stocks methods of ingestion, most common method one of most common drinks used by indigenous people effects of DMT may be taken as drink, but highly ineffective b/c it is rapidly metabolized by monoamine oxidase\*-ayahuasca\* (DMT + additional substance), used in religious ceremonies and PTSD-most common method of administration is inhalation/smoking-effects: excitability, other worldly experience, numbness of the limbs, twitching of the facial muscles, nausea bufotenine chemical name found in what animal side effect unique 5-hydroxy-DMT used as a hallucinogen by indigenous peoples in the form of snuff (yopa and cohoba) found in toads, may have something to do with the use of toads in witch's brews?, also used in Simpson's toad cyanosis, skin turns purplish blue is there evidence that it produces self administration, cyp, little evidence in serotonergic hallucinogens for both-not reinforcing at all neurotransmitter norepinephrine have what kind of properties hallucinogenic properties neurotransmitter amphetamine has what kind of properties combination of stimulant and hallucinogenic properties prime example of serotogenic hallucinogens and norepinephrine hallucinogens serotogenic -> LSD norepinephrine -> mescaline mescaline the prototypical naturally occurring norepinephrine type hallucinogen mescaline is the active alkaloid in what the peyote cactus contains stimulating and rewarding properties characteristics of the cactus small, spineless green grey pin cushion\* or button looks like a pin cushion most of the plant is underneath, the important part is above when used in the natural condition...in this form, the active ingredients remain the button is sliced and often dried to form hard brownish discs they remain potent indefinitely is mescaline absorbed

readily/passes easily? mescaline is absorbed readily from the digestive tract but passes poorly through the blood brain barrier how much needed to produce euphoric effects and how much for hallucinogenic 3 mg for euphoric/stimulant effects 5 mg for hallucinogenic effects peak effects how long, effects last how long, normal duration of effect peak effects occur in about 60 min, effects may last 4-16 hours but the avg is about 10 hrs-effects are long lived peyote intoxication characterized by coloured visions in kaleidoscopic movement, often accompanied by auditory, taste, olfactory, and tactile hallucinations-user experiences sense of weightlessness, alterations in time perception-nausea, vomiting, headaches, and hangover what 3 drugs are indistinguishable mescaline, psilocybin, and LSD since it is related to norepinephrine, mescaline produces sympathetic arousal including dilated pupils, increased heart rate, blood pressure, and body temperature does tolerance develop is there cross tolerance yes, occurs rapidly yes, cross tolerance to other hallucinogens although mescaline resembles the neurotransmitter norepinephrine it seems that it produces hallucinogenic effects in essentially the same manner as the serotonergic hallucinogens, that being an agonist action at 5-HT<sub>2</sub> receptors -> basis of cross tolerance past year use of mushrooms and mescaline in grades 9-12 to be what % higher use in females or males the peak past year of mushrooms/mescaline occurred when and was at what % 4% higher use in males than females peak past year use occurred in late 1990s/early 2000s and was about 17% common methylated amphetamines (4)-DOM or STP-MDA-MDMA-Myristicin and elemicin DOM, street name first used as STP potential appetite suppressant STP stands for “serenity, tranquility, and peace” or “super terrific psychedelic” doses greater than 10 mg produces

strong hallucinations lasting 16-25 hours DOM affects what (2 things) serotonin and dopamine DOM is associated with what due to what adverse reactions/bad trips more than any other hallucinogens due to the long duration of action and the difficulty convincing the panicked users that the effects will wear off 2 related compounds of DOM are DOI and DOB MDA, most common street name first synthesized when and as what EVE 1910, as an appetite suppressant, antidepressant, and in the treatment for Parkinson's disease MDA is more like a what a typical hallucinogen as compared to an amphetamine like drug MDA is a metabolite of MDMA (ecstasy) and my account for many of MDMA's effects-MDMA is metabolized to MDAMDA effects enhances emotions and empathy and tends to promote a strong emotional link MDA and MDMA are useful psychotherapeutic adjunct b/c it allows people to get in touch with their inner feelings root of the sassafras tree formerly used as what contains what that can be converted to what formerly used in the production of root beer but now banned b/c of carcinogenic properties, contains a substance, safrole or shikimo, that can easily be converted to MDAMDMA, commonly known as what was synthesized when and as what 3, 4-methylenedioxymethamphetamine, ecstasy or XTC synthesized in the early 1900s as a potential appetite suppressant onset of effects occur when, duration of effects when within one hour, duration of effects around 4-6 hours symptoms positive mood changes, increased energy, and higher dose hallucinations-undesirable effects include sweating, tension in the jaw, teeth grinding MMDMA has more toxic effects than most other hallucinogens b/c it produces very high body temperatures (malignant hyperthermia) and dehydration, a combination that has produced some deaths speak use when, it

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was legal until when what % use amongst students peak use mid 1980s, drug was legal until 1986 3% MDMA causes release of what what is better at releasing the others serotonin and dopamine and is particularly effective in releasing serotonin, cocaine better at releasing dopamine MDMA and MDA seem to block what the reuptake of serotonin thereby causing a prolonged and intense period of serotonergic activation can this cause permanent damage yes, excess activation of serotonin and dopamine can cause permanent damage to these neurotransmitter systems MDMA has been shown to be a potent serotonergic neurotoxin in what animals rats, nonhuman primates, even evidence from humans when people had used MDMA between 80-100 times compared to people who had not used the drug, what happened the major serotonin metabolite was lower in people who had 50 or more lifetime use there was reduced serotonin binding and transporter densities, effects more pronounced in women in brain areas involving learning and memory damaged nerve endings do not regrow which results in a permanent lowering of serotonergic functioning, underfunctioning in brain areas involving sleep and appetite there is excess regeneration resulting in excess serotonin release chronic use of MDMA, problems include memory impairment, decision making deficits, loss of impulse control, recurrent paranoia, and depression Myristicin and elemicin, found where what spices are derived are they strong or weak taken in the form of what initial effects include what, duration lasts how long found in the fruit of the tree *Myristica fragrans*-nutmeg and mace are spices derived from the dried seed and seed coat-fairly weak hallucinogens-tea-nausea and vomiting, after 2 hours u get a weak hallucinogenic effect mescaline show self administration and cyp? mescaline not self administered by monkeys-difficult

to obtain self administration with this prototypical phenethylamine  
hallucinogen-little evidence that it produces CPP too-these finding similar to  
those with LSD in comparison, many of the methylated amphetamine type  
hallucinogens, notably MDMA and MDA are self administered and produce  
CPP miscellaneous hallucinogens 4 types-a variety of other drugs that produce  
hallucinogenic, psychedelic, deliriant or dissociative experiences-  
phencyclidine and ketamine-dextromethorphan-salvia divinorum-  
thujone phencyclidine and ketamine-called dissociative anesthetics-produce  
total anesthesia, but at lower doses they produce a feeling of detachment  
(dissociation) from the environment and self, a type of out of body/out of  
world experience-drugs also produce analgesia and amnesia Phencyclidine  
developed when, marketed when under what trade name it had the desirable  
effects as what but also what kind of undesirable side effects ketamine was  
developed when developed in 1926 and marketed until the 1950s under the  
trade name Sernyl desirable effects as an anesthetic, undesirable effects of  
hallucinations and seizures and was removed from human use in 1960s as a  
drug with potentially fewer unwanted side effects most common street  
names for phencyclidine angel dust PCP, angel dust, and horse tranks  
(reference to the drug's use in vet medicine as a tranquilizer)-> sprinkle on  
marijuana to increase potency PCP may be taken how peak concentrations  
occur when vs. peak concentrations if smoked orally in the form of a powder  
or liquid orally takes 2 hrs, smoking takes few seconds effects include  
effects last how long but may persist for how long  
euphoria, numbness (due to anesthetic effect), loss of motor coordination,  
catatonia (not moving, waxy flexibility), initial nystagmus (eyes moving back  
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and forth quickly) but eventually a fixed stare (doll's eyes), distortion of body image with the perception of parts the body, auditory hallucinations, extreme mood changes going from almost no emotional affect to outbursts, aggressive hostility, and stereotypies, violent and aggressive behaviour, out of control

effects last 4-6 hours but may persist for days

lethal dose how much are bad trips common 2-5 times the recreational dose occur frequently, in 80% of users does tolerance and physical dependence develop yes in lab animals, not really in humans b/c of infrequency with which humans subject themselves to sufficient dosing withdrawal symptoms in rhesus monkeys include tremors, oculomotor hyperactivity, bruxism, fearfulness, vocalizations, diarrhea, emesis and convulsions Ketamine, street name, popular where effects compared to PCP Special K, popular drug at rave similar but generally shorter, lasting 2 hours users often describe the effects of ketamine by saying they have entered the "K-hole" suggested that ketamine may provide a more useful model of psychosis and schizophrenia PCP and K bind to what sigma opiate receptors and antagonistic NMDA glutamate receptors-PCP and K act as "use dependent" antagonists when glutamate acts to open an ion channel PCP or K attaches to a site within the channel, PCP and K would thus suppress the level of neuronal activity produced by glutamate which is the major excitatory neurotransmitter in the CNS the antagonism of the NMDA receptor is thought to also produce an increase in dopaminergic activity in brain reward centers both K and PCP are self administered by do they support CPP Rats, monkeys, and dogs do not support CPP PCP and K have reinforcing

and aversion (conditioned place aversion) effects  
Dextromethorphan, common use, is it more or less potent than PCP/KDM is a common cough suppressant which stimulates sigma opiate receptors and blocks NMDA receptors, the same two actions of PCP and K, cough syrup, less potent  
what else is a really good cough suppressant  
opiates  
DM abuse occurs with who young drug experimenters  
recreational use of DM is referred to as robo-copping, roboing, robo-tripping, b/c Robitussin is one of the main sources of abuse  
medical dose is what, how much needed to produce an euphoric effect  
15-30 mg, euphoric effect needs 200 mg  
doses at 400 mg and 600 mg produce what  
400 produce more intense euphoria, vivid, imagination, closed eye hallucinations  
600 produces strong alterations in consciousness, out of body experiences, and psychotic like reactions  
also some recreational use with over the counter medications, 2 examples  
benadryl (cold medication)  
Gravol (motion sickness)-if you take enough Gravol you will experience a hallucinogenic effect  
Salvia divinorum, member of what, also called what, is it illegal  
member of the mint family, also called Diviner's Sage, no it's legal  
naturally occurring plant substances that are used to produce mystical or religious experience are now commonly called entheogens  
most common method of ingestion is chewing a quid (not effective), smoking a cigarette, or making a tea  
it has a potency roughly equal to what  
psilocybin containing mushrooms  
active ingredient is what  
salvinorin A, shown to exert an agonistic action at kappa opioid receptors, produces a hallucinatory effect  
use of pure salvinorin A is reported to produce what  
very strong effects, to the point that most users have no desire to use it a second time  
salvia produces CPP in what and CPA in what  
CPP in zebra fish and CPA in rats  
salvia controlled by what, sold where  
controlled by Health Canada, sold in

convenience store thujone, where is it found a variety of plants including sage, but most notable in wormwood wormwood was used to produce what became popular where and when the alcoholic drink Absinthe (60-80% alc content) became popular in France in the mid 1800s heavy consumers of Absinthe were noted to do what act strangely and report hearing voices and seeing things thujone is what an GABA antagonist, and since GABA is an inhibitory neurotransmitter blocking its action increases neural activity which may produce convulsions banned in NA when, but now what 1910, but now possible to obtain Absinthe that has a controlled amount of thujone Absinthe sometimes called what Green Fairy b/c of green liquid common method of taking Absinthe is to do what pour the green liquid into a small, tulip shaped glass, then place a special slotted spoon on the glass and put a sugar cube that has been soaked in Absinthe on it. sugar cube is ignited, burned for awhile and then dropped into the Absinthe and finally water is poured in to douse the flames. showed in what movie Johnny Depp from Hell, he takes opium and Absinthe Oscar Wilde characterizes how people feel about Green Fairy after 1st glass, you see things as you wish they were after 2nd, you see them as they are not after 3rd, you see them as they really are, which is the most horrible thing in the world