## **Psychiatry Medications**

Drug	Indication	MOA	Side Effects	Notes
		Antipsych	otics	
<b>1<sup>st</sup> generation</b> - Chlorpromazine - Haloperidol.	<ul> <li>Functional psychosis:</li> <li>Schizoaffective disorders.</li> <li>Schizoaffective disorders.</li> <li>Schizophreniform disorder.</li> <li>Brief psychotic disorder.</li> <li>Mania (Chlorpromazine " sedative").</li> <li>Postpartum psychosis.</li> <li>Psychosis with depressed mood.</li> <li>Delusional disorders.</li> <li>Delirium.</li> <li>Resistant schizophrenia (Clozapine).</li> <li>Bipolar Mood Disorder.</li> <li>Organic psychosis:</li> <li>Psychosis induced by meds (corticosteroids).</li> <li>Substance abuse.</li> <li>Delirium and dementia.</li> <li>Violence/aggression, agitation, and excitement.</li> <li>Agitation induced by alcohol intoxication.</li> </ul>	<ul> <li>High blockade of dopamine receptors type 2 (D2) in the mesolimbic pathway → reduces active psychotic features.</li> </ul>	<ul> <li>Antidopaminergic:</li> <li>Nigrostriatal tract → EPSE (Acute dystonia, parkinsonism, akathisia, rabbit syndrome, tardive dyskinesia) → treat with anticholinergic.</li> <li>Tuberoinfundibular tract → hyperprolactinemia (gynecomastia and amenorrhea).</li> <li>Mesocortical tract → lack of motivation, ↓ concentration.</li> <li>Anticholinergic:</li> <li>Dry mouth, constipation, urinary retention, poor erection, blurred vision, closed-angle glaucoma.</li> <li>Antiadrenergic:</li> <li>Postural hypotension.</li> <li>Inhibition of ejaculation.</li> <li>Antihistaminergic:</li> <li>Sedation and weight gain.</li> <li>Prolonged QT In ECG.</li> </ul>	<ul> <li>Don't treat amenorrhea with dopaminergic medications (Bromocriptine) in psychotic females.</li> <li>Quetiapine = Atypical antipsychotic has no effect on prolactin → doesn't cause amenorrhea</li> <li>Neuroleptic Malignant Syndrome → reaction to antidopaminergics</li> <li>Features: Muscle rigidity, sweating, hyperthermia, akinesia, clouding consciousness.</li> <li>Labs → ↑ CPK, ↑ potassium, neutrophilia.</li> </ul>
2 <sup>nd</sup> Generation - Olanzapine - Quetiapine - Clozapine - Risperidone - Paliperidone		<ul> <li>Blockade of 5HT A2 in the mesocortical tract → ↑ dopamine function (5HT inhibits dopamine) → improve negative symptoms of psychosis: ↓ initiation, lack of motivation.</li> </ul>	<ul> <li>Mesolimbic &gt; Nigrostriatal → Less EPSE, antiadrenergic, anticholinergic.</li> <li>High risk of metabolic syndrome.</li> <li>Agranulocytosis (Clozapine).</li> <li>Hyperprolactinemia (Risperidone).</li> </ul>	<ul> <li>Treatment Medical ICU". bromocriptine to reduce the risk of acute renal failure tha may result from ↑ myoglobin</li> </ul>
<b>3<sup>rd</sup> Generation.</b> Dopamine System Stabilizers • Aripiprazole	<ul> <li>Amphetamine intoxication (Olanzapine)</li> </ul>	<ul> <li>Partial D2 agonist; in mesolimbic it competes with dopamine → less active symptoms</li> </ul>	<ul> <li>Agitation, anxiety, insomnia, dyspepsia, nausea, seizures.</li> </ul>	<ul> <li>Doesn't increase weight and is usually non-sedating.</li> </ul>
DEPOT	- Poor compliance patients.	-	-	- IM injections , (slow release)

Drug	Indication	MOA	Side Effects	Notes
Electroconvulsive therapy "ECT"	<ul> <li>Schizophrenia (catatonic, resistant to drugs).</li> <li>Depressive disorder with suicidal risk or delusions.</li> <li>Depressive stupor or marked retardation.</li> <li>1<sup>st</sup> trimester of pregnancy.</li> <li>Elderly.</li> <li>Renal failure.</li> <li>Postpartum psychosis.</li> <li>Schizoaffective disorder.</li> <li>Mania &amp; mixed affective states.</li> </ul>	<ul> <li>ECT Procedure:</li> <li>Bilateral (most common) → fronto-temporal position.</li> <li>Unilateral → less memory impairment but less effective.</li> </ul>	<ul> <li>Headache (due to ↑ ICP).</li> <li>Body aches and myalgias (due to muscle contraction)</li> <li>Memory impairment (due to neuronal hypoxia during seizure).</li> <li>Mania in susceptible depressed patients.</li> <li>Bone fracture and tongue or lip injury.</li> </ul>	<ul> <li>Psychiatric disorders that may show deterioration or no response to ECT:</li> <li>Phobic disorders.</li> <li>Conversion disorder.</li> <li>Primary hypochondriasis</li> <li>Depersonalization disorder.</li> <li>Contraindications:</li> <li>↑ ICP (Absolute)</li> <li>Cardiac infarct &lt; 3 months.</li> <li>Arrhythmias.</li> <li>Hx of cerebral infarction.</li> <li>Brain tumor.</li> </ul>
		ANTIDEPRES	SANTS	
SSRIs: - Paroxetine - Fluoxetine - Citalopram - Escitalopram - Sertraline - Fluvoxamine	<ul> <li>Depressive disorders.</li> <li>GAD → Fluoxetine, Venlafaxine.</li> <li>Social phobia → Fluoxetine, Venlafaxine.</li> <li>Panic disorders → Paroxetine.</li> <li>OCDs.</li> </ul>	- Selectively inhibit serotonin reuptake into presynaptic neuron.	<ul> <li>Sexual dysfunction (delayed orgasm).</li> <li>Insomnia (Fluoxetine).</li> <li>Sedation (Fluoxamine).</li> <li>Prolong bleeding time by inhibiting platelet aggregation → ↑ risk of stroke.</li> <li>Hyponatremia (paroxetine)</li> <li>Withdrawal syndrome (paroxetine).</li> <li>Serotonin syndrome; due drugs combination with MOAIs (which inhibit the catabolism of serotonin) Features → myoclonus, nystagmus, tremor, and hyperpyrexia.</li> </ul>	<ul> <li>Relatively safe in overdose.</li> <li>Treatment of OCD often requires high doses of SSRIs.</li> <li>Play good role in controlling blood sugar in DM.</li> <li>Citalopram = effective in Post-Stroke depression but may increase the bleeding tendency.</li> </ul>
<b>SNRIs:</b> - Venlafaxine. - Desvenlafaxine - Duloxetine.	<ul> <li>Trichotillomania.</li> <li>Tic disorders.</li> <li>Premature ejaculation (SSRI).</li> <li>Post stroke depression (Citalopram).</li> </ul>	- Selective-Serotonin-Norepinep hrine Reuptake Inhibitors	<ul> <li>Dry mouth, anorexia, asthenia, blurred vision,</li> <li>Abnormal ejaculation or orgasm.</li> <li>Erectile disturbances.</li> <li>Sweating</li> <li>↑ BP (Venlafaxine &gt; 225 mg).</li> </ul>	<ul> <li>Higher rates of remission.</li> <li>Desvenlafaxine has fewer side effects.</li> <li>Duloxetine has positive benefits in neuropathy in DM.</li> </ul>

Drug	Indication	MOA	Side Effects	Notes
Mirtazapine	- Depression + insomnia (highly sedating).	- ↑ NE and 5HT.	<ul> <li>↑ Appetite.</li> <li>Weight gain.</li> <li>Sedation.</li> </ul>	- Combined with SSRIs or venlafaxine to counteract serotonergic side effects (nausea, agitation & insomnia).
Bupropion	- Antidepressant.	- Norepinephrine and dopamine reuptake inhibitor.	<ul> <li>Dry mouth, constipation.</li> <li>Weight loss.</li> <li>HTN.</li> </ul>	- Combined with SSRIs or venlafaxine to counteract sexual side effects (delayed ejaculation), sedation, weight gain.
<b>TCAs:</b> - Amitriptyline. - Imipramine. - Clomipramine.	<ul> <li>Depressive disorders.</li> <li>Anxiety, phobic.</li> <li>Panic disorders.</li> <li>OCDs (Clomipramine).</li> <li>Nocturnal enuresis (imipramine).</li> <li>Depression in RA patient (Amitriptyline).</li> <li>Pruritus (H1 blockade e.g. doxepin).</li> <li>Gastric ulcer (H2 blockade e.g. amitriptyline).</li> </ul>	- Tricyclic Antidepressants	<ul> <li>Anticholinergic:, urinary retention, impaired visual accommodation, glaucoma. Delirium.</li> <li>Antiadrenergic: Postural hypotension, delayed ejaculation and drowsiness.</li> <li>Sweating.weight gain, arrhythmia, tremor.</li> <li>Mania in susceptible patients.</li> </ul>	<ul> <li>If a patient has insomnia,(amitriptyline or doxepin)</li> <li>Tricyclics are avoided in suicidal patients and cardiac patients.</li> <li>Can be given in the 2<sup>nd</sup> and 3<sup>rd</sup> trimester of pregnancy.</li> </ul>
Monoamine Oxidase Inhibitors (MAOIs)	<ul> <li>Patients who have not responded to other antidepressants.</li> <li>Atypical depression</li> <li>Phobic and panic disorders.</li> <li>Narcolepsy.</li> </ul>	-	<ul> <li>Interactions with tyramine – containing foodstuffs.</li> <li>Sexual dysfunction.</li> <li>Headache/ Dizziness/ Tremor.</li> <li>Sleep disturbances. Weight gain</li> <li>Ankle edema.</li> <li>Hepatotoxicity.</li> <li>Hypertensive crisis.</li> </ul>	<ul> <li>Patients shouldn't start another type of antidepressant for at least a two- week after last dose of any MAOI</li> <li>Moclobemide (RIMA)&gt; freedom from tyramine reactions and its better tolerated.</li> </ul>

Drug	Indication	MOA	Side Effects	Notes			
	MOOD STABILIZERS						
LITHIUM	<ul> <li>Bipolar Mood Disorder.</li> <li>Prophylaxis in recurrent unipolar depression.</li> </ul>	- Stabilizes neuronal activities.	<ul> <li>Fine tremor, Metallic taste.</li> <li>Fatigue /Weight gain.</li> <li>Renal complications.</li> <li>Reversible hypothyroidism and nephrogenic diabetes insipidus.</li> <li>Fetal cardiac anomalies (1<sup>st</sup> tri).</li> </ul>	<ul> <li>Labs → RFT, electrolytes / TFT/ ECG/ Pregnancy test</li> <li>Drug interactions → Thiazide/ NSAID/ACEi.</li> <li>May potentiate the effect of muscle relaxants.</li> <li>Plasma concentrations:</li> <li>0.9 - 1.2 (acute phase).</li> <li>0.4 - 0.8 (prophylaxis).</li> </ul>			
Carbamazepin	<ul> <li>Not responsive to lithium.</li> <li>Acute mania</li> <li>Prophylaxis,</li> <li>Controlling impulsive and aggressive behavior in persons who are not psychotic.</li> </ul>	-	<ul> <li>CNS (sedation, vertigo, blurred vision and ataxia).</li> <li>SIADH.</li> <li>hepatitis, pancreatitis, serious skin reactions (Stevens-Johnson syndrome).</li> <li>Agranulocytosis and aplastic anemia.</li> </ul>	<ul> <li>↓ serum concentrations of numerous drugs ( OCP, warfarin, haloperidol, valproate)</li> </ul>			
Valproate	- Manic episode with mood and schizoaffective disorders.	-	<ul> <li>CNS (sedation, drowsiness, dysarthria, and ataxia).</li> <li>Hepatotoxicity, pancreatitis.</li> <li>Fetal neural tube defects (spina bifida).</li> </ul>	-			
	Anxiolytics						
Benzodiazepine short acting: - Alprazolam - Lorazepam.	<ul> <li>Panic disorder (for rapid onset).</li> <li>Social phobia (small doses).</li> <li>Specific phobia (before exposure session).</li> </ul>	- They act on benzodiazepine receptors which linked with	<ul><li>Dizziness and drowsiness.</li><li>Release of aggression.</li></ul>	DC 2-3 weeks in detoxification to prevent dependency. Withdrawal Syndrome:			
<b>Benzodiazepine</b> Long acting - Diazepam. - Clonazepam.	<ul> <li>OCD, ASD, PTSD.</li> <li>Conversion disorder (Diazepam).</li> <li>Alcohol detoxification (long acting)</li> </ul>	GABA receptors in the C.N.S → ↑ GABA action which has an inhibitory effect.	<ul> <li>Dependence and withdrawal (Short acting).</li> <li>Intoxication (Similar to alcohol): slurred speech -incoordination</li> </ul>	<ul> <li>2 - 3 days after cessation of short acting.</li> <li>7 days after cessation of long acting.</li> </ul>			

Drug	Indication	MOA	Side Effects	Notes
Buspirone	- GAD.	<ul> <li>Stimulates 5HT – 1A receptors and reduces 5HT (serotonin) transmission.</li> </ul>	<ul> <li>Headache.</li> <li>Irritability.</li> <li>Nervousness.</li> <li>Lightheadedness.</li> </ul>	<ul> <li>Gradual onset of action (days <ul> <li>weeks) → not effective on</li> <li>(as required) basis.</li> </ul> </li> <li>It doesn't lead to <ul> <li>dependence.</li> </ul> </li> </ul>
<b>Beta Blockers</b> - Propranolol	<ul> <li>Social phobia.</li> <li>Specific phobia (before exposure session).</li> <li>GAD</li> <li>Neuroleptic-induced akathisia</li> <li>Lithium-induced postural tremor.</li> <li>Control of aggressive behavior.</li> </ul>	- 30-60 minutes before the anxiety-provoking situation	- Depression.	Caution in patients with: - Asthma - Insulin- dependent diabetes. - Cardiac diseases. - Propranolol = uterine contraction

	Cognitive enhancing medications					
Class Drug Indication MOA				Side Effects	Notes	
Cholinesterase	- Donepezil	- All stages of Alzheimer's	-	<ul><li>Diarrhea, weight loss</li><li>Bradycardia, syncope</li></ul>	-	
	- Rivastigmine - Galantamine	- Mild to moderate Alzheimer's	-	<ul> <li>Anorexia, fatigue, somnolence, dizziness</li> </ul>	<ul> <li>Rivastigmine is available as skin patches</li> </ul>	
NMDA receptor antagonist	- Memantine (Ebix)	<ul> <li>Mild to moderate</li> <li>Alzheimer's</li> <li>Lewy body dementia</li> </ul>	- It acts on the glutamatergic system by blocking NMDA receptors to protect neurons from neurodegenerative process induced by glutamate excitotoxicity	<ul> <li>Confusion, dizziness, drowsiness, headache, insomnia, agitation, hallucinations</li> <li>Less common: anxiety, hypertonia, cystitis, increased libido</li> </ul>	-	

	Substance Abuse					
Class	Drug	Intoxication	Withdrawal	Treatment	Notes	
CNS	Alcohol Alcohol - Irritability - Incoordination - Apathy - Ataxia - slurred speech, - Alcoholic coma mg/ml)	<ul> <li>Incoordination</li> <li>Apathy</li> <li>Ataxia</li> <li>slurred speech,</li> <li>Alcoholic coma (&gt;300</li> </ul>	<ul> <li>Stage I (6-8h):</li> <li>Autonomic hyperactivity: tremors, tachycardia</li> <li>Stage II (10-30h):</li> <li>Hallucinations</li> <li>Stage III (12-48h):</li> <li>Grand mal seizures</li> <li>Stage IV (2-3 days):</li> <li>Delirium Tremens: Delirium, autonomic &amp; electrolytes disturbance, dehydration.</li> <li>Worsens at night.</li> </ul>	Conscious patient: - Observation & support - Agitation, hyperactivity: Haloperidol Unconscious patient: - Hospitalization - Protection of airway, - Forced diuresis & alkalization of urine Detoxification: - Benzodiazepines (diazepam, Chlordiazepoxide) Delirium Tremens: - Admission in ICU, benzodiazepines +/- Mg sulfate, rehydration, Thiamine (B1) → glucose metabolism.	<ul> <li>Give vitamin B1 (thiamine).</li> <li>Anticonvulsants for seizures</li> <li>Disulfiram (antabuse): blocks oxidation of alcohol causing accumulation of acetaldehyde: flushing, choking sensation, headache, N/V, tachycardia, anxiety.</li> <li>(Citrate Calcium Carbimide can be used)</li> <li>Screening: CAGE</li> </ul>	
Suppressants Sec - Benz - Zolpi Inl Aromat (acet	Sedatives - Benzodiazepines - Zolpidem, zopiclone	<ul> <li>Slurred speech</li> <li>Incoordination</li> <li>Unsteady gait</li> <li>Nystagmus</li> <li>Impaired attention</li> </ul>	<ul> <li>Autonomic hyperactivity</li> <li>N/V, anorexia</li> <li>Insomnia</li> <li>Perceptual disturbance</li> <li>Seizures, delirium</li> </ul>	-	<ul> <li>A patient should not be deprived of a benzodiazepine drug when it is clinically indicated. (insomnia, anxiety, akathisia)</li> </ul>	
	Inhalants Aromatic hydrocarbons (acetone, benzene, toluene)	<ul> <li>Small doses: euphoria, excitement, pleasant floating sensation, disinhibition.</li> <li>High doses: disturbed consciousness, impulsiveness, ataxia, nystagmus.</li> </ul>	-	- Treat psychiatric complications. - Education.	<ul> <li>Inexpensive, legal.</li> <li>Symptoms: unusual breath, rash around nose &amp; mouth, residues.</li> <li>Complications: multi-organ damage, depression, respiratory depression, aspiration, cardiac arrhythmia.</li> </ul>	

	Class	Drug	Intoxication	Withdrawal	Treatment
	Opioids Opium, heroin, morphine, codeine, pethidine, methadone	<ul> <li>Euphoria, analgesia, relaxation</li> <li>Apathy, dysphoria, drowsiness, slurred speech, ↓ sexual desire.</li> <li>Pupillary constriction (pupillary dilatation occurs in severe overdose)</li> </ul>	<ul> <li>Rhinorrhea</li> <li>Lacrimation</li> <li>Pupillary dilatation</li> <li>Yawning, insomnia</li> <li>Fever, sweating</li> <li>Muscle/joint ache</li> <li>N/V, Diarrhea</li> <li>Dysphoric mood</li> <li>Craving</li> </ul>	<ul> <li>Intoxication:</li> <li>ICU, Open airway, O2, fluids</li> <li>Antidote (Naloxone)</li> <li>Withdrawal:</li> <li>Short term: painkillers, sedatives, observation, Clonidine.</li> <li>Long term: Methadone harm reduction strategies (methadone patches can be used for heroin addicts)</li> </ul>	<ul> <li>Interaction with endogenous opioids (enkephalins, endorphins and dynorphins) and opiate receptors (mu, kappa and delta).</li> <li>Heroin may cause vein thrombosis, PE</li> <li>Tolerance may develop &amp; diminish rapidly (6h): withdrawal is not fatal but taking a large does can cause respiratory arrest.</li> </ul>
CNS Stimulants	Amphetamine (captagon) - Overce - Aggres	<ul> <li>Hypervigilance</li> <li>Overconfidence</li> <li>Aggression</li> <li>Insomnia</li> </ul>	-	- Antipsychotic medications (Olanzapine)	- It can be indistinguishable from functional psychosis
	Cocaine (crack)	- Euphoria - Hallucinations		- Antidepressants	- Positive finding in urine screen
Cannabis	- Marijuana - Hash - Hashish	<ul> <li>Euphoria, Dry mouth</li> <li>Red conjunctiva</li> <li>Mild tachycardia</li> <li>↑ Appetite, Brief psychosis.</li> <li>Impaired cognitive functions</li> <li>Anxiety +/- depersonalization &amp; derealization.</li> </ul>	- Depressive features	<ul> <li>Outpatient</li> <li>Antipsychotic medications</li> </ul>	- Chronic use of cannabis can lead to a state of apathy and amotivation (amotivation syndrome)

- TB treatments (INH, cycloserine) can cause many psychiatric disorders
- Antihypertensive (beta blocker), steroids, chemotherapy may induce depression
- Amitriptyline is used to treat depression, anxiety, and pain symptoms in rheumatoid arthritis.
- Interferon therapy for hepatitis C can cause severe depression and suicidal ideation.
- Pethidine (narcotic) = addicted in SCA

Neurotransmitter	Diseases	Notes
<b>Serotonin</b> - Raphe nuclei. - L-tryptophan. - Metabolized by MAO-A	<ul> <li>Anxiety</li> <li>Depression</li> <li>Bulimia nervosa</li> <li>OCD</li> </ul>	SHT1: Frontal cortex $\rightarrow$ impulsive behaviour and anxiety SHT2A: 1- Basal ganglia $\rightarrow$ movement. 2- Brainstem $\rightarrow$ deep sleep. 3- Spinal cord $\rightarrow$ sexual responses and orgasm.SHT2A-C: Limbic system $\rightarrow$ anxiety and panic feelings. SHT3: 1- Brainstem chemoreceptors $\rightarrow$ vomiting. 2- Hypothalamus $\rightarrow$ weight gain and appetite.Peripheral serotonergic receptors 3,4 and 7 Intestinal secretion
<ul> <li>Dopamine</li> <li>From tyrosine</li> <li>D1&gt; -ve symptoms</li> <li>D2 blockade produces parkinsonian like effects so motor.</li> </ul>	Schizophrenia	<ul> <li>Mesolimbic path: Reward system → positive symptoms of schizophrenia → <u>nucleus</u> accumbens → reinforcement is achieved through the effects of nicotine, caffeine and CNS stimulants</li> <li>Mesocortical path: Cognition and mental arousal → negative symptoms of schizophrenia         <ul> <li>a. Pathological causes of this path could be due to → Dopamine neuron defect, Glutamate excitation, Serotonin overactivity.</li> <li>Nigrostriatal: Motor and parkinsonian like symptoms</li> <li><u>Tuberoinfundibular:</u> Production of prolactin</li> </ul> </li> </ul>
Noradrenaline - Hydroxylation of dopamine from Locus Cerulus	<ul> <li>Ejaculation and orgasm through the sympathetic system</li> <li>Stimulation of immune system in low doses and inhibition in high doses</li> </ul>	1- Frontal cortex:         • Mood through Beta 1 receptors         • Cognition through Alpha 2 receptors         2- Limbic system         3- cerebellum         4- cardiovascular
ACH - Nucleus basalis of meynert.	<ul> <li>Erection through parasympathetic system</li> <li>Other activators include NO</li> </ul>	
GABA - Synthesized by glutamate	Suppresses seizure activity, anxiety and mania	Inhibitory exclusively found in the brain
<b>Glutamate</b> - Synthesized by <mark>glutamine</mark>	Neuronal degeneration in chronic schizophrenia with negative features	<ul> <li>Excitatory neurotransmitter</li> <li>Sigma receptors 1&amp;2 are related to glutamate receptors NMDA and involved in <u>enhancement</u> <u>of memory and cognition</u> and hence when fluvoxamine is used it alleviates the negative symptoms in schizophrenia</li> </ul>

Neurotransmitter	Diseases	Notes
Substance P	<ul> <li>Initiation of pain.</li> <li>Migraine.</li> <li>Cluster headaches.</li> <li>Chronic pain.</li> <li>Huntington's disease.</li> <li>Mood disorders</li> </ul>	Excitatory neurotransmitter
<b>Melatonin</b> - Pineal gland	Sleep wake cycle	
Histamine - Hypothalamus	Allergies	<ul><li>H1- arousal and appetite.</li><li>H2- gastric ulcer.</li></ul>
Endogenous opioids	Pain	Learning , memory , mood and perception of pain

CBT	<ul> <li>Panic disorders.</li> <li>Agoraphobia.</li> <li>Social phobia.</li> <li>GAD.</li> <li>Acute stress disorder.</li> <li>Adjustment disorder crisis intervention.</li> </ul>	
Behavioural therapy	<ul> <li>Social phobia.</li> <li>Thought stopping OCD.</li> <li>Relaxation for anxiety.</li> <li>Exposure (with desensitization and flooding).</li> <li>Token economy for chronic schizophrenia</li> <li>Assertiveness training for dependant and avoidant personality</li> </ul>	
<b>Psychodynamic</b> - Patients behaviour is determined by an unconscious process (personality disorders, anxiety or chronic depression )		
Martial	- Problem solving - Behavioural - Dynamic	

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