



Mental Health Meds

WHAT, WHY, WHEN & HOW

Dr Tim Worthley 22.10.25

Arch.



Outline

- What are mental health medications?
- Neurotransmitters
- Questions to ask before prescribing/taking
- Run through of the main types of medications
 - How they work
 - Potential side effects
 - Useful (hopefully) info



This is not(!)

- An in-depth biochemical explanation of how they all work
- A list of every single mental health medication
- A conversation about psychiatry vs anti-psychiatry, the evils of drug companies, or a profound comment on what is normal and what are we even medicating anyway



What is MH medication

- Prescription drug that affects the brain to alter mood, behaviour, thoughts, or perceptions, often used to treat mental health conditions like depression, anxiety, and psychosis.
- These medications work by influencing brain chemistry, specifically neurotransmitters like dopamine, serotonin, and noradrenaline.
- Common types include antidepressants, antipsychotics, mood stabilisers, stimulants, sedatives & anxiety medication.

Neurotransmitters

Serotonin

BRAIN: ↑mood, sleep, ↓libido

BODY: stimulates gut, insulin, vasoconstriction. Pain & Temperature regulation

Noradrenaline

BRAIN: stress, attention, focus; alertness, arousal

BODY: Raised BP, pulse, sugar, slows intestines

Dopamine

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BODY: fine tuning of movements, GI both, CV both

GABA “off switch”

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Acetylcholine

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Excitatory – pain, sensory information, nerve activity

Histamine

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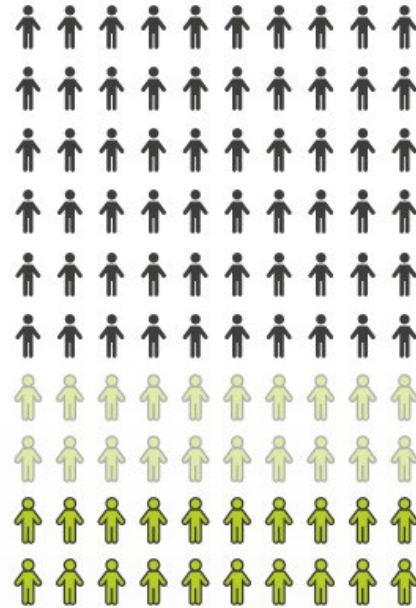
Antidepressants

- Questions to ask
 - Depression vs Distress
 - Other chemical influences in brain
 - Moderate-Severe Depression
 - First 2 weeks
 - Take regularly?
 - Impact of side effects, any contraindications?
 - Risk of overdose
 - What's the plan?
 - Risk of discontinuation/withdrawal
 - Placebo

How well can **antidepressants** relieve symptoms?

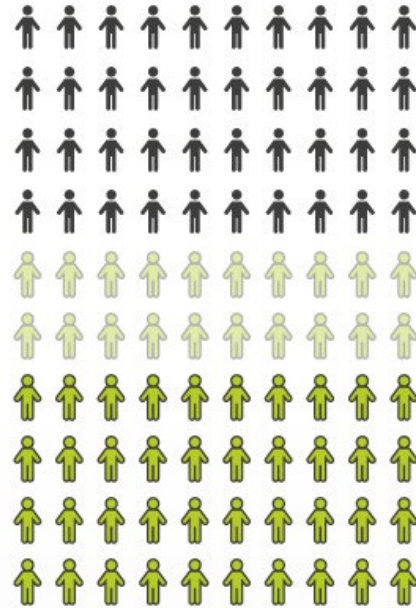
Studies of people with moderate or severe depression showed:

Without antidepressant



 = About **20 to 40** out of 100 people who took a placebo noticed an improvement of their symptoms within six to eight weeks

With antidepressant



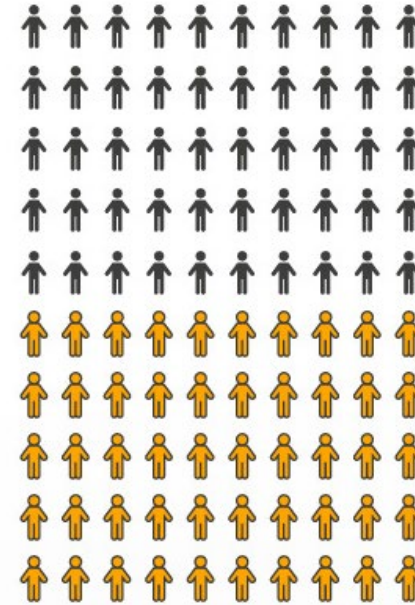
 = About **40 to 60** out of 100 people who took an antidepressant noticed an improvement of their symptoms within six to eight weeks

That means: Antidepressants improved symptoms in about 20 out of 100 people.

How well can **antidepressants** prevent relapses?

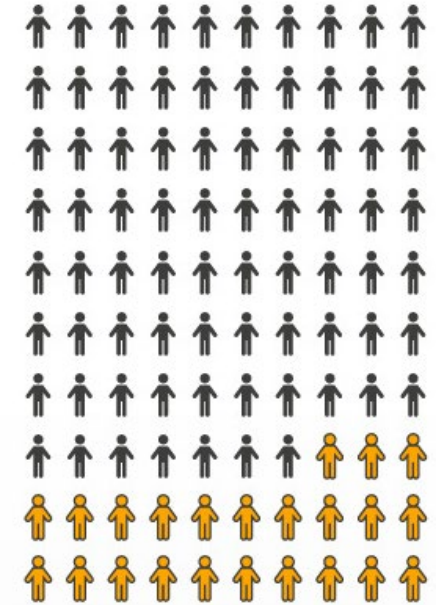
Studies showed that antidepressants cannot completely prevent relapses, but that they can lower the risk:

Without antidepressant



 = About **50** out of 100 people who took a placebo had a relapse within one to two years

With antidepressant



 = About **23** out of 100 people who took an antidepressant had a relapse within one to two years

That means: Taking an antidepressant for a longer period successfully prevented a relapse in 27 out of 100 people on average.

Tricyclic Antidepressants (Amitriptyline)

- 1st Generation
- Not very selective - NA & Serotonin, BUT ALSO block ACh, histamine
- Mood, Sleep, Pain, Migraines, Agitation
- **BUT**
- Side effects – dry you up and knock you down
- Very risky in overdose
- Consider if other options fail, or other effects needed
- Very careful dose management

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SSRI Antidepressants

- Serotonin Selective Re-uptake Inhibitors
- Citalopram, Sertraline, Fluoxetine, Escitalopram, Paroxetine
- More selective. But other NTs are impacted
- Can also be useful for anxiety & sometimes chronic pain
- Beware – reflux, weight, GI, sexual side effects, serotonin syndrome, discontinuation (esp Paroxetine)
- “Take every morning after breakfast”

Other Antidepressants

SNRI's

Venlafaxine, Duloxetine

- More stimulating
- Can be useful in chronic pain

Mirtazapine

NA, Serotonin, Histamine

- Sedative – as much at 15mg as 45mg
- Weight gain ++

Trazodone

NA, Serotonin, Dopamine, Ach, Histamine

- Unselective, therefore more side effects
- Drowsiness, NA side effects

Pause



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Antipsychotics

1st Generation – “typical”

eg Haloperidol, Chlorpromazine

2nd Generation – “atypical”

eg Risperidone, Olanzapine, Quetiapine, (Aripiprazole)

CLASSES

* TYPICAL (1st GENERATION)

~ HIGH or LOW POTENCY → AMOUNT of DRUG REQUIRED to MINIMIZE SYMPTOMS

~ NOT SELECTIVE TO D₂ RECEPTORS in MESOLIMBIC PATHWAY → WORSENING NEGATIVE SYMPTOMS

* ATYPICAL (2nd GENERATION)

~ BLOCK BOTH D₂ RECEPTORS & SEROTONIN 5-HT_{2A} RECEPTORS → ↓↓ NEGATIVE SYMPTOMS

SIDE EFFECTS

* HIGH-POTENCY, 1ST GENERATION

~ EXTRAPYRAMIDAL SYMPTOMS

~ TARDIVE DYSKINESIA

~ NEUROLEPTIC MALIGNANT SYNDROME



* LOW-POTENCY, 1ST GENERATION

~ DRY MOUTH

~ CONSTIPATION

~ SEDATION

~ DIZZINESS



* 2nd GENERATION

~ WEIGHT GAIN

~ DRUG-INDUCED TYPE 2 DIABETES

~ TIREDNESS



	Extrapyramidal	Sedation	Weight gain	Hyperglycaemia	Anticholinergic	Orthostatic hypotension
Atypical antipsychotics						
Risperidone	●●	●● initially	●●	●●	●	●● initially
Quetiapine	●*	●●●	●●	●●●	●●	●●
Olanzapine	●	●●●	●●●	●●●	●●●	●
Clozapine	●	●●●	●●●	●●●	●●●	●●
Amisulpride	●●*	●	●	●	●	●
Aripiprazole	●	●	●	●	●	●
Ziprasidone	●	●●	●	●	●	●●
Typical antipsychotics						
Haloperidol	●●●	●	●●	●●	●	●
Chlorpromazine	●●	●●●	●●●	●●●	●●●	●●●

Approximate frequency of adverse effects: ● (<2%) = negligible or absent; ● (>2%) = infrequent; ●● (>10%) = moderately frequent; ●●● (>30%) = frequent. * rarely a problem at usual therapeutic doses

Antipsychotics and Prolactin Levels

- Prolactin-elevating antipsychotics
 - Risperidone and paliperidone
- Variable effect on prolactin
 - Olanzapine, lurasidone, asenapine, ziprasidone
- "Prolactin-sparing" antipsychotics
 - Iloperidone, quetiapine, clozapine, aripiprazole



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~ CONSTIPATION

~ SEDATION

~ DIZZINESS



* 2nd GENERATION

~ WEIGHT GAIN

~ DRUG-INDUCED TYPE 2 DIABETES

~ TIREDNESS



Pause



ADHD Medication

- Stimulants
 - Methylphenidate
 - Lisdexamfetamine
 - Slow and steady wins the race, esp for adults
 - Extraordinary BUT risky especially in substance use
- Non-stimulants
 - Atomoxetine
 - (Guanfacine, Clonidine)



Sedatives

- Benzodiazepines
 - Stimulate GABA (like alcohol)
 - Addiction, memory loss, depression, falls, possibly dementia, death
 - Never more than 2 weeks
 - Opportunity cost
- Z-drugs – questionable as to whether they are any better
- Antihistamines – promethazine. Lower histamine and ACh

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Mini Pause



Mood Stabilisers

- Recycled & Lithium
- Anticonvulsants (Epilepsy Medications)
 - Beware Pregabalin and Gabapentin!!
 - Special mention to Lamotrigine
 - Also Sodium Valproate (Epilim), Carbamazepine (Tegretol)
- Antipsychotics – particularly in mania
- Antidepressants - with caution



Lamotrigine rash (SJS)



Anxiety Medication

- Recycled & Difficult
- “When required”
 - Non-sedative – propranolol (beware asthma)
 - Sedative
 - promethazine, BDZ (only very short-term)
 - low dose Quetiapine but only if severe – usually Psychiatrists
- Regular
 - Antidepressants
 - Beware the more stimulating options
 - Sometimes need higher doses
 - Antipsychotics – Only if severe, usually Psychiatrists, only 2nd Gen
 - (Buspirone)



Mini Pause



Closing thoughts

- Complex
- Potential for more harm than good
- Can have profoundly positive impact
- Lack of available lifestyle interventions results in increased prescribing
- Effective prescribing requires:
 - Understanding of the person and good relationship
 - Regular monitoring
 - Avoidance of long-term if possible, unless SMI
 - Changing when situation changes
- De-prescribing can be hugely impactful





Thank you
Any Questions?



Arch.



Frontline
Network
Partner