

Psychopharmacology

Hamish McAllister-Williams

Reader in Clinical
Psychopharmacology

Department of Psychiatry, RVI

Intro

- NOT a comprehensive review of everything you need to know about drug treatments of mental illness!
- Remember:
 - 30% of patients visiting their GP have mental health problems
 - 20-50% of patient in hospital out-patient clinics have mental health problems
 - This is NOT some highly specialised area that you only need to have a vague knowledge of
 - All doctors need to know something about psychotropic drugs

Plan

- General comments
- Brief overview of major groups of psychotropics
 - Antidepressants
 - Hypnotics/anxiolytics
 - Antipsychotics
 - “Mood Stabilisers”
- NOT
 - Anti-dementia drugs
 - Drugs used in addictions
 - Drugs used for ADHD

Why is the use of drugs in psychiatry different from using drugs in other branches of medicine?

- Fundamental nature of psychiatric illnesses
- Attitudes and beliefs of patients (and doctors)
- Pharmacological complexity of psychotropic drugs
- Drugs are but one treatment modality

“The study of MEDICINE is
prosecuted under two relations,
namely as a *Science* and as an *Art*”

The Science and Practice of Medicine

W.Aitken

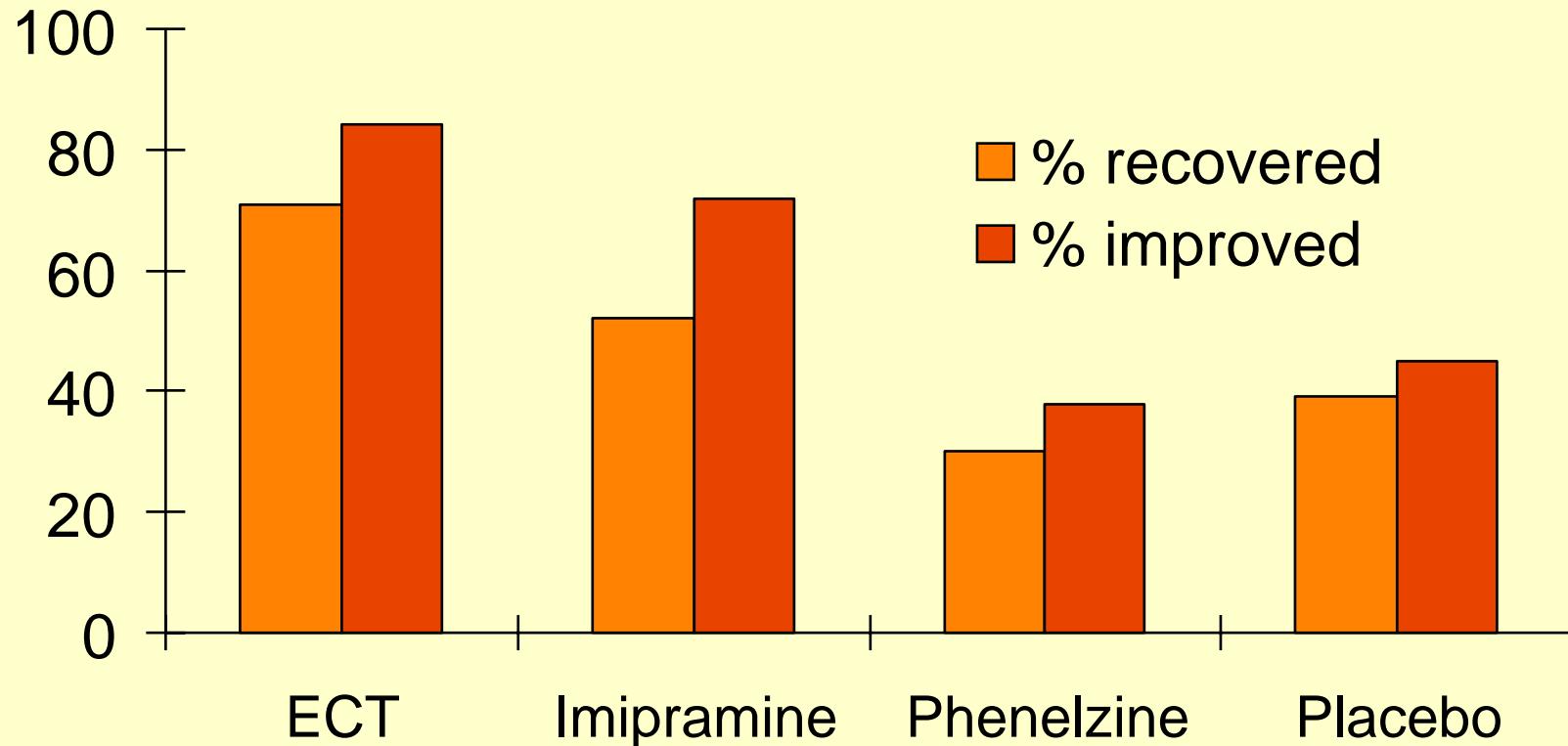
1872

The science of psychopharmacology

Three classic studies

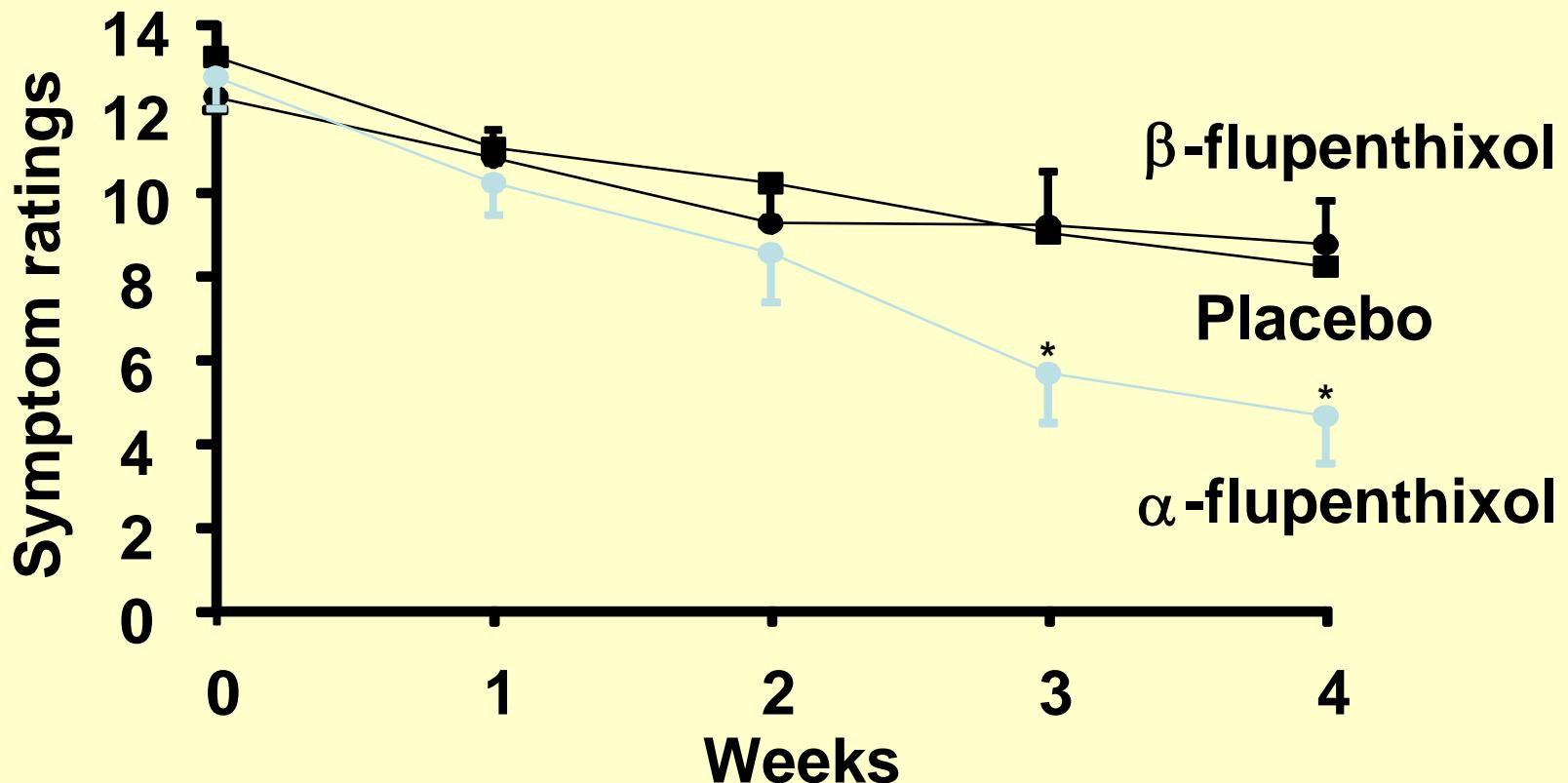
Efficacy of Drug Treatments - Antidepressants

MRC Antidepressant Trial, 1965

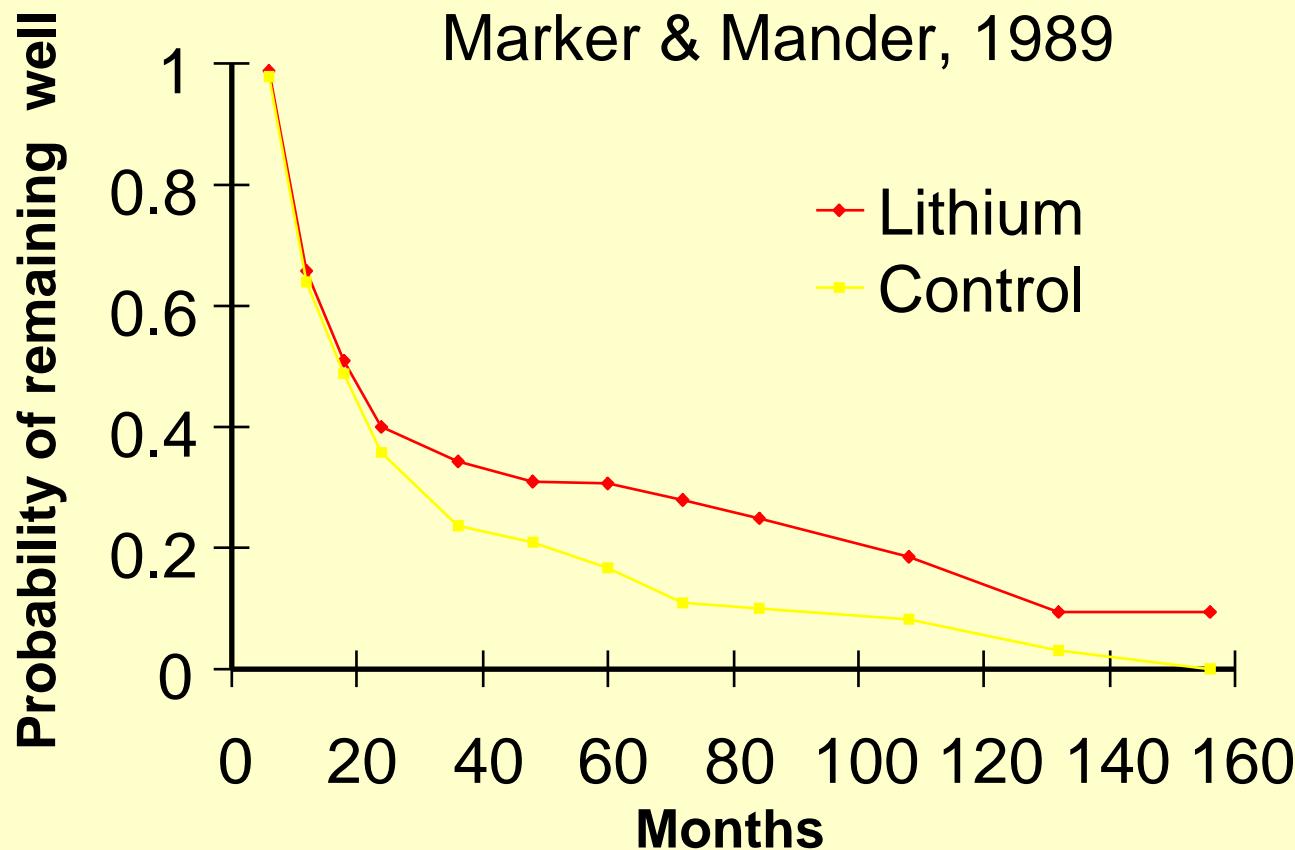


Efficacy of Drug Treatments - Antipsychotics

Johnstone et al. 1978 - Treatment of Schizophrenia



Efficacy of Drug Treatments - Lithium Prophylaxis



Pharmacological complexity of psychotropic drugs: When science becomes an art

The clinical use of psychotropic drugs

- Idiosyncratic reactions
 - e.g. amphetamine, SSRIs and migraine
- Lack of therapeutic ranges for drugs
 - e.g. most antidepressants
- Enormous dosage ranges
 - e.g. 10mg – 1500mg chlorpromazine per day
- High rates of non-response
- Multiple classes of drugs
- Polypharmacy

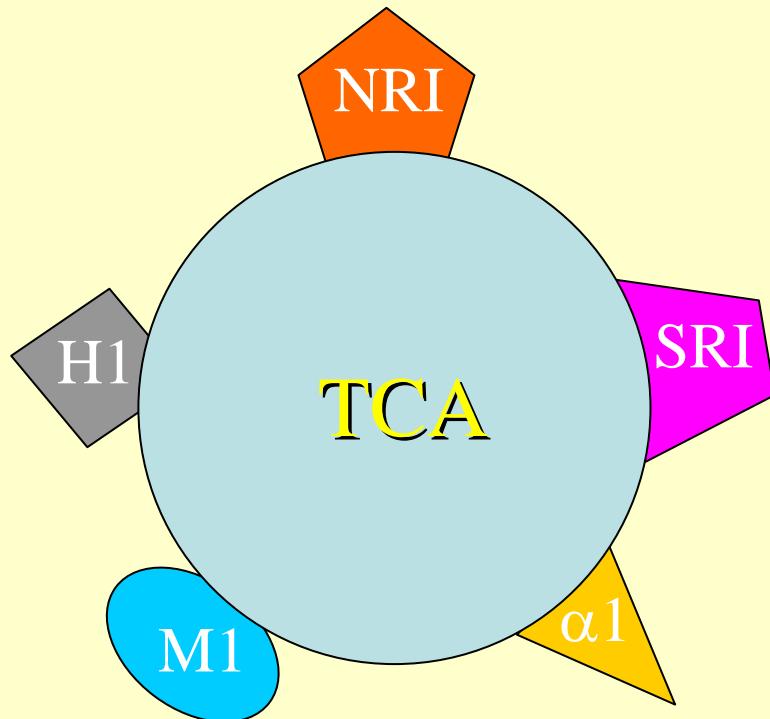
Antidepressants

- TCAs
- SSRIs
- NaRIs
- SNRIs
- Antagonists
- MAOIs
-and others currently available plus ones on the way

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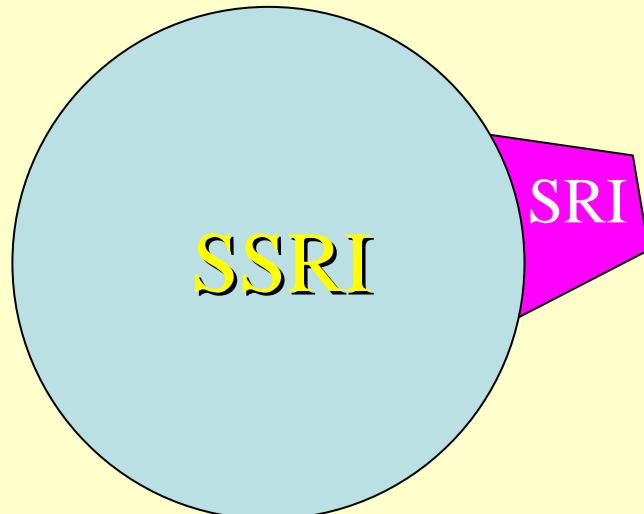
Tricyclic Antidepressants (TCAs)



N.B. also effects on cardiac and neuronal membrane excitability

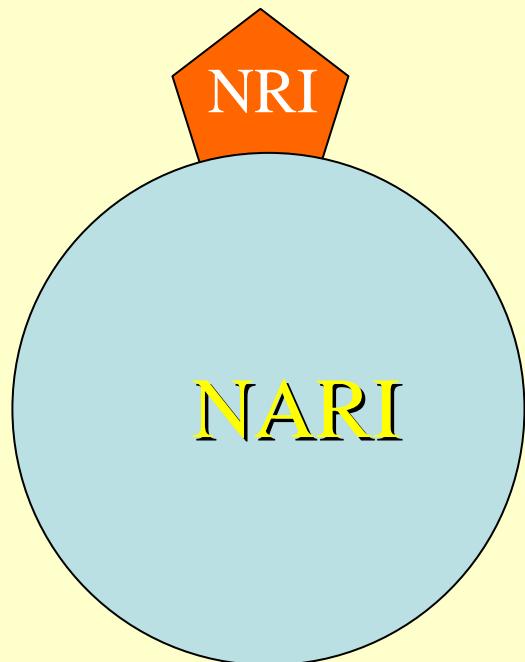
- e.g. amitriptyline, lofepramine, imipramine
- Inhibit 5-HT and NA uptake
 - Produces therapeutic effect
- Block of M_1 , H_1 , α_1 receptors produces side effects
- Poorly tolerated and toxic in overdose (except lofepramine)
- ?Amitriptyline more potent than SSRIs for severe depression

Selective Serotonin Reuptake Inhibitors (SSRIs)



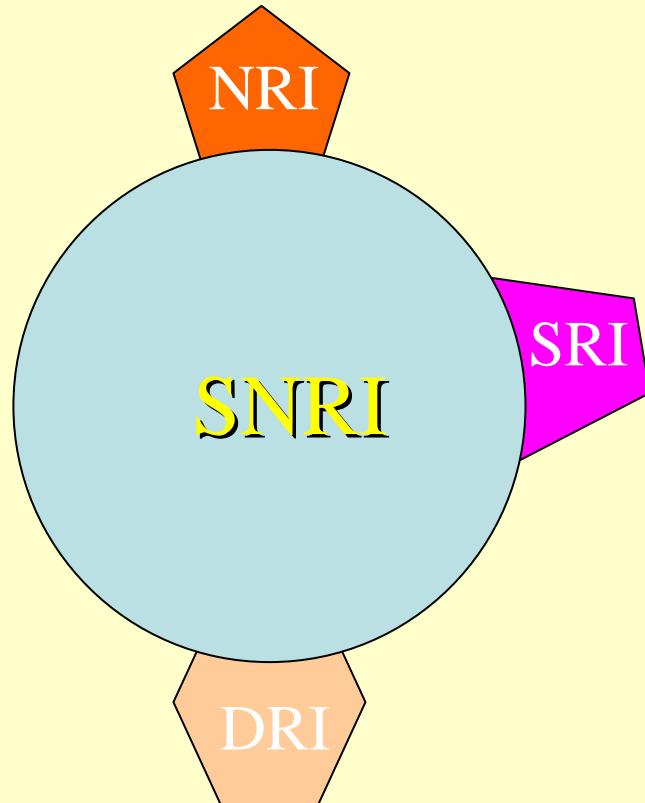
- e.g. fluoxetine, paroxetine, sertraline, citalopram, escitalopram
- Inhibit 5-HT uptake
 - Produces therapeutic benefit
 - depression
 - OCD, Panic, anxiety
 - Produces side effects
 - Nausea
 - Early increased anxiety
 - Sexual dysfunction
- Well tolerated and good first line treatments

Noradrenaline Reuptake Inhibitors (NaRIs)



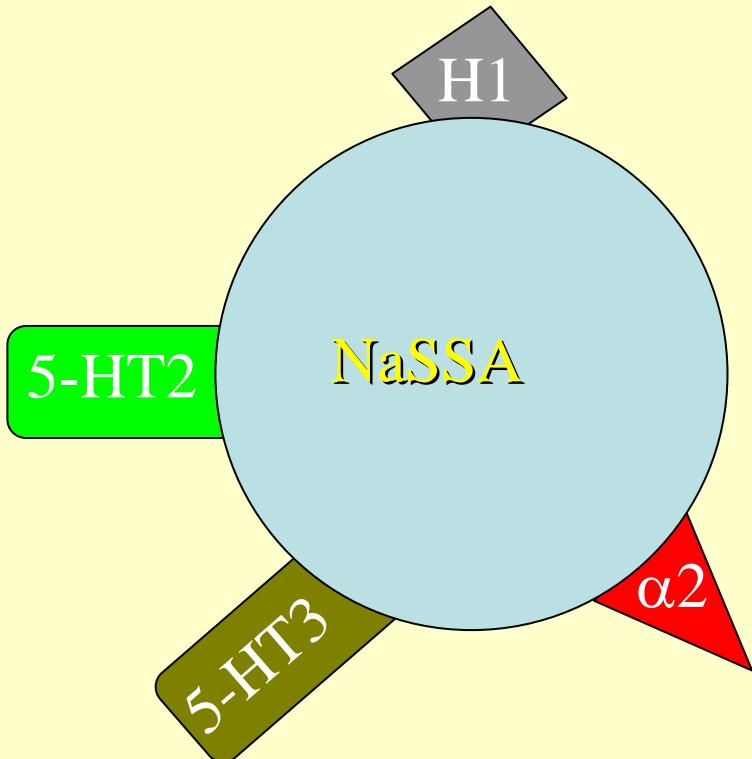
- e.g. reboxetine (lofepramine)
- Inhibit NA uptake
 - Produces therapeutic effect
 - Produces side effects
- Well tolerated
- Alternative class for patients who fail an SSRI

Serotonin and Noradrenaline Reuptake Inhibitors (SNRIs)



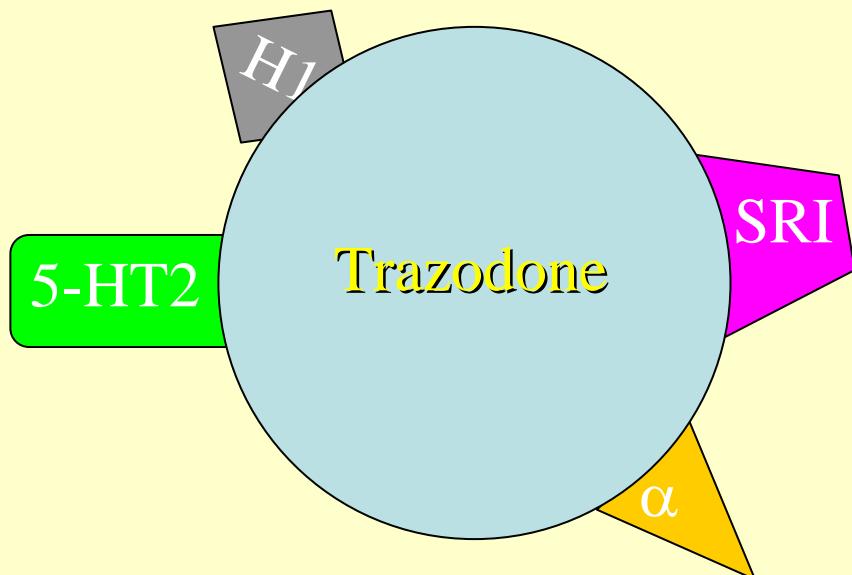
- e.g. venlafaxine, duloxetine
- Inhibit 5-HT and NA (and DA) uptake
 - Produces therapeutic effect
 - Produces side effects
 - Similar to SSRI
- Better tolerated than TCAs and ? more effective than SSRIs for severe depression therefore good second/third line treatment

Noradrenaline and Serotonin Selective Antagonist



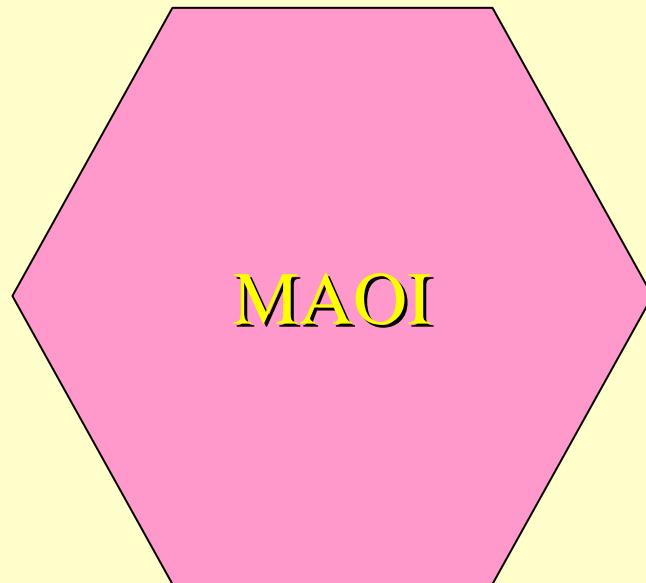
- e.g. mirtazepine
- Blocks α_2 receptors
 - Produces antidepressant effect (Increases 5-HT release)
- Blocks 5-HT₂ receptors
 - Produces decreased anxiety
- Blocks H₁ receptors
 - Produces sedation
- ? more potent than SSRIs plus lacks sexual side effects, but causes marked weight gain.
Used second line

5-HT₂ antagonist



- e.g. Trazodone
- Weakly blocks 5-HT uptake
 - ? effect
- Blocks 5-HT₂ receptors
 - ? Main way produces benefit on depressive and anxiety symptoms
- Blocks H1 receptors
 - Produces sedation
- ? Potency as mainline treatment but often used to augment SSRIs or SNRIs

Monamine Oxidase Inhibitors



- Traditional
 - e.g. phenelzine, tranylcypromine
 - Food & drug interaction
- RIMA
 - e.g. moclobemide
- Increase levels of 5-HT, NA (and dopamine -传统inals)
 - produces therapeutic benefit
- Second line for atypical depression, third line treatments for severe depression

NICE Clinical Guideline 23

December 2004

**Depression: management of
depression in primary and
secondary care**

Guidance

- Good practice points for all
- **Stepped care**
- Step 1: Recognition of depression
- Step 2: Depression in primary care – mild depression
- Step 3: Depression in primary care – moderate to severe
- Step 4: Mental health services – refractory, recurrent, atypical and psychotic depression
- Step 5: Depression requiring inpatient care

Step 2 – Mild depression

- Antidepressants
 - Not recommended for initial treatment
 - Use if
 - symptoms persist after other interventions
 - depression associated with psychosocial problems
 - past history of moderate to severe depression
- Recommended interventions
 - Sleep and anxiety management
 - Watchful waiting
 - Structured exercise
 - Guided self-help

Step 3 - moderate to severe depression

- Antidepressants
 - Antidepressants should be routinely offered
 - Address common concerns
 - Inform about potential side effects and risk of discontinuation/withdrawal symptoms (particularly with paroxetine and venlafaxine)
 - Inform about time delay in response
 - Continue for at least 6 months from remission
 - After 6 months review need for medication

Step 3

- Choice of antidepressant
 - SSRI in routine care
 - fluoxetine or citalopram
 - If response inadequate consider increasing dose to BNF limits
 - If not effective switch antidepressant [C]
 - Reasonable alternative to SSRIs = mirtazepine, but consider moclobemide, reboxetine, lofepramine

Step 4 – Atypical, psychotic and recurrent depression

- Atypical depression
 - Hypersomnia, increased appetite, hypersensitive
 - consider an MAOI (phenelzine)
- Psychotic depression
 - Augment with an antipsychotic
- Recurrent depression
 - Drugs
 - If 2 or more episodes consider ADs for 2 years
 - Use same dose of AD as for acute treatment
 - Psychotherapies
 - Consider CBT

Step 4 - Refractory depression

- Failure to respond to 2 or more ADs
- Consider the following options:
 - ADs plus CBT
 - Lithium augmentation
 - Venlafaxine up to BNF limits
 - SSRI + mianserin or mirtazepine [C]
 - Consider phenelzine

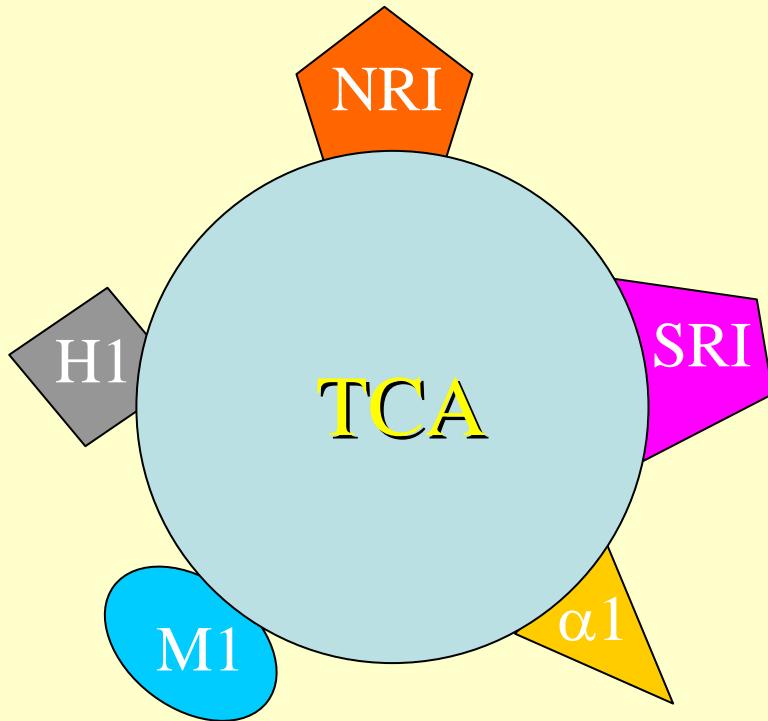
Anxiolytics/hypnotics

- Benzodiazepines e.g. diazepam, lorazepam, chlordiazepoxide, temazepam
- Act on GABA_A receptor complexes
- Relieve anxiety immediately, good for short term use
- Hypnotics are simply BZs with shorter **half lives**
- S/E's - very few except dependency
- **However** antidepressants are the drugs of choice for treating anxiety but take longer to work

Antipsychotics

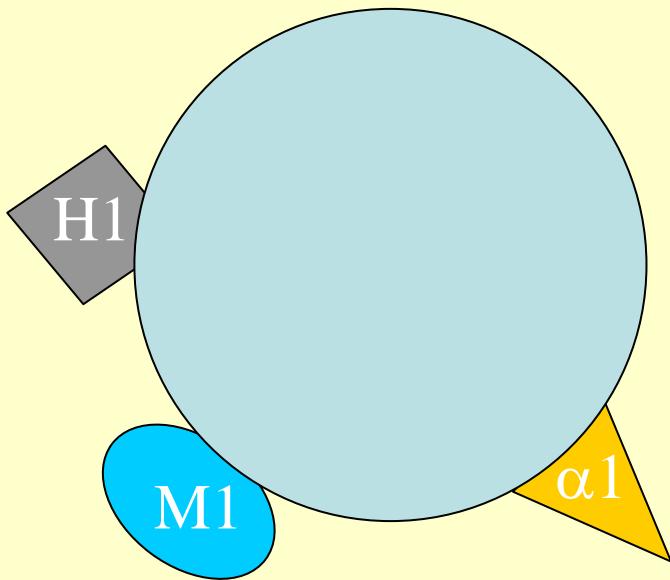
- Typicals/conventionals/first generation antipsychotics
 - e.g. chlorpromazine, haloperidol
- Atypicals/second generation antipsychotics
 - Clozapine
 - Olanzapine
 - Risperidone
 - Quetiapine
 - Aripiprazole
 - Amisulpiride/sulpiride

Typical antipsychotics



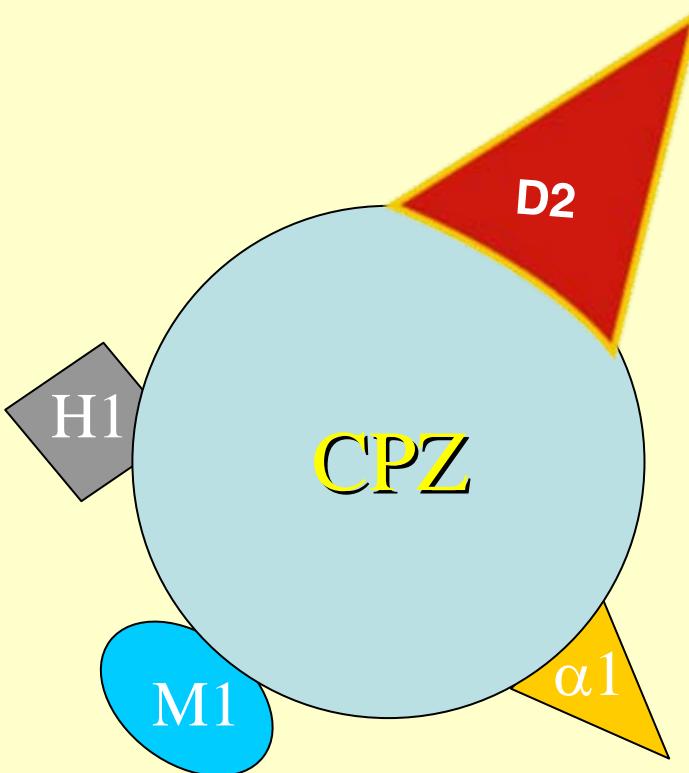
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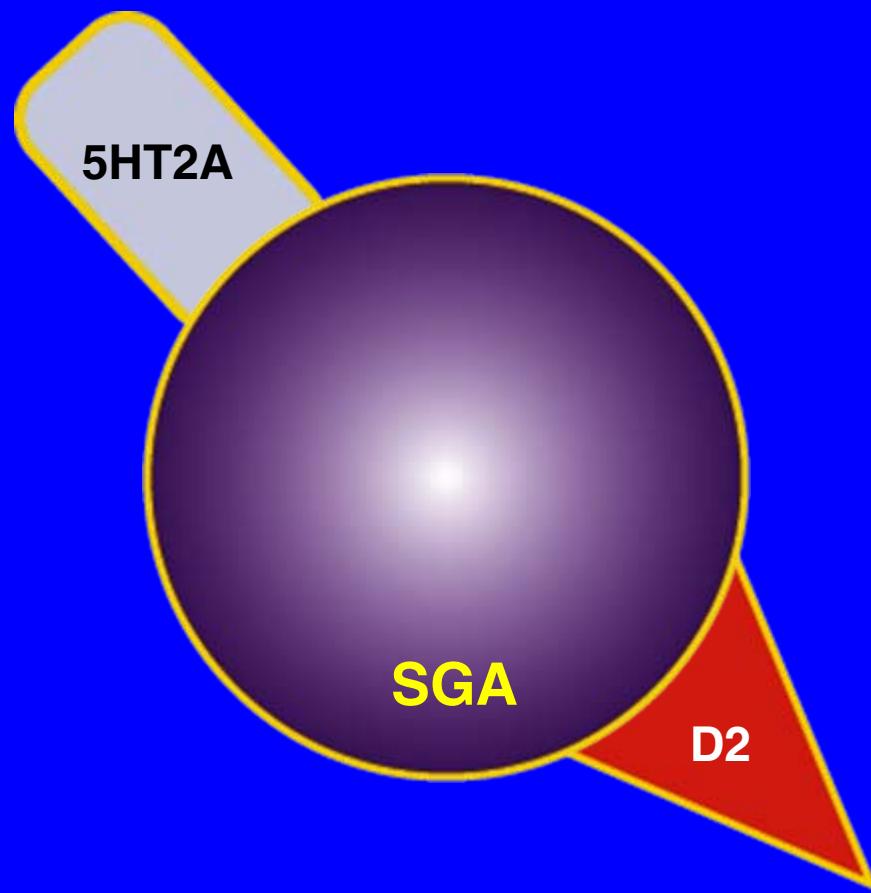
N.B. also effects on cardiac and neuronal membrane excitability

- E.g. chlorpromazine, haloperidol
- Block D₂ receptors
 - Therapeutic effects
 - EPS
- Also antagonise histamine, NA and acetylcholine receptors causing side effects
- Antipsychotic and sedative
- Used in schizophrenia, mania, psychotic depression
- NICE no longer recommend them first line

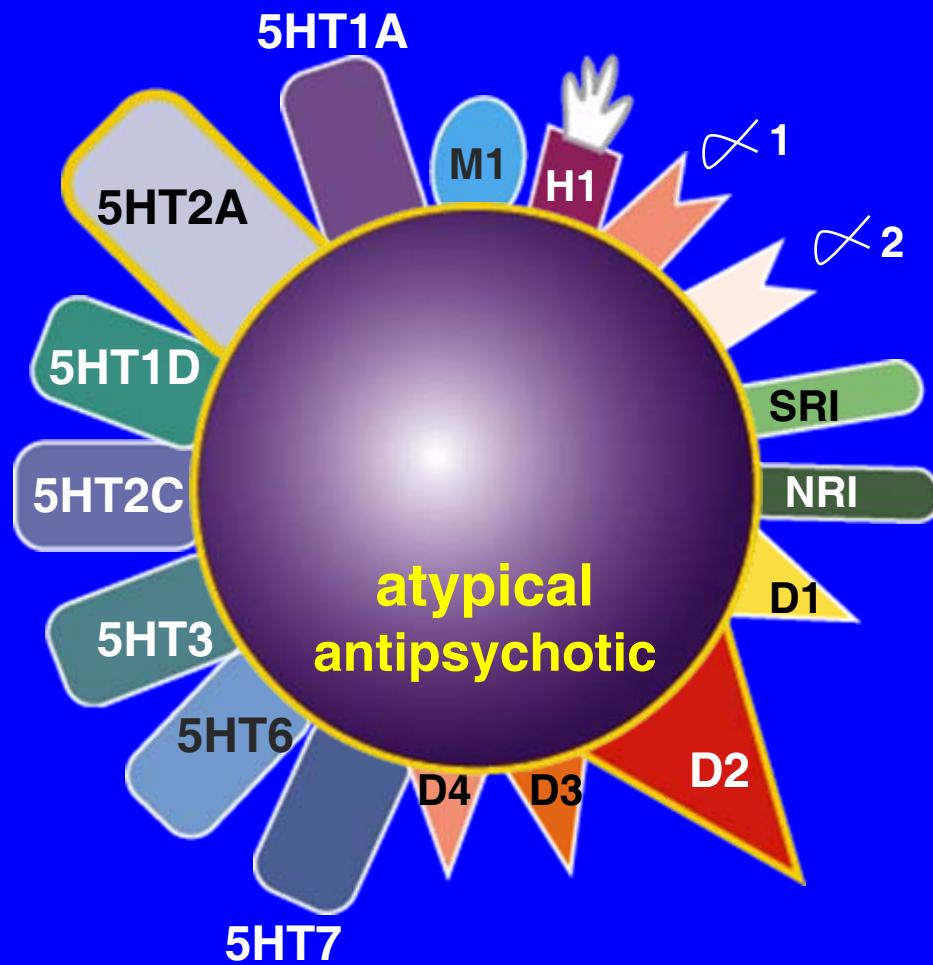
Haloperidol



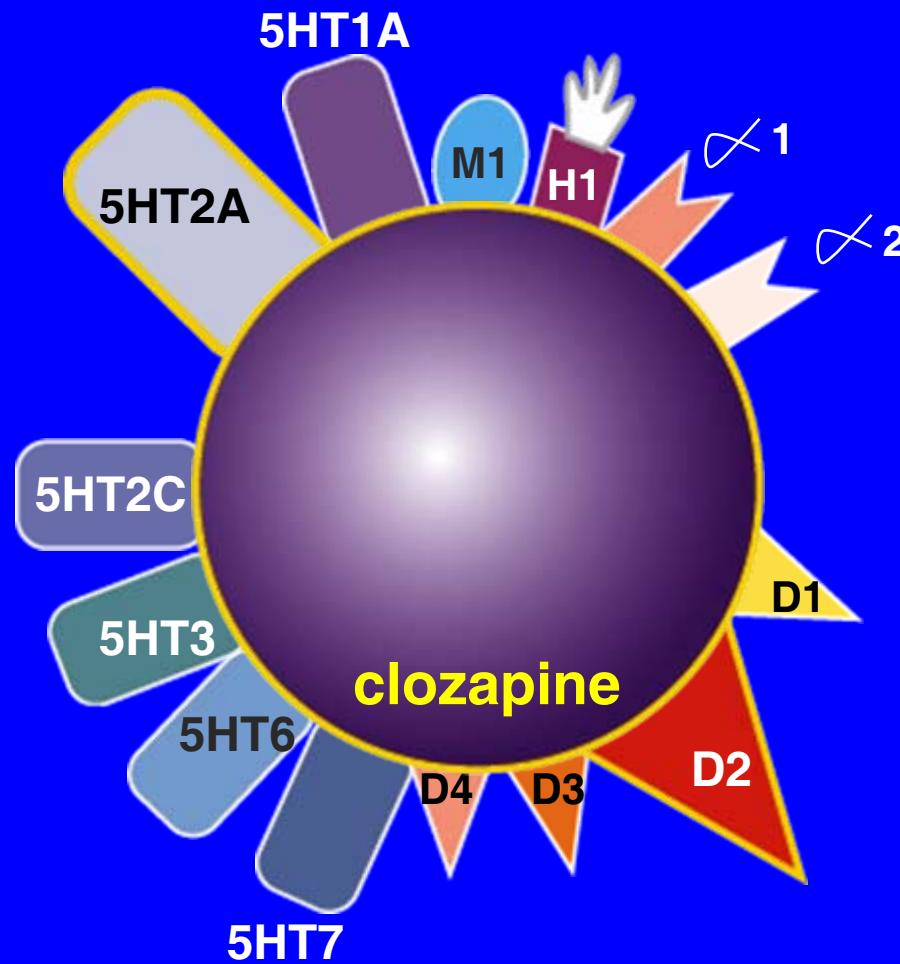
Atypical – core pharmacology



Atypical – “rich” pharmacology



Clozapine – the archetypal atypical



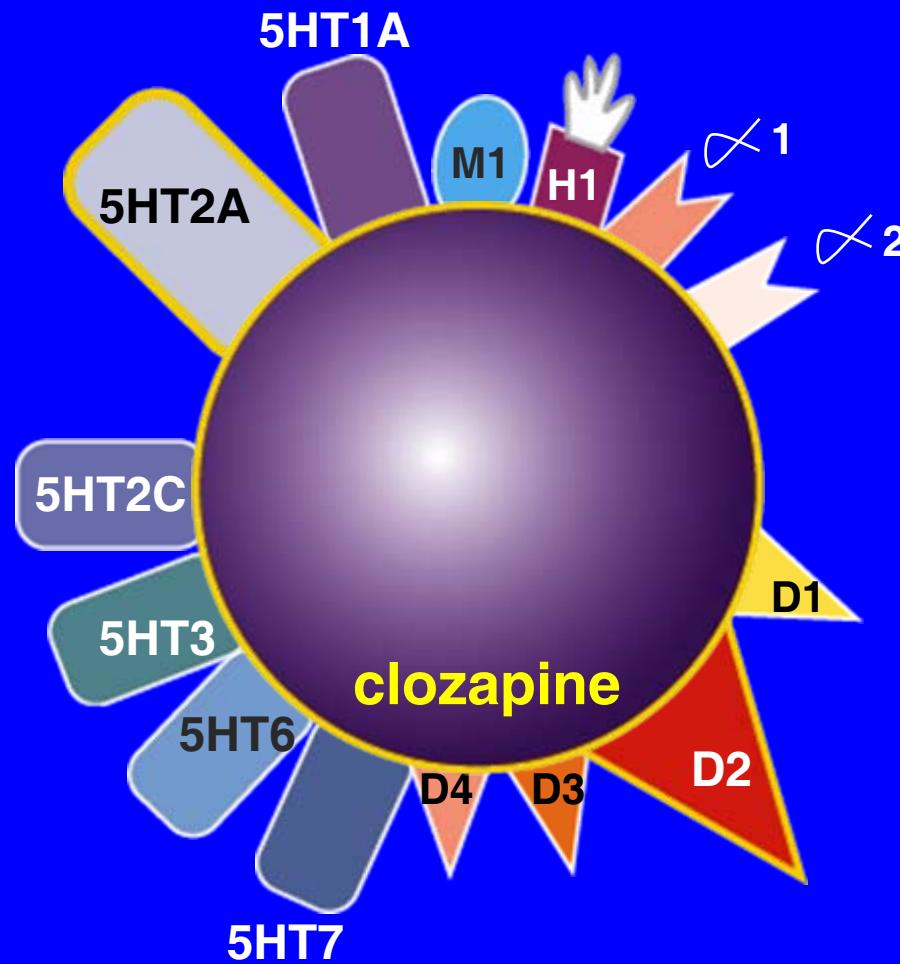
Clozapine

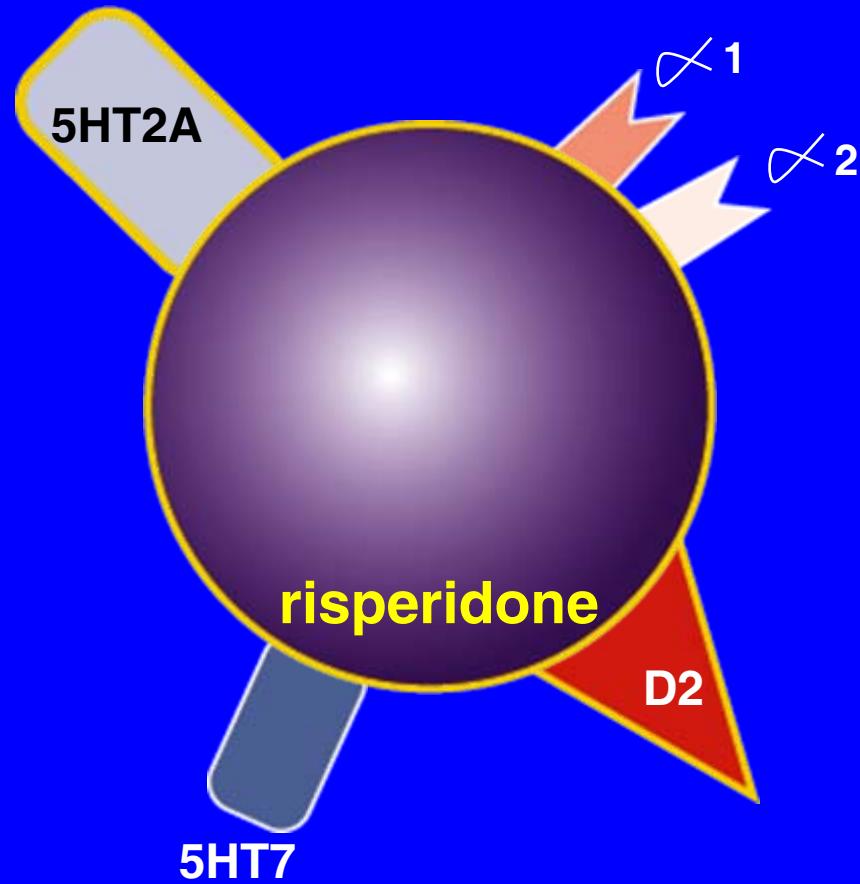
- Introduced in Europe in 1975 (but not UK)
- Less extrapyramidal symptoms than typicals
- Withdrawn due to cases of fatal neutropenia
- Kane et al. (1988) - large trial of 'treatment resistant' schizophrenic patients (300+)
 - failed at least three antipsychotics before entry
 - all given high dose haloperidol - those who failed to respond randomised to chlorpromazine or clozapine
 - 30% response to clozapine c.f. 4% to chlorpromazine
 - significant effect on 'negative symptoms'

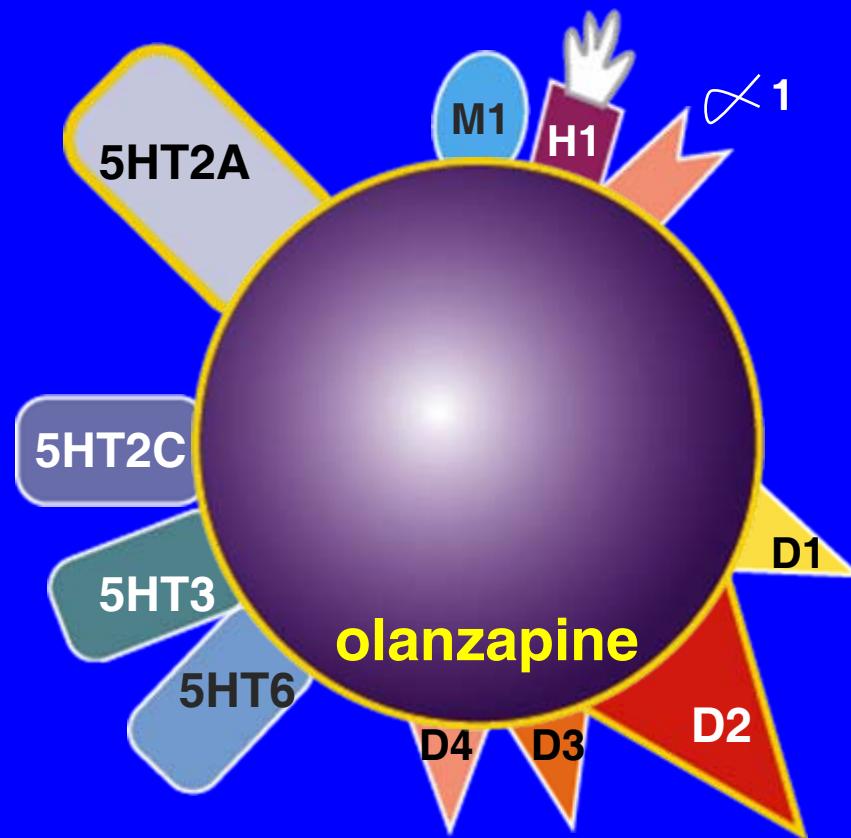
Newer atypicals

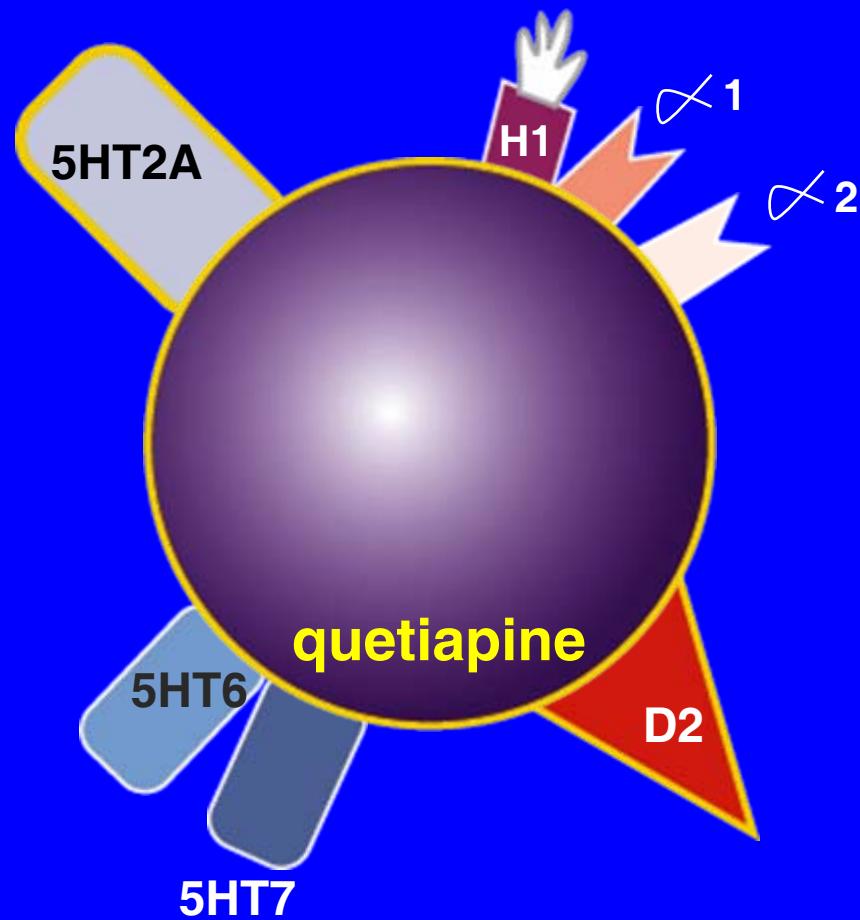
- More recently introduced drugs
- Some more dopamine selective
 - e.g. sulpiride, amisulpride
- Most recent dopamine partial agonist
 - e.g. aripiprazole
- Most combined 5-HT₂ and D₂ antagonists
 - e.g. risperidone, olanzapine, quetiapine
 - All pharmacologically differ

Clozapine – the archetypal atypical









Atypicals

- Now first line antipsychotics for schizophrenia and mania (+ other effects in bipolar disorder)
- Less EPS
 - quetiapine, olanzapine < risperidone
 - amisulpiride, sulpiride, aripiprazole < conventional antipsychotics
- Hyperprolactinaemia
 - Not with quetiapine and olanzapine
 - Yes with risperidone, amisulpiride and sulpiride
- Risk of weight gain, metabolic syndrome and diabetes
 - clozapine, olzapine > quetiapine, risperidone > amisulpiride, sulpiride, aripiprazole
- ? Differences in potency
 - Clozapine used for TRS, but risk of blood disorders
 - Olanzapine and risperidone possibly more potent than quetiapine in schizophrenia
 - Quetiapine has evidence bipolar depression
- Depot risperidone available
- Expensive

NICE Clinical Guideline 1

December 2002

**Core interventions in the
treatment and management
of schizophrenia in primary
and secondary care**

WWW.NICE.org.uk



*National Institute for
Clinical Excellence*

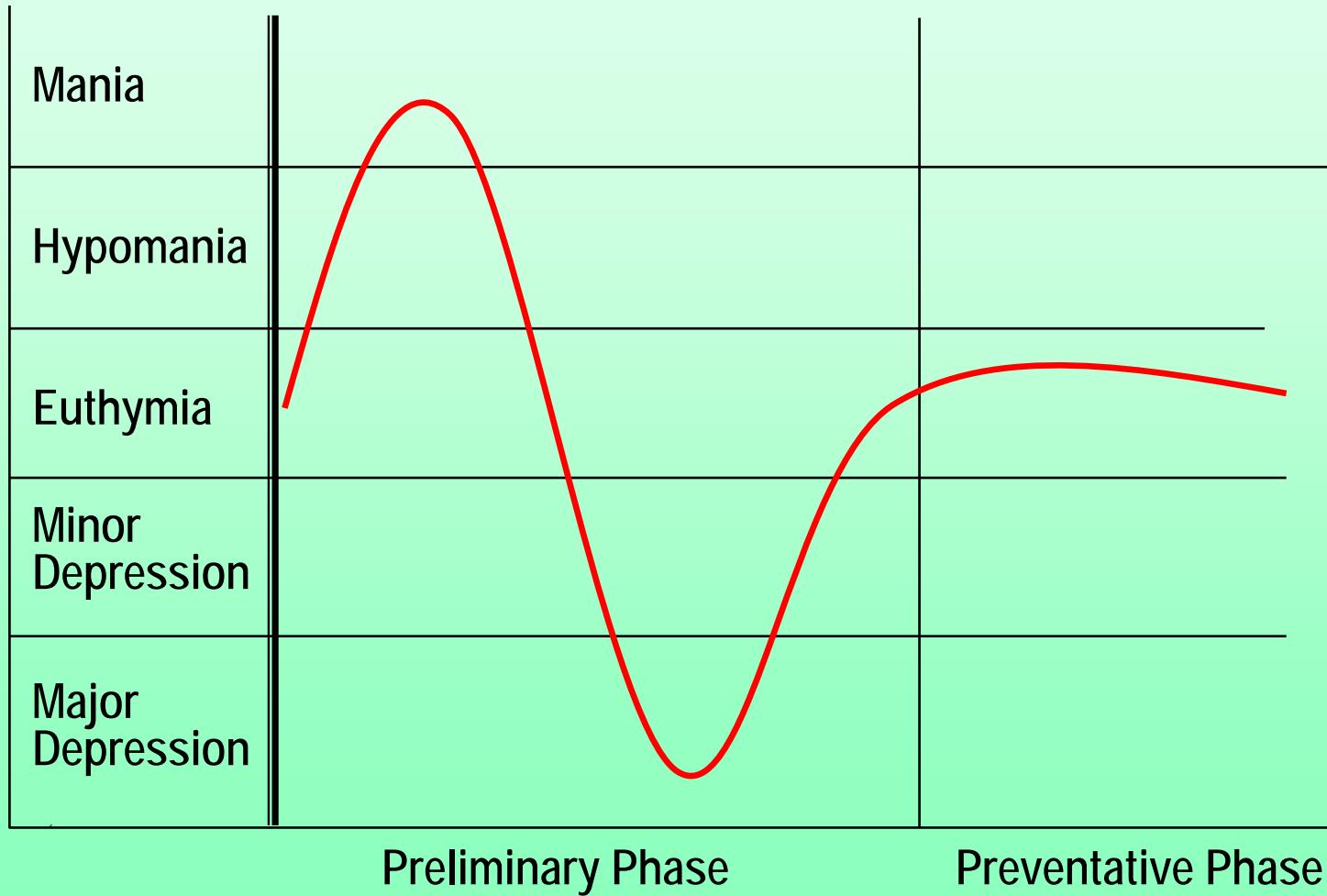
Acute Episode

- Informed choice by patient
 - If not able to do this then atypical
- If on typical and SEs are problematic or Sx control is inadequate, then atypical (otherwise remain on conventional)
- Single drug within BNF limits
 - Avoid high doses and loading doses
- Treatment trials should be for periods of 6-8 weeks
- Progress, SEs and user satisfaction should be monitored closely
- Treat for 1-2 years, withdraw slowly and monitor for 2 years after withdrawal

What is a “Mood Stabiliser”

- Treats depression plus mania without making either pole worse and/or has prophylactic effects for both mania and depression
- Absence of a consensus definition
- NICE
 - “The guideline avoids the term ‘mood stabiliser’, because there is no agreed definition. The terms ‘antimanic agent’ or ‘antimanic medication’ are used for treatment of an acute episode, and ‘prophylactic agent’ or ‘prophylactic medication’ for long-term maintenance treatment”

The course of Bipolar Disorder



NICE Clinical Guideline 38

July 2006

**Bipolar Disorder: The
management of bipolar
disorder in adults, children
and adolescents, in primary
and secondary care**

Acute Mania

- Atypical antipsychotic (olanzapine, risperidone, quetiapine) for those with severe mania
 - If ineffective consider adding Li or valproate
- Valproate or Li if previous good response and compliance
 - Avoid valproate in women of child bearing potential
 - Li only if less severe

Acute Depression

- First line: SSRI plus antimanic agent
- If on antimanic: SSRI or quetiapine (if not on antipsychotic)
- If recent unstable mood: avoid antidepressants – increase antimanic and consider lamotrigine
- Taper antidepressants after symptoms reduced for 8 weeks

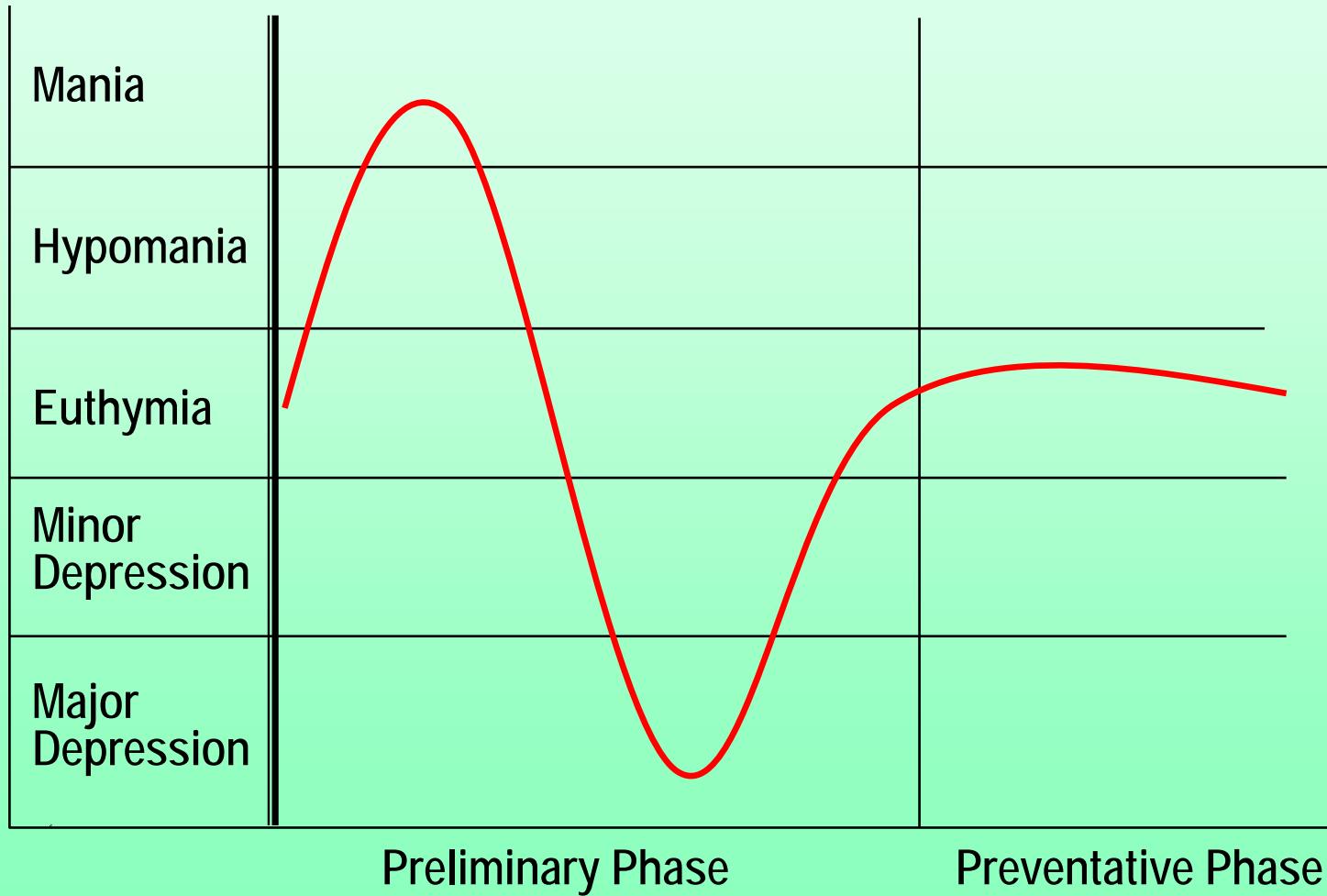
Long-term Treatment

- First line: lithium, olanzapine or valproate
- If fails monotherapy over 6 months
 - Li + valp, Li + olanz, Valp + olanz
- If combination fails
 - Consider lamotrigine (esp. freq. depressions), carbamazepine
- NOT antidepressants routinely (unless no mania X 5 yrs)
- Normally treat for at least 5 years

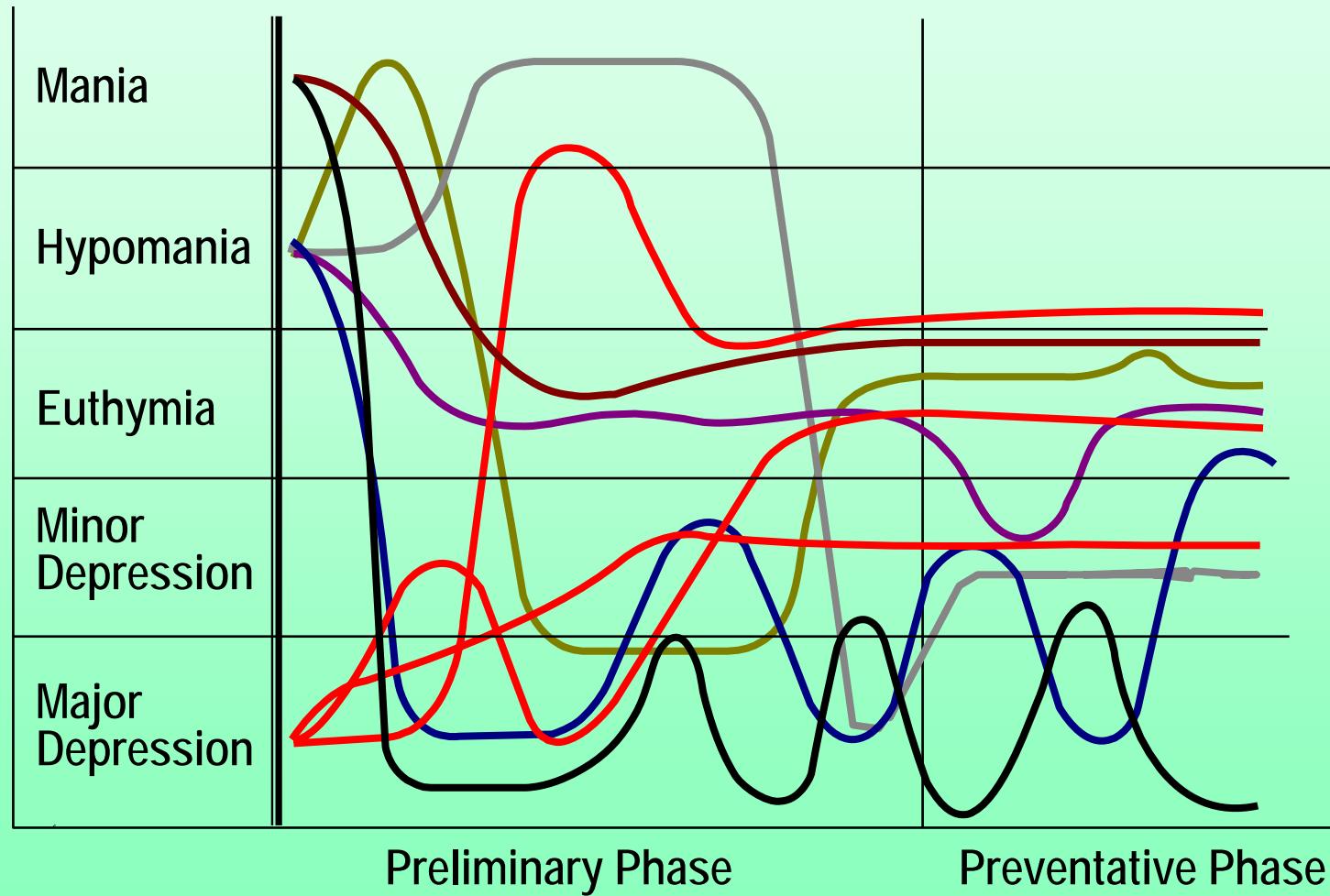
Bipolar treatments: some issues

- Lithium
 - Narrow therapeutic index – monitor levels 2 monthly
 - Renal and thyroid dysfunction – renal function + TFTs 6 monthly
 - Sudden discontinuation – 50% risk of mania
 - Teratogenic – Epsteins anomaly
- Valproate
 - Not for women under 18 or of child bearing potential
 - Teratogenicity (neural tube), polycystic ovary
 - Levels if ineffective, poor adherence or toxicity
- Lamotrigine
 - Risk of Stevens-Johnson syndrome
 - Slow dose titration

The course of Bipolar Disorder



The course of Bipolar Disorder



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Clinical use of Drugs in Psychiatry

- Fundamental principles:
 - Assessment of risk/benefits
 - Consideration of costs
 - Full discussion with patient
 - Informed choice by patient
 - Repeated monitoring and re-assessment
 - Integration with other treatments

Conclusions

- Drugs are often the first line treatment in psychiatric illness
- However drugs are not the only mode of treatment
- Good evidence supports their use
- The use of psychotropics is complicated by:
 - professional perceptions of illness
 - patient perceptions of treatment
 - Complexity of their pharmacology
- “Normal” good clinical practice is essential