

# Old Age Module

A large, stylized orange bracket that spans across the width of the slide, positioned below the main title.

## Older People with Psychosis

Developing people  
for health and  
healthcare

## **Expert Led Session**

# **Older People with Psychosis**

Dr Salman Karim

Consultant Old Age Psychiatrist & Honorary  
Senior Lecturer

Lancashire Care NHS Foundation Trust

# Epidemiology of Psychosis in Elderly

- 20% of patients with schizophrenia (SZ) have their first onset at or after the age of 40, i.e. late-onset psychosis (LOP) (Maglione, Thomas, & Jeste, 2014)
- 0.1-0.5% prevalence of schizophrenia in >65 (Castle, 1993 and Copeland 1998)
- Estimated incidence of schizophrenia after the age of 65 years is 7.5 per 100,000 person-years (Stafford, 2018)

## Increased risk of psychosis in elderly

- Age-related changes in fronto-temporal cortices
- Neuro-chemical changes with ageing
- Sensory deficits
- Cognitive decline - significantly greater risk of developing dementia [relative risk (RR) 2.29; 95% confidence interval (CI) 1.35–3.88] (Palmer, 2001)
- Social isolation
- Polypharmacy

# Consequences of psychotic symptoms

- Disruptive and aggressive behaviour
- Neglect and abuse
- Carer distress
- Institutionalisation
- Financial burden

(Schneider et al, 1997 ; Stern, 1997)

# Psychosis in the elderly

## Organic:

- Delirium
- Dementias
- Approx 60% of psychosis in elderly is *secondary* to other disorders (Holroyd S, 1999 and Manepalli, 2007)

## Functional:

- Schizophrenia
- Affective Disorder
- Delusional Disorder

# Schizophrenia – historical perspective

- Kraepelin (1894)
  - Dementia Praecox (disorder of emotion/volition)
  - Paraphrenia (insidious delusional system)
- Bleuler (1911) - Schizophrenia
- Bleuler (1943) - Late onset schizophrenia (onset after age 40)
- Roth and Morrisey (1952) - Late paraphrenia (onset after age of 55)

# SCIZOPHRENIA IN THE ELDERLY

## International Consensus Classification

- Chronic Schizophrenia (graduates)
- Late Onset Schizophrenia (onset after age 40)
- Very Late onset Schizophrenia (onset after age 60)

(Howard et al, 2000)



# Schizophrenia in the elderly

- Overall community prevalence – 0.1- 0.5%
- **Chronic schizophrenia** – 85% of the total
- **Late onset schizophrenia** – 23.5% develop the illness after age of 40
- **Very late onset** – 4% develop the illness after age 60

(Howard et al, 2000 ; Harris & Jeste, 1998)

# Early & Late-Onset Schizophrenia

## Similarities:

- **Genetic** risk
- Presence and severity of **positive symptoms**
- Early psycho-social maladjustments
- **Subtle brain abnormalities**

## Differences:

- **Fewer negative symptoms**
- **Better neuropsychological performance**
- **Better response to antipsychotics**

(Howard et al, 2004 ; Palmer et al, 2001)

# Very-late onset schizophrenia

## **Higher likelihood/risk:**

- Female gender
- Associated sensory impairment
- Social isolation
- Tardive dyskinesia

## **Lesser likelihood/risk:**

- Formal thought disorder
- Affective blunting
- Family history

(Lisa et al, 2002 ; Tune & Salzman, 2003)

# Comparison by age of onset

	Early onset	Late onset	Very late onset
Age of onset	<40	40-60	>60
Paranoid subtype	Common	Very common	Common
Negative symptoms	Marked	Present	Absent
Thought disorder	Present	Present	Absent
Organic brain pathology	Absent	Absent	Present
Family history	Present	Present	Absent
Childhood maladjustment	Present	Present	Absent
Cognitive impairment	Present	Present	Progressive
Information retention	Normal	Normal	Impaired?
Risk of tardive dyskinesia	Present	Present	Marked
Antipsychotic dose	High	Low	Lower

# Very late onset

## Higher likelihood/risk:

**Female** gender

Associated **sensory impairment**

**Social isolation**

**Tardive dyskinesia**

## Lesser likelihood/risk:

**Formal thought disorder**

Affective **blunting**

**Family history**

(Lisa et al, 2002 ; Tune & Salzman, 2003)

# Biology of schizophrenia in elderly

## **Female Gender:**

Higher brain volume loss in parietal lobes

Excess of dopamine receptors

Loss of anti-dopamine action of oestrogens?

(Jeste et al, 1997; Madhusoondanan et al, 2000)

# Biology of schizophrenia in elderly

## Hypothesis 1:

Genetic susceptibility

Neuronal loss due to aging/vascular changes

Manifestation of symptoms

## Hypothesis 2:

No genetic risk

Single event (vascular?) precipitating symptoms

(Karim& Burns, 2003; Pearlson G, 1995)

# Clinical Features

Reduction of positive symptoms “burning out”

High levels of depression:

**2 out of 5** clinically depressed  
physical problems, poor support

**Smoking** rate twice than general population

Alcohol consumption lower in older people

(Bridge et al, 1978; Adler,1995)



# Clinical features

## Physical problems unrecognized in 50%

Psychiatrists miss half of the physical problems

Higher rates of IHD, Diabetes, respiratory problems, peptic ulcers.

(Koran, 1989; Koryni, 1979; Karim et al, 2006)

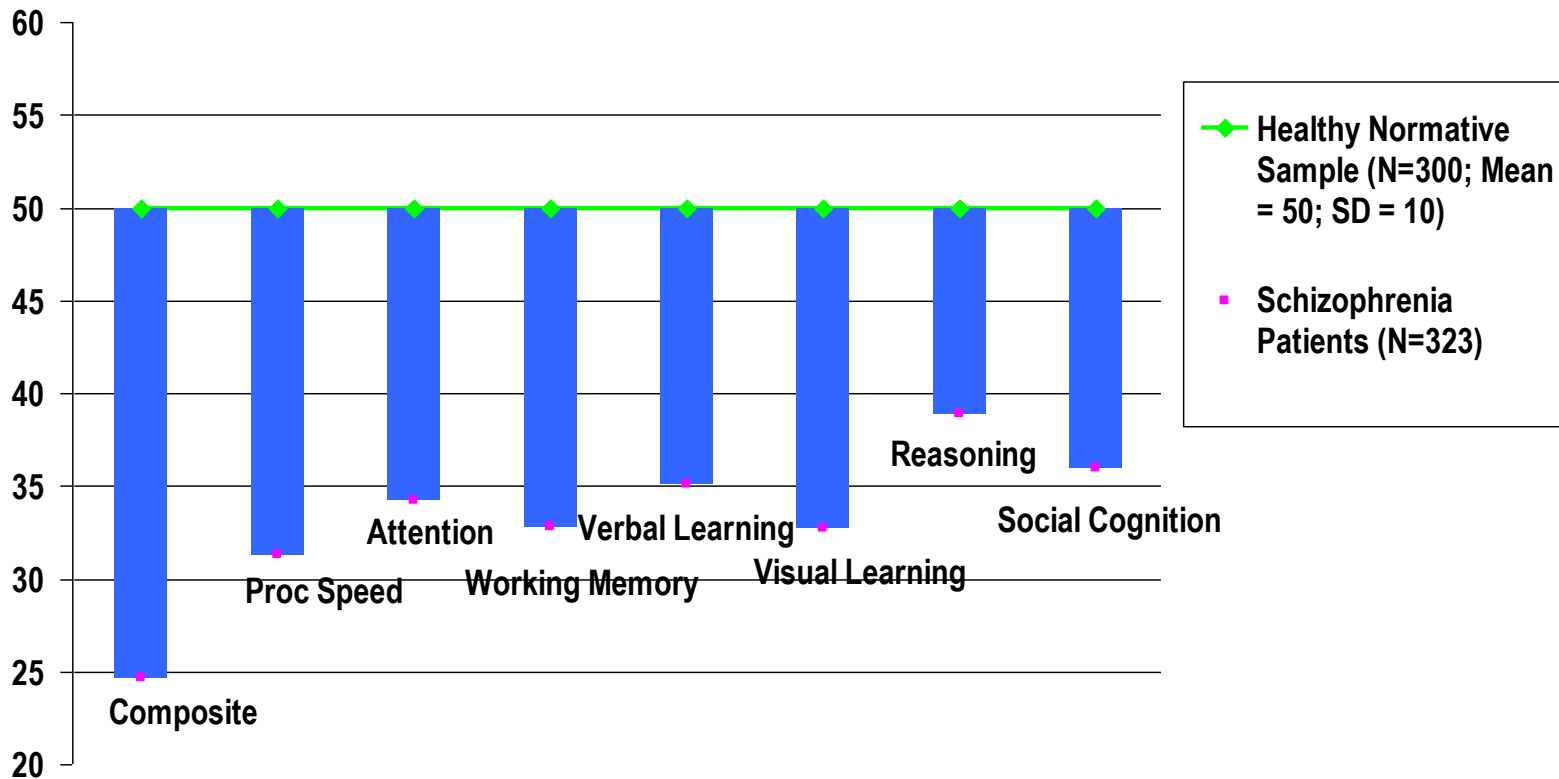
# Cognitive Deficits

- Predictor of poor outcome
- Specific deficits:
  - Use of Language
  - Executive functioning
  - Memory
- Comparison with AD (Gabrovska et al, 2002)
  - More impaired on visuo-spatial task
  - Less impaired on verbal
  - Right hemisphere atrophy on MR

# Cognitive Deficits

- Role of antipsychotics?
- Cerebrovascular disease?
- Treatment Implications:
  - Failure of social rehabilitation
  - Poor community living skills
  - Poor self care
  - Higher numbers in nursing homes

# Cognitive Deficits

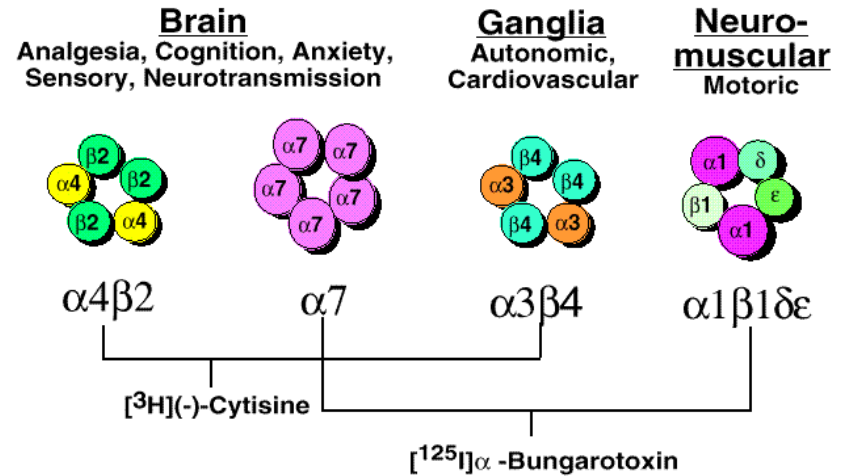


## Neuronal Nicotinic Receptors – cholinergic hypothesis

Growing support for aberrant nicotinic, cholinergic signalling in psychosis

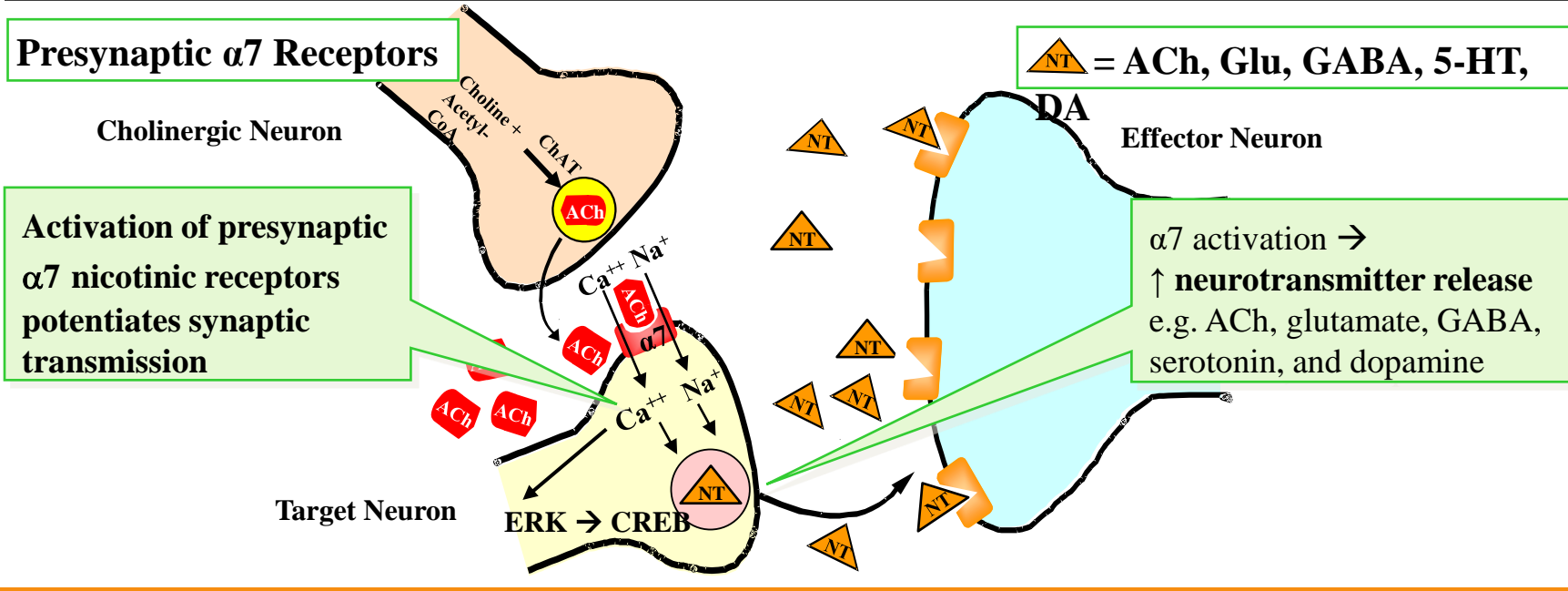
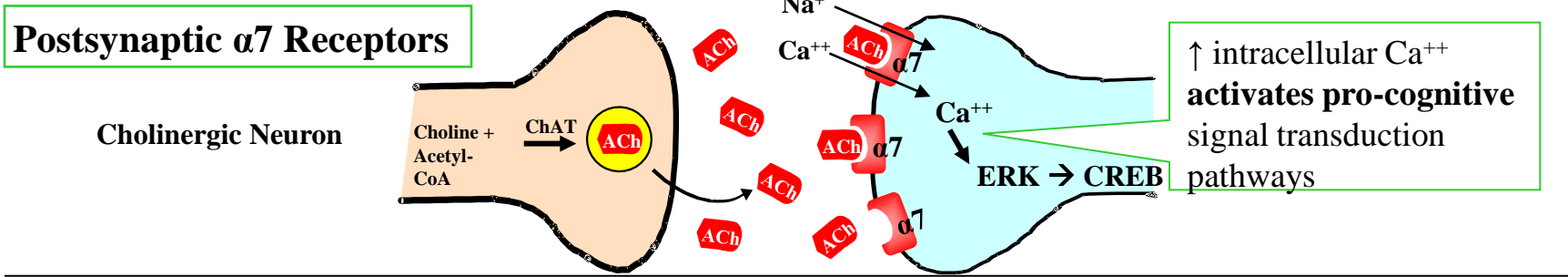
Cause of the cognitive deficits observed in schizophrenia

Low availability of the  $\alpha 7$ -nAChR in Hippocampus may be linked to recent onset of psychosis



Future - novel classes of  $\alpha 7$ -nAChR-modulating drugs acting in the hippocampus?

# α7 Receptors: Pre- and Postsynaptic Mechanisms



Activation of presynaptic α7 nicotinic receptors potentiates synaptic transmission

α7 activation → ↑ neurotransmitter release e.g. ACh, glutamate, GABA, serotonin, and dopamine

# Social disabilities

- Improved coping skills
- Deficits in daily functioning in higher domains
- Predictors of abnormal functioning:
  - Cognitive impairment
  - Negative symptoms
  - Movement disorders

(Cohen,1993; Klaplow et al,1997; Cohen et al, 2000)

# Management

## Typical antipsychotics:

- Higher risk of TD
- 37% higher risk of death; risk is dose dependent
- Effective in treating positive symptoms
- Higher risk of disabling side effects

## Atypical antipsychotics:

- Better side effect profile
- Better at treating negative symptoms

Essali A, Ali G: Antipsychotic drug treatment for elderly people with late-onset schizophrenia (Review); 2013 ;The Cochrane Collaboration  
(Correll et al, 2004; Wang et al,2005; Nasrallah,2006)



# Management

## Atypical Antipsychotics:

### Recommended 1st line treatment

#### Risperidone and Olanzapine:

- Most extensively studied in elderly
- Both equally effective
- Fewer adverse events than typicals
- Risk of EPS higher with Risperidone
- Improvement with switching from typical

(Ritchie et al, 2006 & 2003; Barak et al, 2004)

- ATLAS trial: **low-dose amisulpride** is effective and well tolerated as a treatment for VLOSLP (Howard, R 2018)

# Service Needs

85% reside in community

Service utilization comparable to AD (Shaw et al, 2000)

Concern about standard of services (McNulty et al, 2003):

- Low spending

- Shortage of consultants

- Lack of policy

# Old Age Module

MCQs

# Old Age Module

## MCQs

1. **A 76 year old lady is diagnosed with ‘late paraphrenia’. Which of the following delusions is the GP most likely to find compared to younger adults?**
  - A. Hypochondriacal
  - B. Delusions of misidentification
  - C. Religious delusions
  - D. Delusions of reference
  - E. Persecutory delusions

# Old Age Module

## MCQs

- 1. A 76 year old lady is diagnosed with ‘late paraphrenia’. Which of the following delusions is the GP most likely to find compared to younger adults?**

  - Hypochondriacal
  - Delusions of misidentification
  - Religious delusions
  - Delusions of reference
  - Persecutory delusions**

# Old Age Module

## MCQs

2. **Very late onset schizophrenia is characterised by onset after:**
- A. 40 years
  - B. 60 years
  - C. 65 years
  - D. 70 years
  - E. 80 years

# Old Age Module

## MCQs

2. **Very late onset schizophrenia is characterised by onset after:**
- A. 40 years
  - B. 60 years**
  - C. 65 years
  - D. 70 years
  - E. 80 years

# Old Age Module

## MCQs

- 3. Which antipsychotic is most likely to cause postural hypotension:**
- A. Aripiprazole
  - B. Risperidone
  - C. Haloperidol
  - D. Quetiapine
  - E. Sulpiride



# Old Age Module

## MCQs

3. Which antipsychotic is most likely to cause postural hypotension:
- A. Aripiprazole
  - B. Risperidone
  - C. Haloperidol
  - D. **Quetiapine**
  - E. Sulpiride

# Old Age Module

## MCQs

4. Which of the following drugs should not be used in renal failure?
- A. Amisulpride
  - B. Aripiprazole
  - C. Chlorpromazine
  - D. Olanzapine
  - E. Quetiapine

# Old Age Module

## MCQs

4. Which of the following drugs should not be used in renal failure?
- A. Amisulpride
  - B. Aripiprazole
  - C. Chlorpromazine
  - D. Olanzapine
  - E. Quetiapine

# Old Age Module

## MCQs

5. **‘Sensitivity to antipsychotics’ is linked to which disorder?**
- A. Alzheimer’s Disease
  - B. Dementia with Lewy Bodies
  - C. Late onset Schizophrenia
  - D. Organic mood disorder
  - E. Huntington’s Disease

# Old Age Module

## MCQs

5. **‘Sensitivity to antipsychotics’ is linked to which disorder?**
- A. Alzheimer’s Disease
  - B. Dementia with Lewy Bodies**
  - C. Late onset Schizophrenia
  - D. Organic mood disorder
  - E. Huntington’s Disease

# Old Age Module

Any Questions?

Thank you

Please provide feedback/suggestions on this presentation to the module lead [Anthony.Peter@lancashirecare.nhs.uk](mailto:Anthony.Peter@lancashirecare.nhs.uk)