

Year 6 Integrated Medicine Course:
Respiratory Lecture 2:
Management of common airway diseases

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Year 6 Integrated Medicine Course

Respiratory Lectures

- **Lecture 1:**
Diagnosis and approaches to the breathless patient
- **Lecture 2:**
Management of Common Airway Diseases
- **Lecture 3:**
Less Common Lung Diseases and preparation for Paces

Year 6 Integrated Medicine Course

Respiratory Lectures

- This is revision
- Most already covered in Year 3
- E Learning Modules are available to cover this ground
- All available from the Year 6 page on the Undergraduate intranet

Imperial College
London



Respiratory Medicine

Asthma

Enter

Imperial College
London



Respiratory Medicine

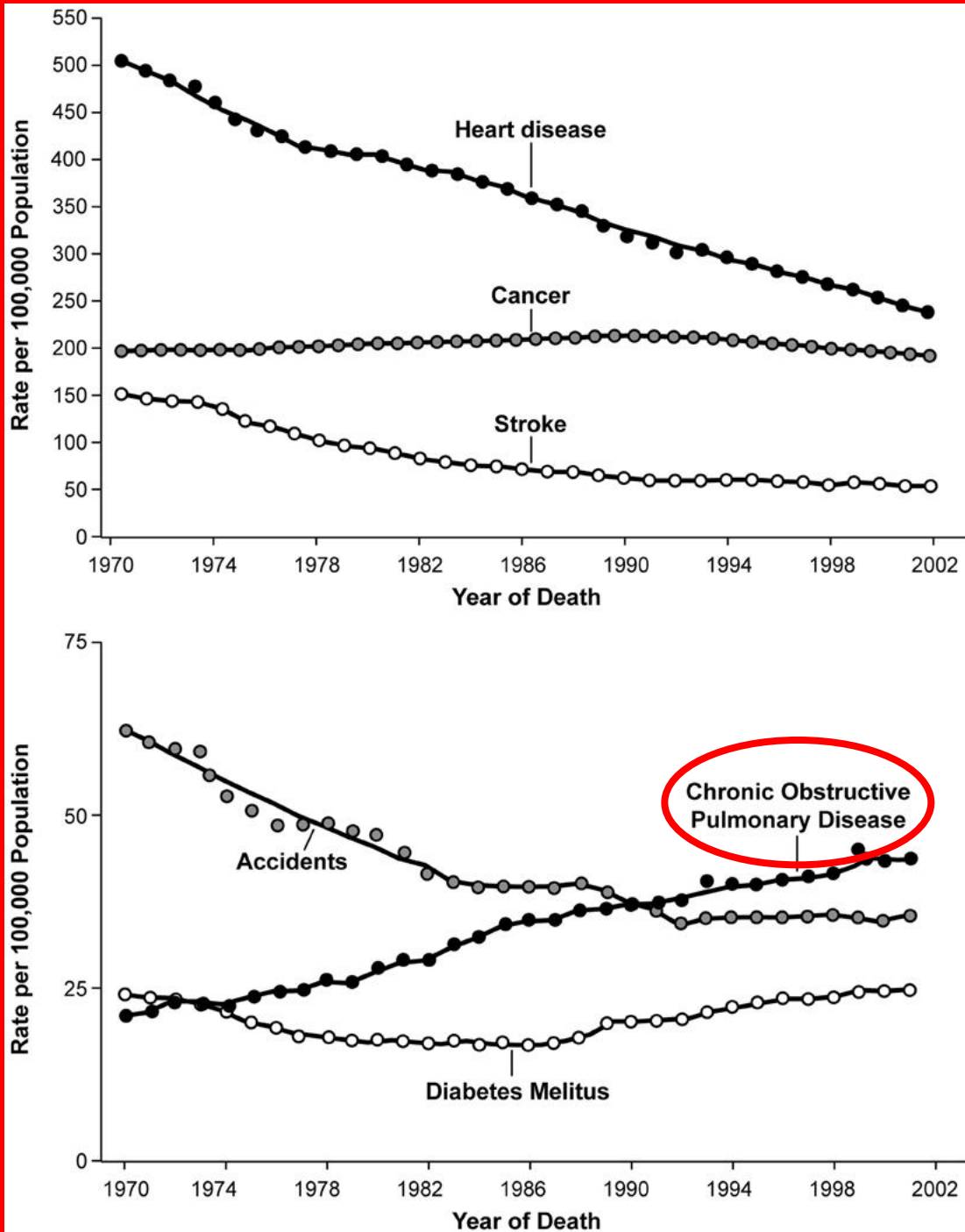
COPD

Enter

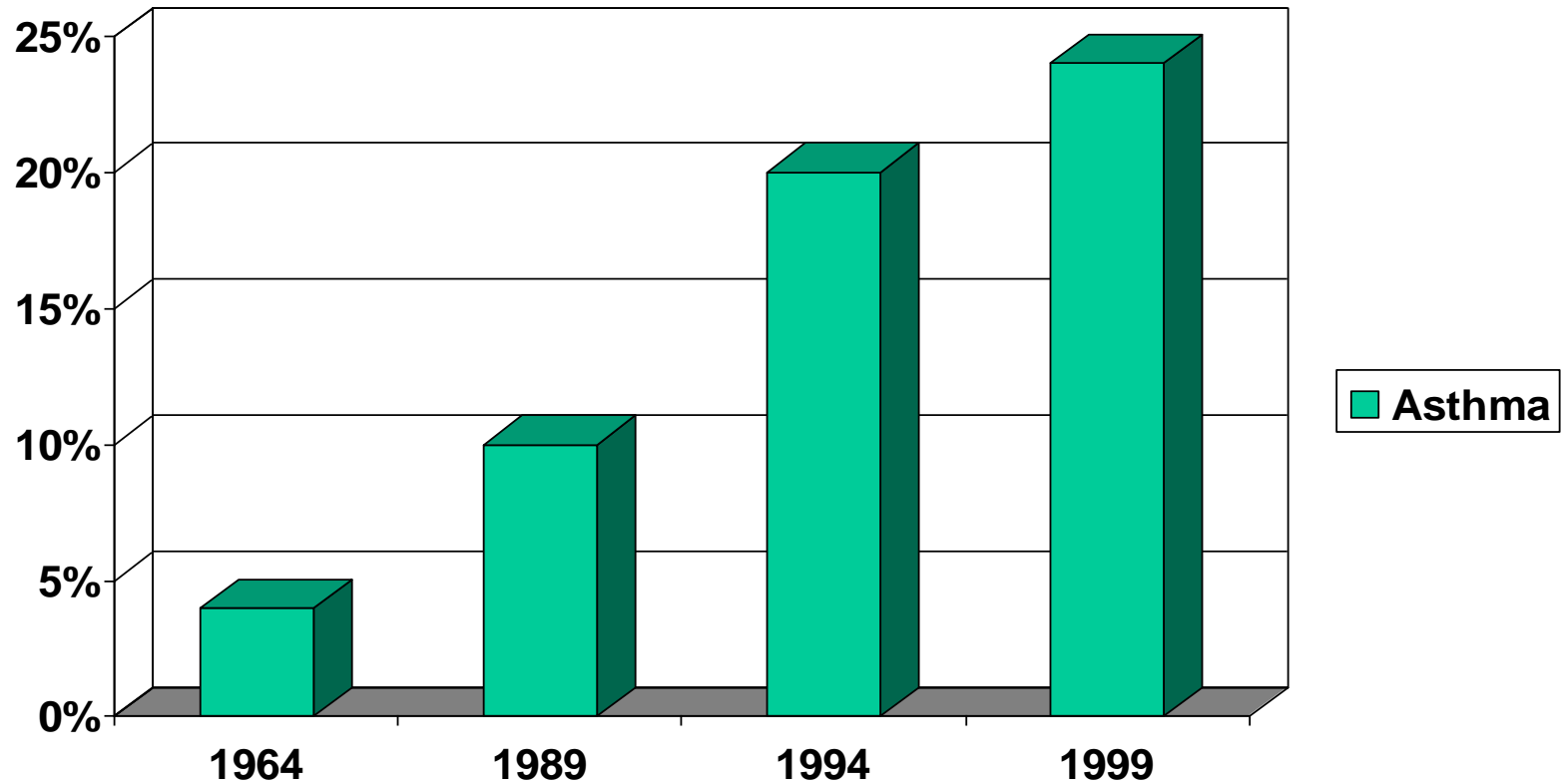


Of the six leading causes of death in the United States, only COPD has been increasing steadily since 1970

Source: Jemal A. et al. *JAMA* 2005



Increasing prevalence of asthma (9-12 year olds)



Devenny A et al BMJ 2004 329: 489-90

A Venn diagram illustrating the relationship between four respiratory conditions. It consists of four overlapping circles. The circle on the far left is dashed and labeled 'Asthma'. The other three circles are solid and labeled 'Chronic Bronchitis', 'Emphysema', and 'Obstructive Airway Disease'. The 'Obstructive Airway Disease' circle is the largest and overlaps with the other three. The 'Chronic Bronchitis' and 'Emphysema' circles overlap each other and both overlap with the 'Obstructive Airway Disease' circle. The 'Asthma' circle overlaps with the 'Chronic Bronchitis' circle.

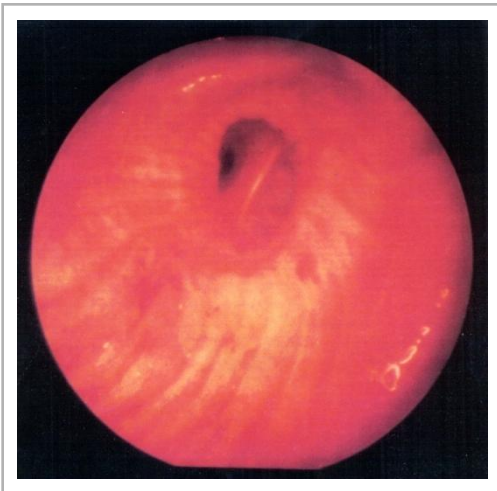
Asthma

Chronic
Bronchitis

Emphysema

Obstructive Airway
Disease

Asthma



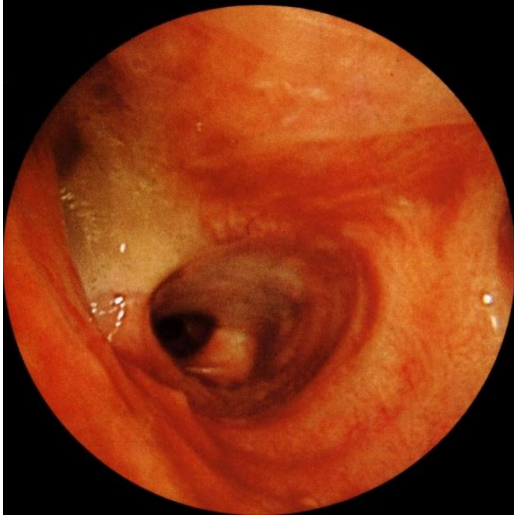
- **chronic inflammatory disorder of the airways ... associated with airway hyper-responsiveness ... and variable, airflow obstruction within the lung**

Asthma



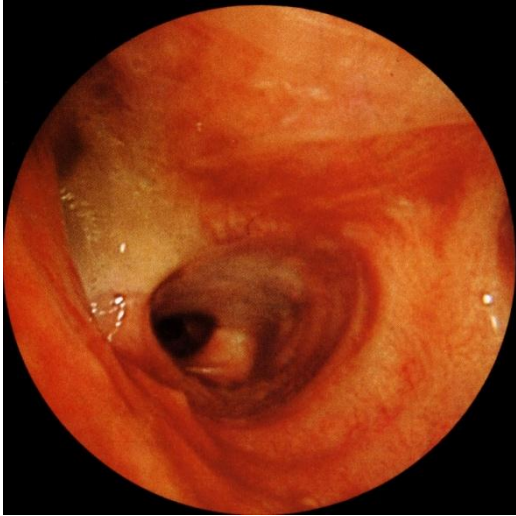
- **chronic inflammatory disorder of the airways ... associated with airway hyper-responsiveness ... and **variable, airflow obstruction** within the lung**

C O P D



- COPD is a preventable and treatable disease with some significant extra-pulmonary effects that may contribute to the severity in individual patients
- Pulmonary component characterized by airflow limitation that is not fully reversible
- Airflow limitation usually progressive and associated with an abnormal inflammatory response of the lung to noxious particles or gases

C O P D

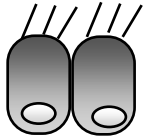


- COPD is a preventable and treatable disease with some significant extra-pulmonary effects that may contribute to the severity in individual patients
- Pulmonary component characterized by **airflow limitation that is not fully reversible**
- Airflow limitation **usually progressive** and associated with an abnormal inflammatory response of the lung to noxious particles or gases

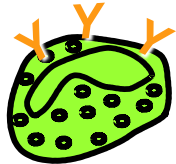


ASTHMA

Allergens



Ep cells



Mast cell



CD4+ cell
(Th2)

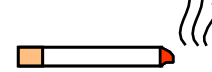


Eosinophil

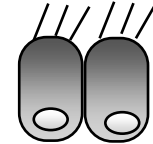
Bronchoconstriction
AHR

COPD

Cigarette smoke



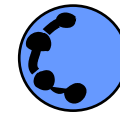
Alv macrophage



Ep cells



CD8+ cell
(Tc1)



Neutrophil

Small airway narrowing
Alveolar destruction

Airflow Limitation

Reversible

Irreversible

Symptoms

Asthma

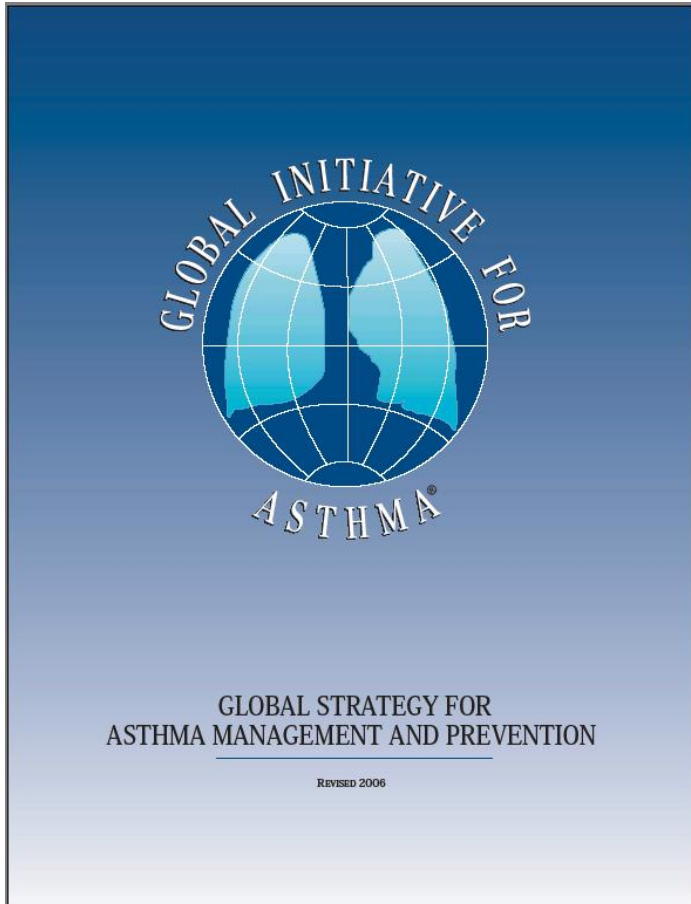
- Variable breathlessness and/or chest tightness
- Wheezing
- Symptoms awaken patient from sleep or are worse in the early morning
- When bad, short of breath at rest or on exertion
- May be completely free of symptoms between attacks

COPD

- Continual symptoms
- Breathless usually on exertion; until advanced often ok at rest
- Cough
- Sputum
- Progressive

Remember last weeks lessons re objective confirmation of diagnosis

Guideline goals for successful asthma management



- Achieve and maintain control of symptoms
- Maintain normal activity levels – including exercise
- Maintain pulmonary function close to normal levels
- Avoid adverse effects from asthma medications
- Prevent asthma exacerbations
- Prevent asthma mortality

Guideline goals for successful asthma management



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Guideline goals for successful asthma management



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- **Prevent asthma exacerbations**
- **Prevent asthma mortality**

Levels of Asthma Control

Characteristic	Controlled (All of the following)	Partly Controlled (Any measure present in any week)	Uncontrolled
Daytime symptoms	None (twice or less/week)	More than twice/week	Three or more features of partly controlled asthma present
Limitations of activities	None	Any	
Nocturnal symptoms/awakening	None	Any	
Need for reliever/rescue treatment	None (twice or less/week)	More than twice/week	
Lung function (PEF or FEV ₁)‡	Normal	< 80% predicted or personal best (if known)	
Exacerbations	None	One or more/year*	
			One in any week†

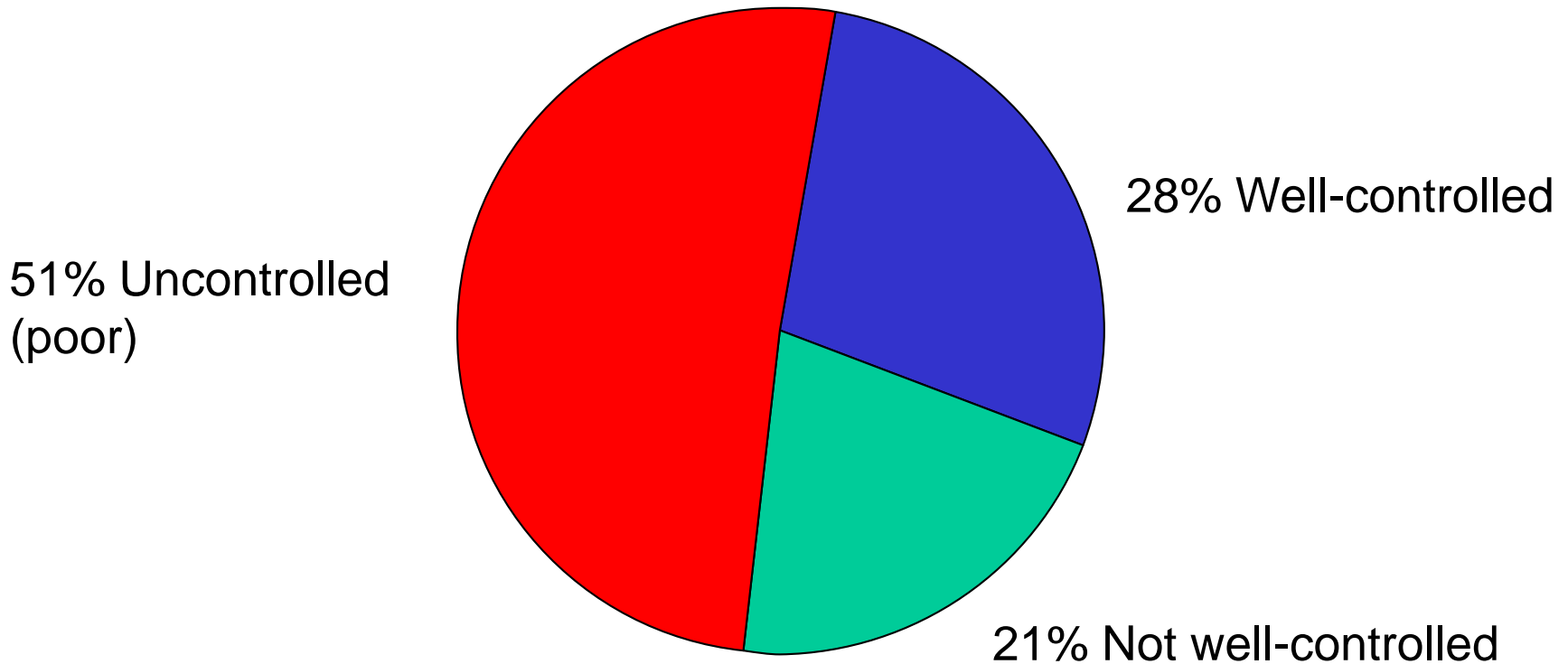
* Any exacerbation should prompt review of maintenance treatment to ensure that it is adequate.

† By definition, an exacerbation in any week makes that an uncontrolled asthma week.

‡ Lung function is not a reliable test for children 5 years and younger.

Despite ICS or ICS/LABA therapy, only 28% of patients were well-controlled according to the ACQ

Partridge MR, BMC Pulmonary Medicine 2006

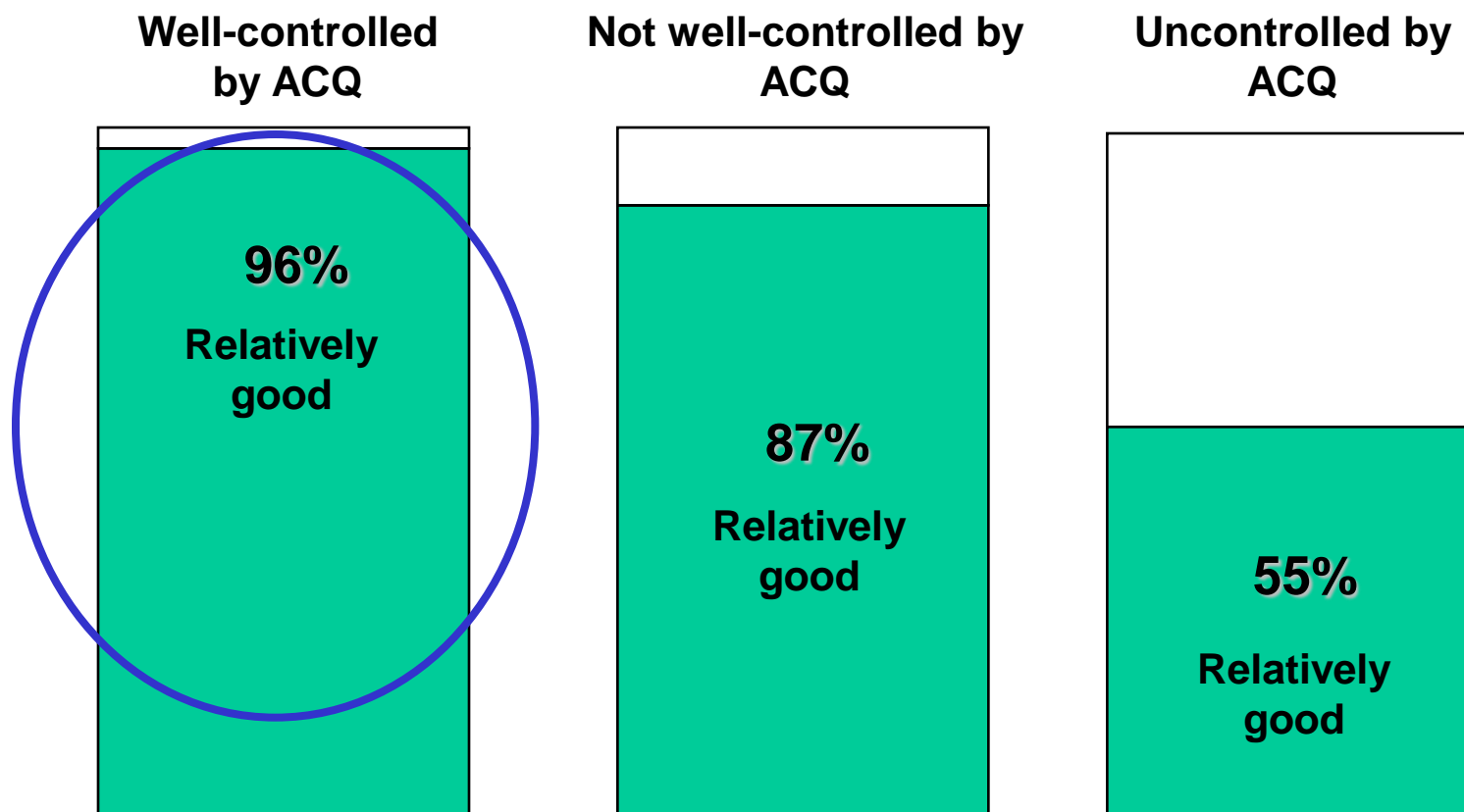


ACQ-6 Summary Score

Well-controlled:	0.0 to 0.74
Not well controlled:	0.75 to 1.5
Uncontrolled:	1.5+

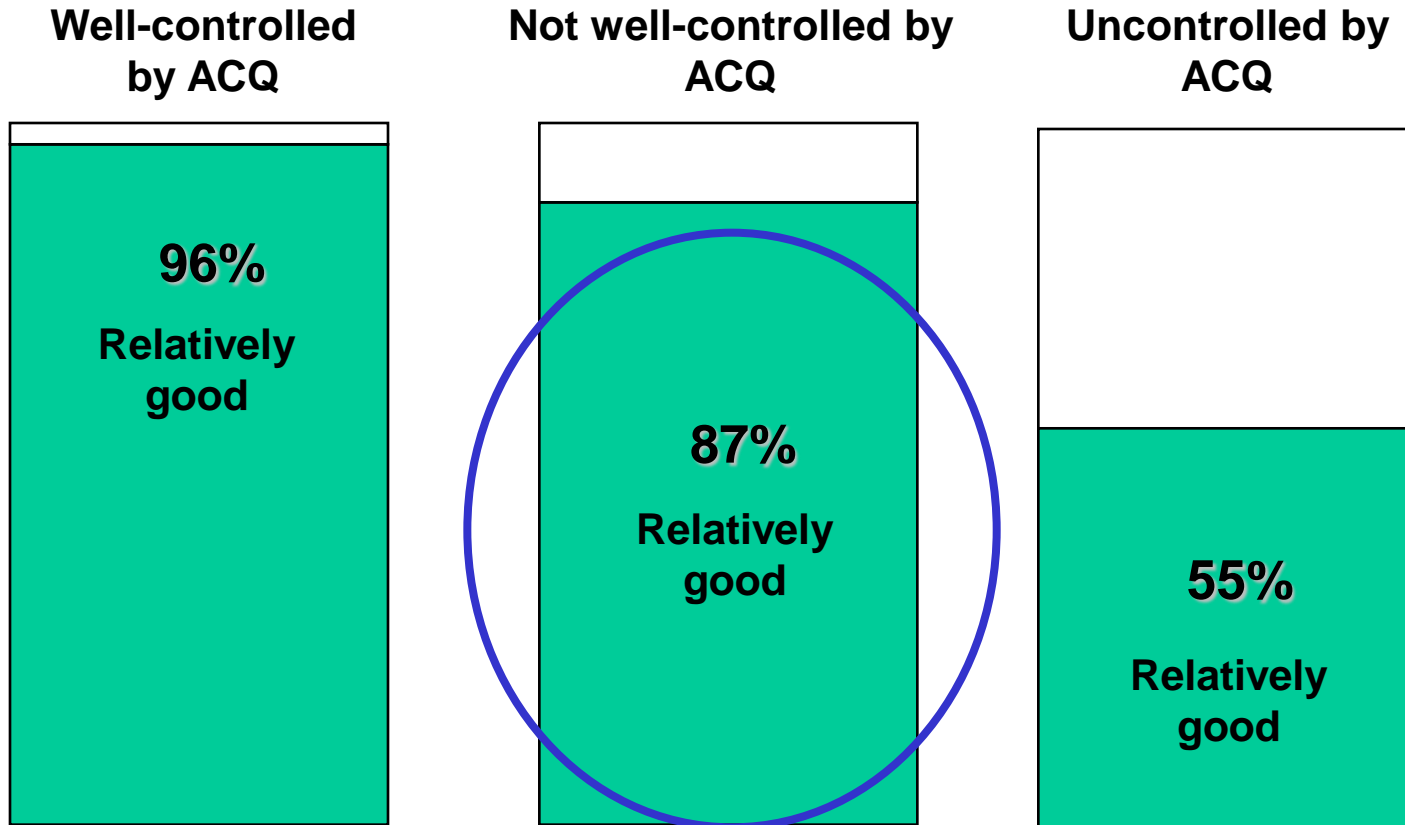
Most patients perceive their asthma to be better controlled than the ACQ showed


Proportion of patients reporting disease 'relatively good' in the past week:



Most patients perceive their asthma to be better controlled than the ACQ showed

Proportion of patients reporting disease 'relatively good' in the past week:






*But if you had
asked me about
last week.....*

Fine!

How are you?



*Great! Next
Patient
Please!*



Assessment: Royal College of Physicians of London three questions

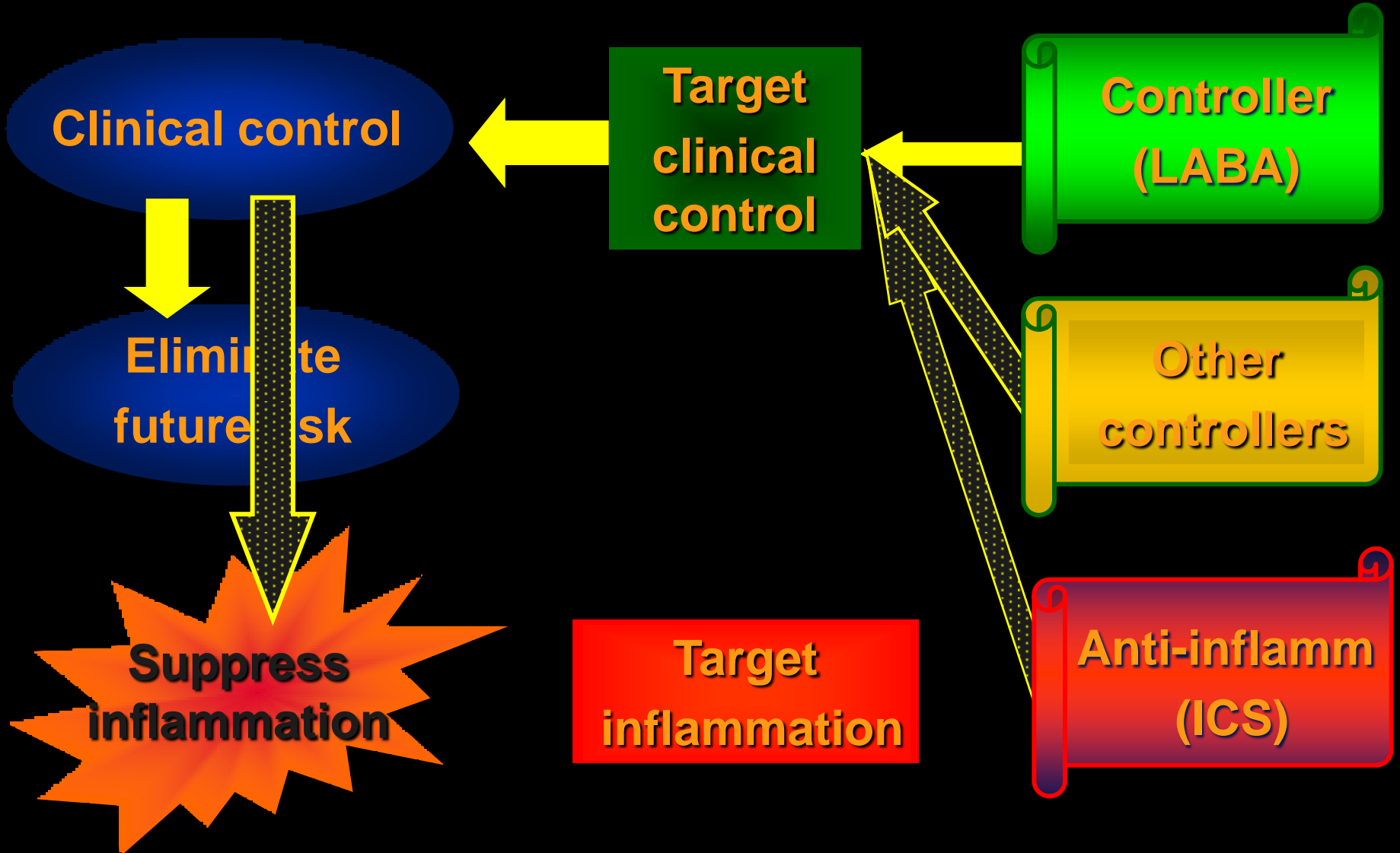
IN THE LAST WEEK / MONTH		
	YES	NO
“Have you had difficulty sleeping because of your asthma symptoms (including cough)?”	<input type="checkbox"/>	<input type="checkbox"/>
“Have you had your usual asthma symptoms during the day (cough, wheeze, chest tightness or breathlessness)?”	<input type="checkbox"/>	<input type="checkbox"/>
“Has your asthma interfered with your usual activities (e.g. housework, work, school, etc)?”	<input type="checkbox"/>	<input type="checkbox"/>
Date ____ / ____ / ____		

- Applies to all patients with asthma aged 16 and over.
- Only use after diagnosis has been established.

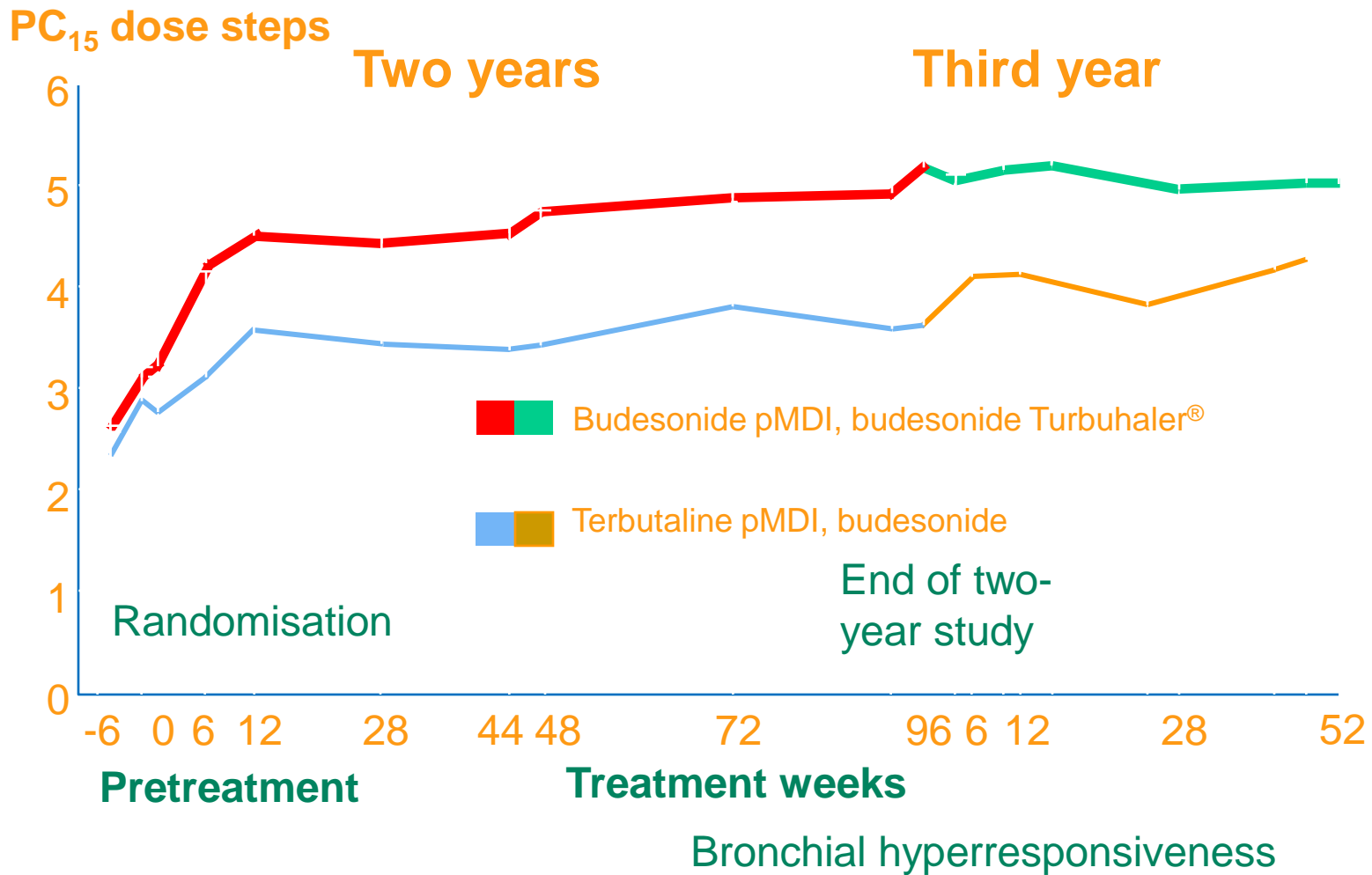
TARGET

STRATEGY

TREATMENT



The importance of inhaled steroids in maintaining optimal airway function.

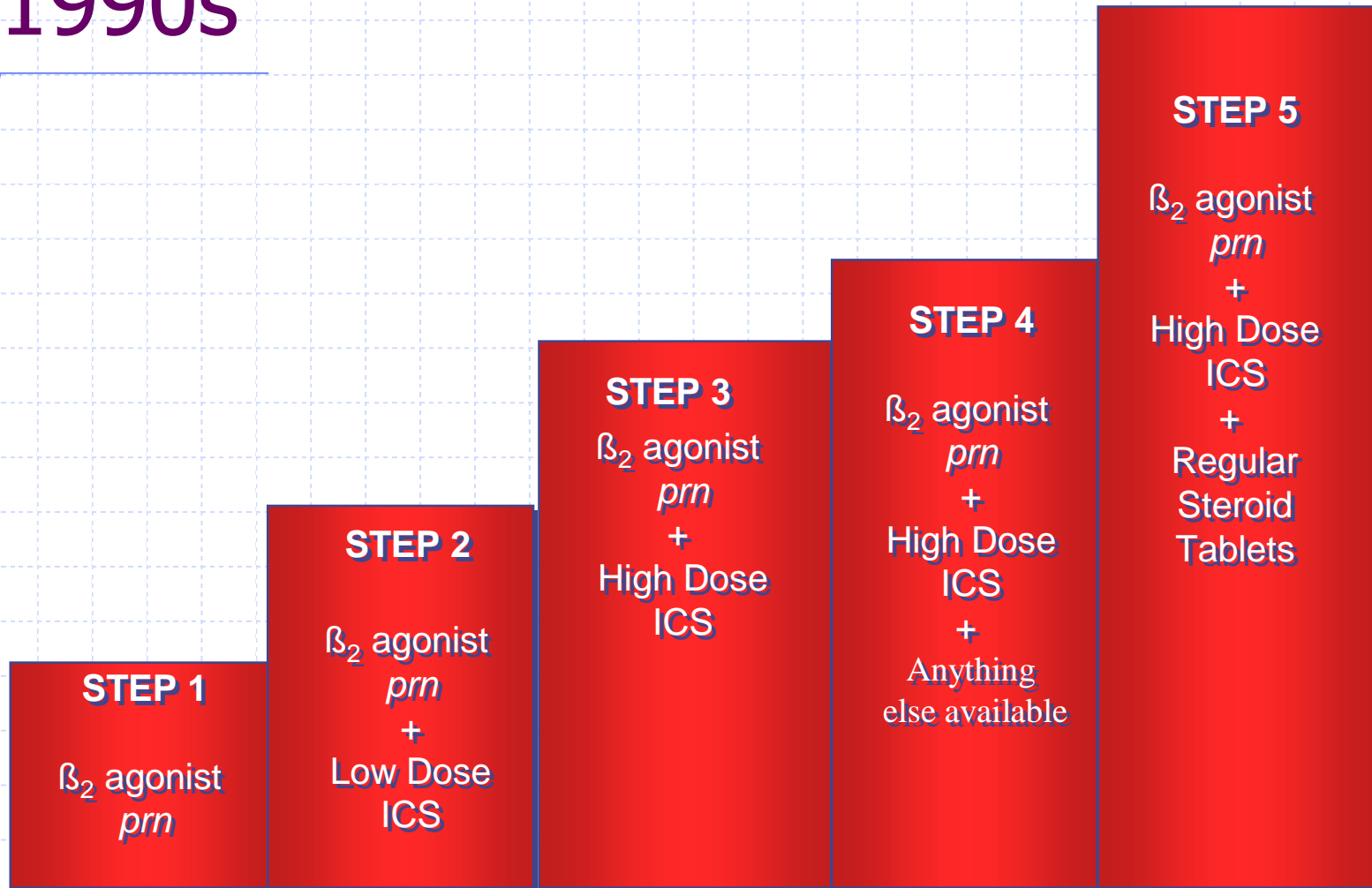


Guideline goals for successful asthma management

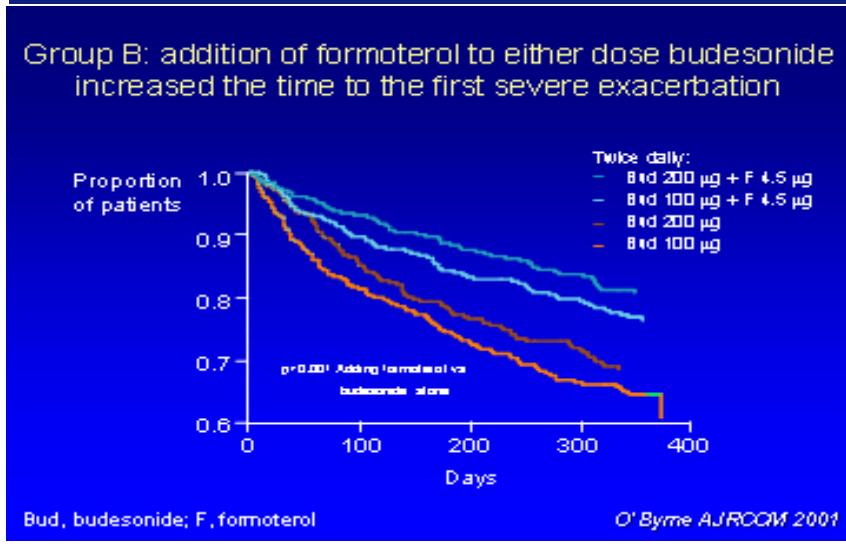
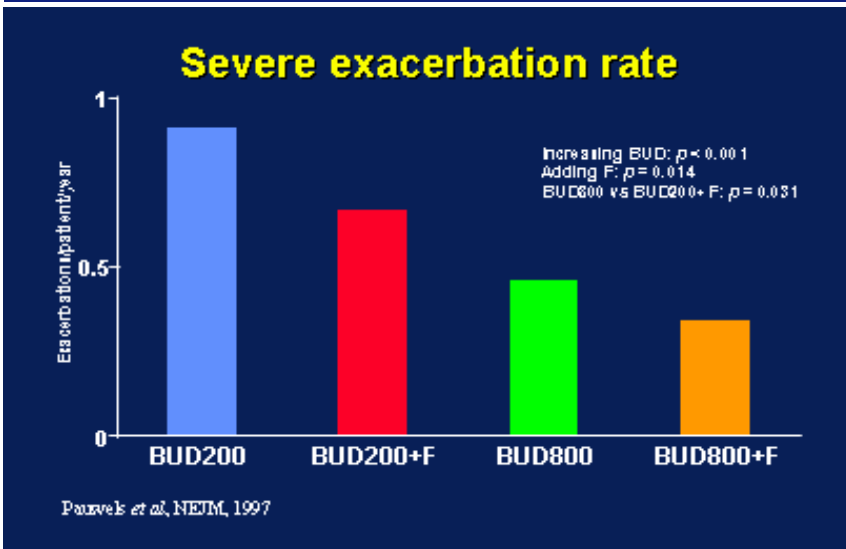
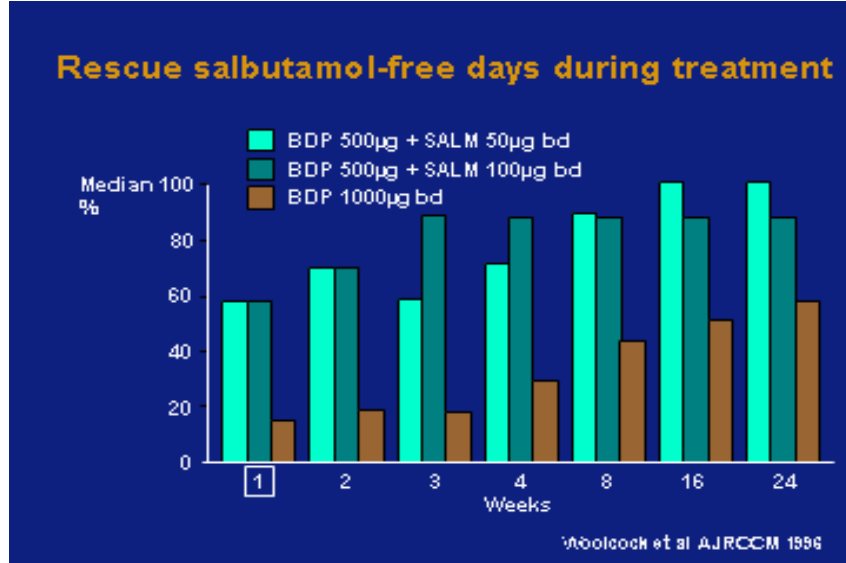
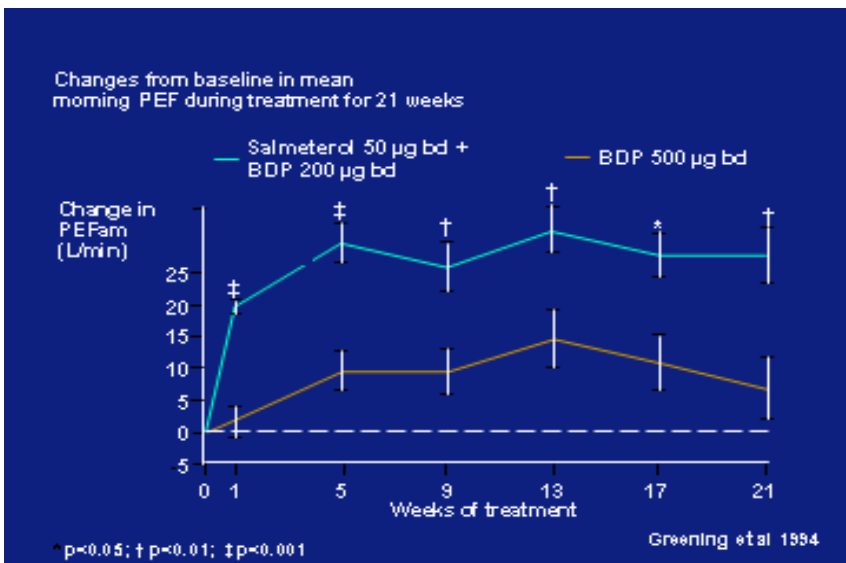


- Achieve and maintain control of symptoms
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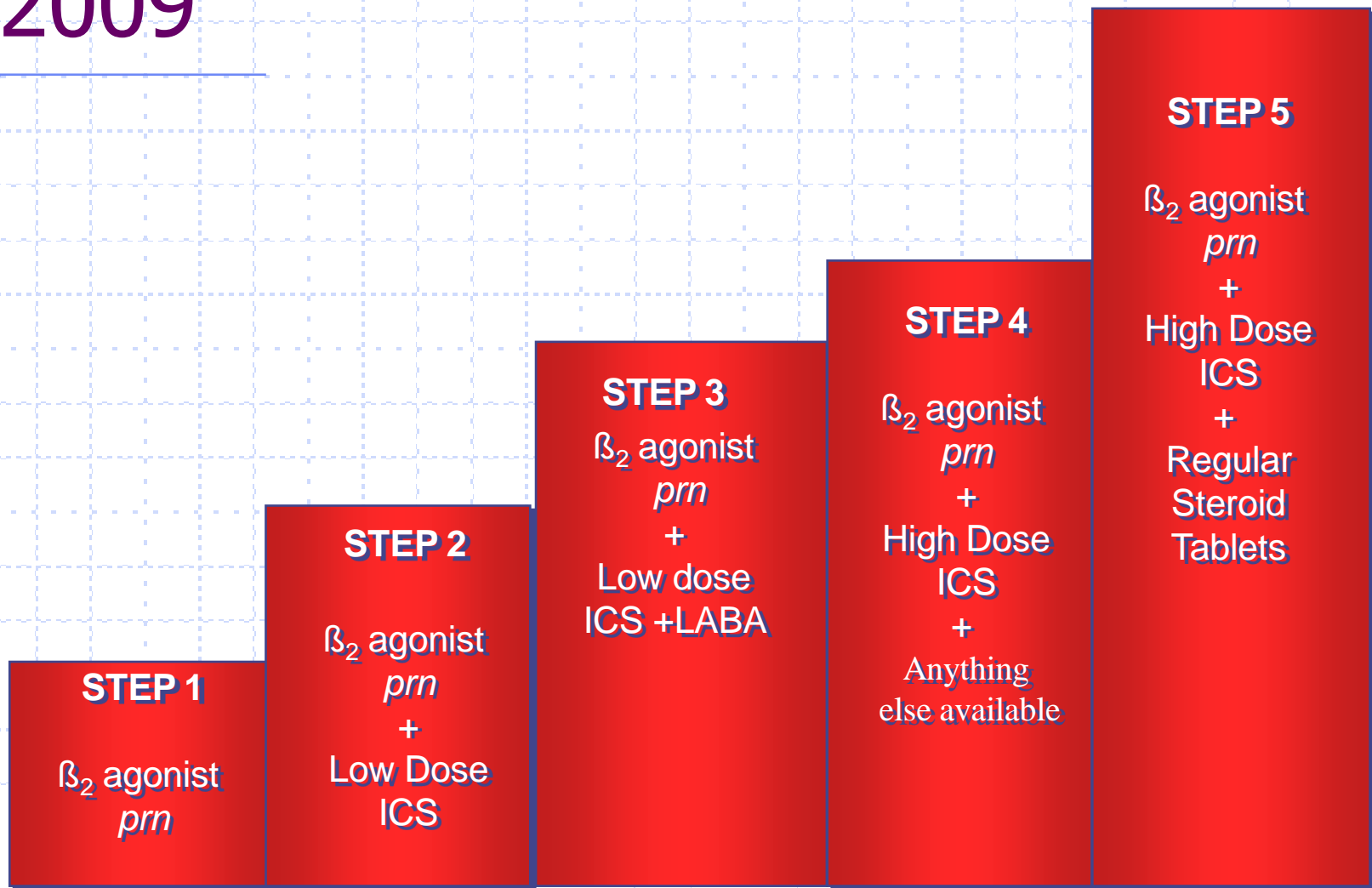
Guidelines on Asthma Management 1990s



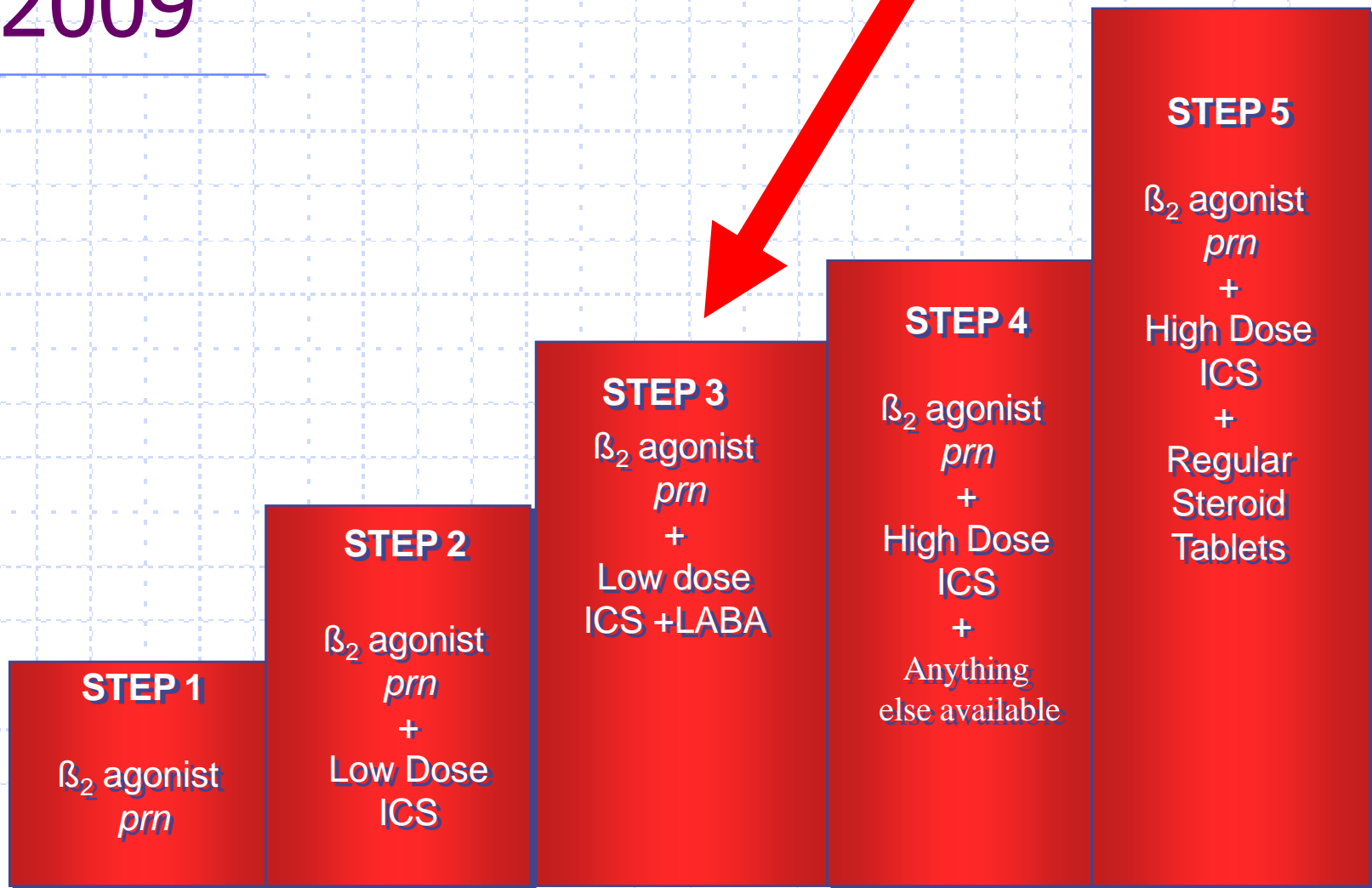
New understanding of asthma treatments 1994-2004



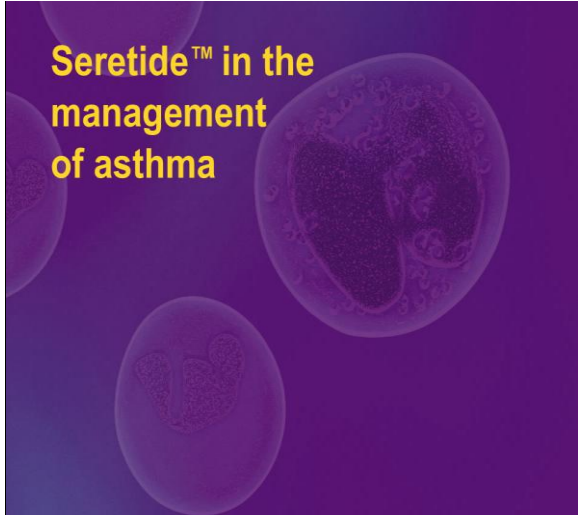
Guidelines on Asthma Management 2009



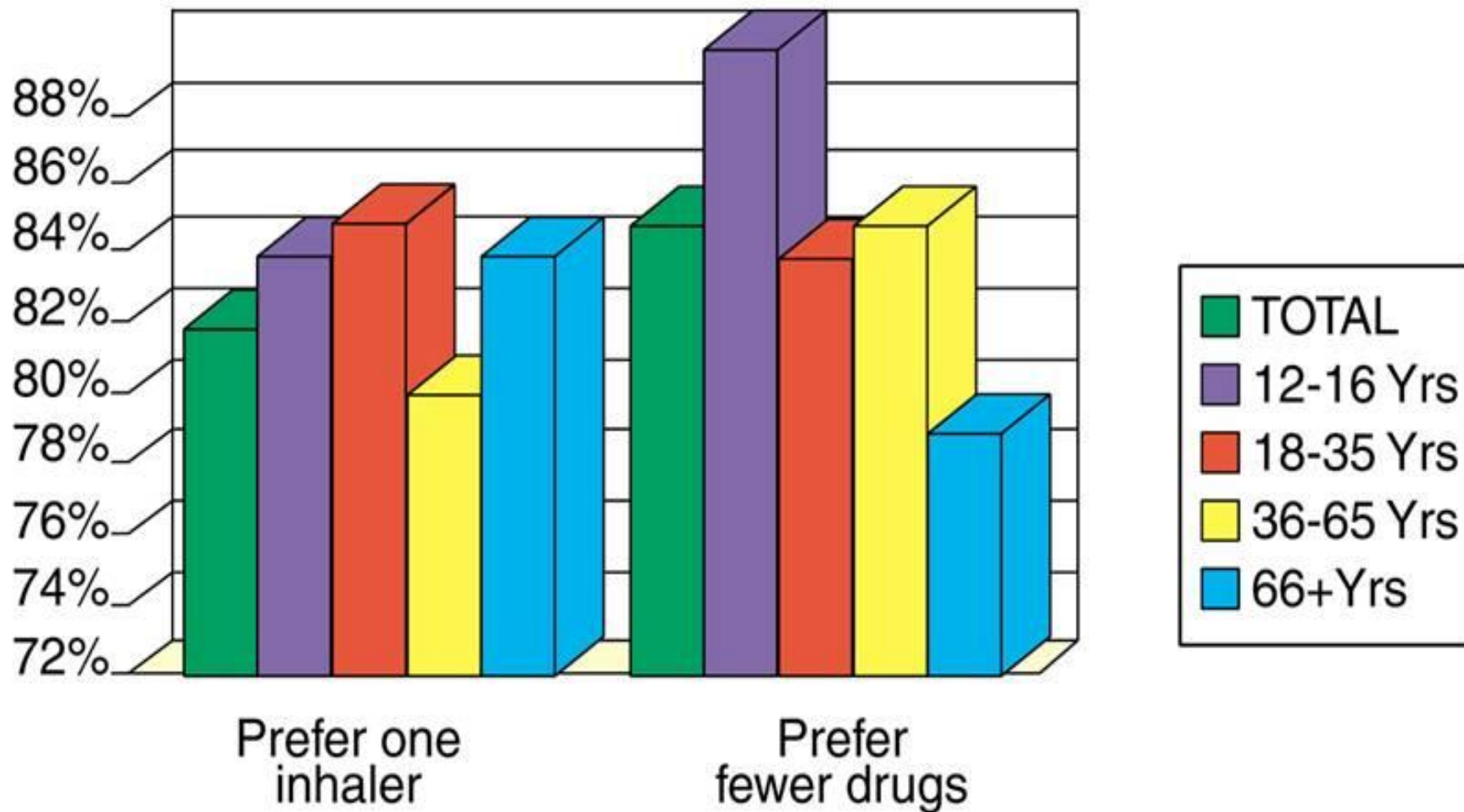
Guidelines on Asthma Management 2009



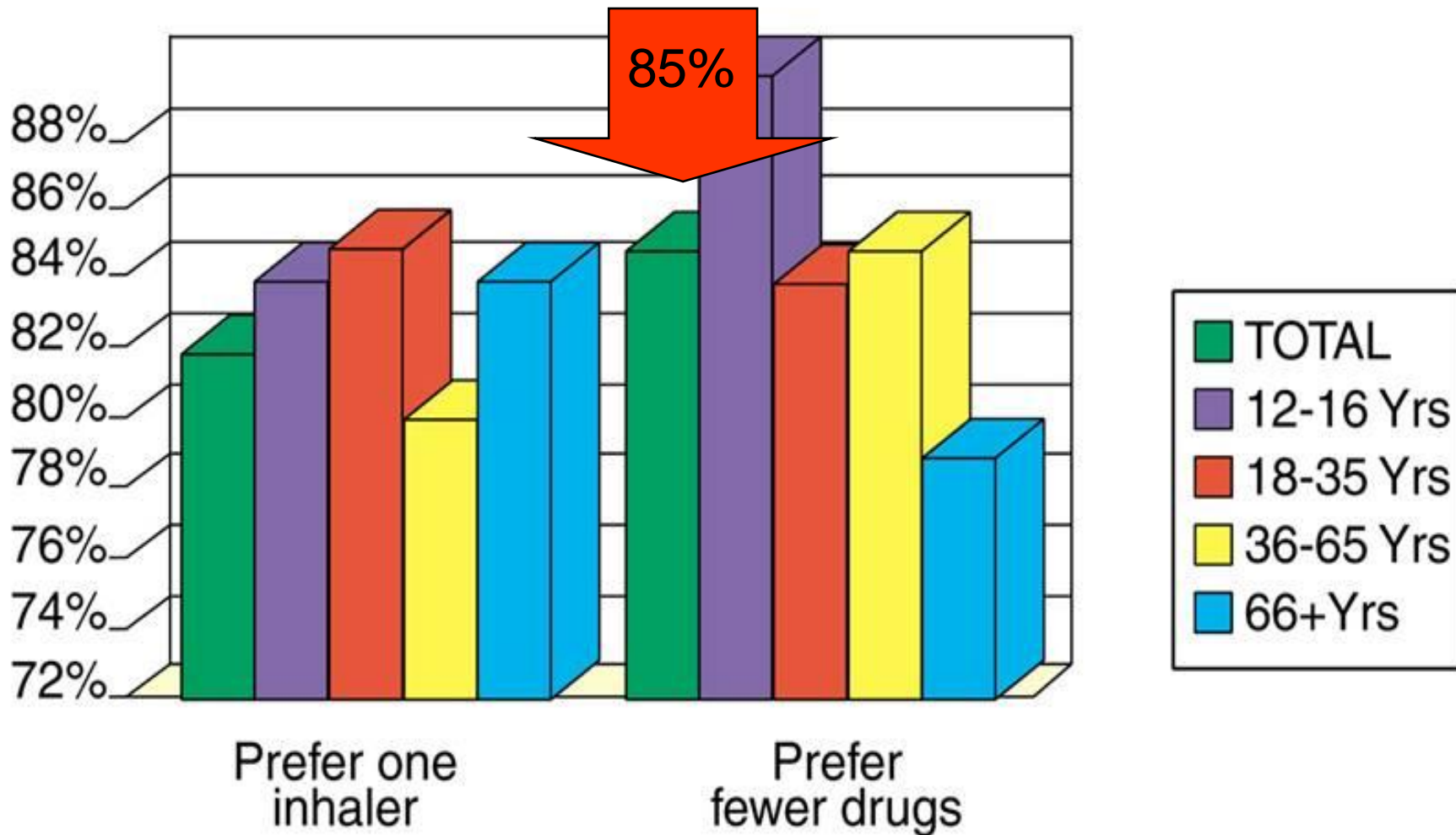
Making things simpler for patients



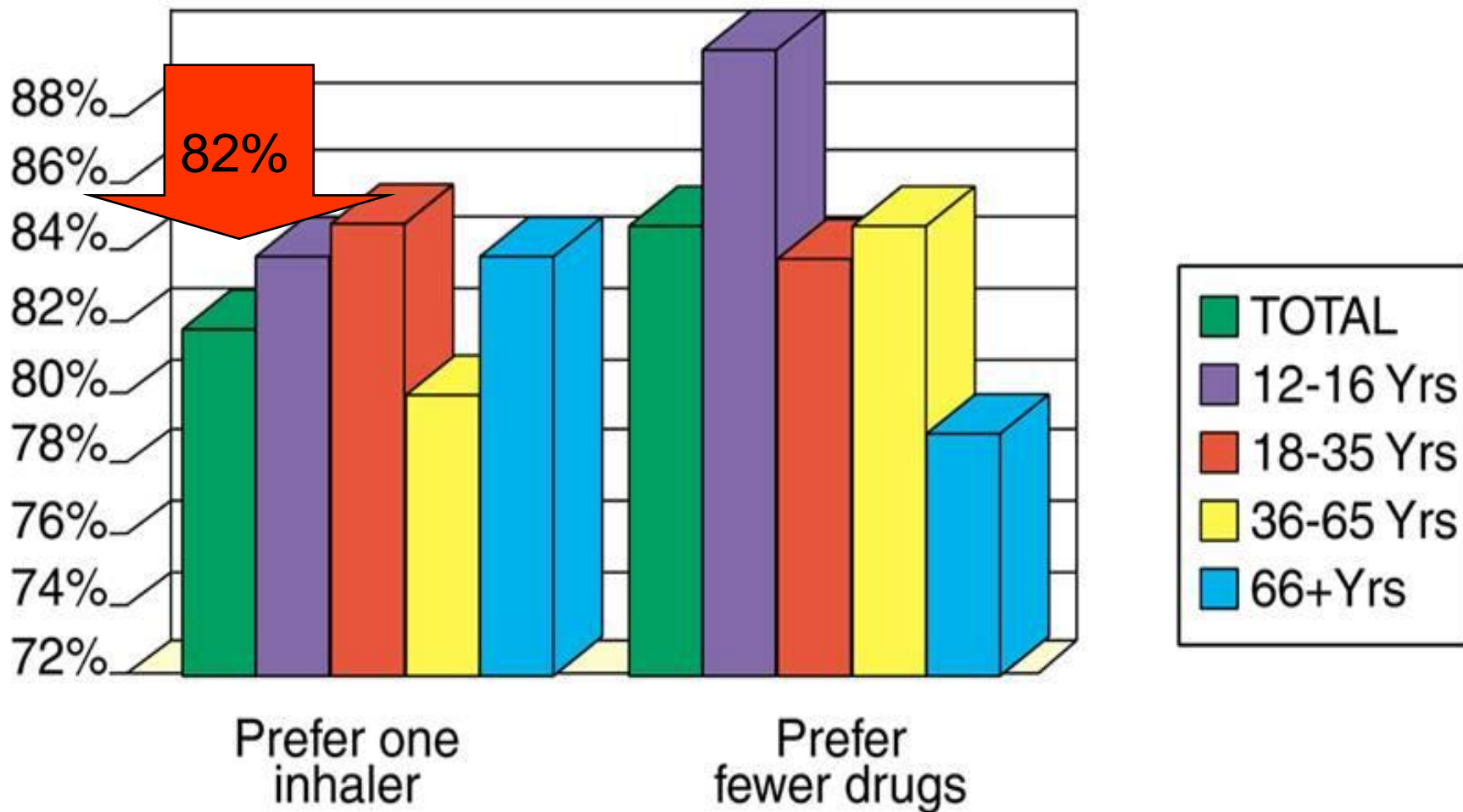
Patient wishes to use just one inhaler and fewer drugs (n=454). Stahl E et al ATS 2002



Patient wishes to use just one inhaler and fewer drugs (n=454). Stahl E et al ATS 2002



Patient wishes to use just one inhaler and fewer drugs (n=454). Stahl E et al ATS 2002



Why might patients prefer combination inhalers?



- Less to carry about
- Less need to understand the type and purpose of each
- Less worry about one prescription finishing before the other
- Less expense
- Less to remind them they have a “disease”

Why might patients prefer combination inhalers?



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**And (from our point of view)
AVOIDS THE RISK OF MONOTHERAPY**

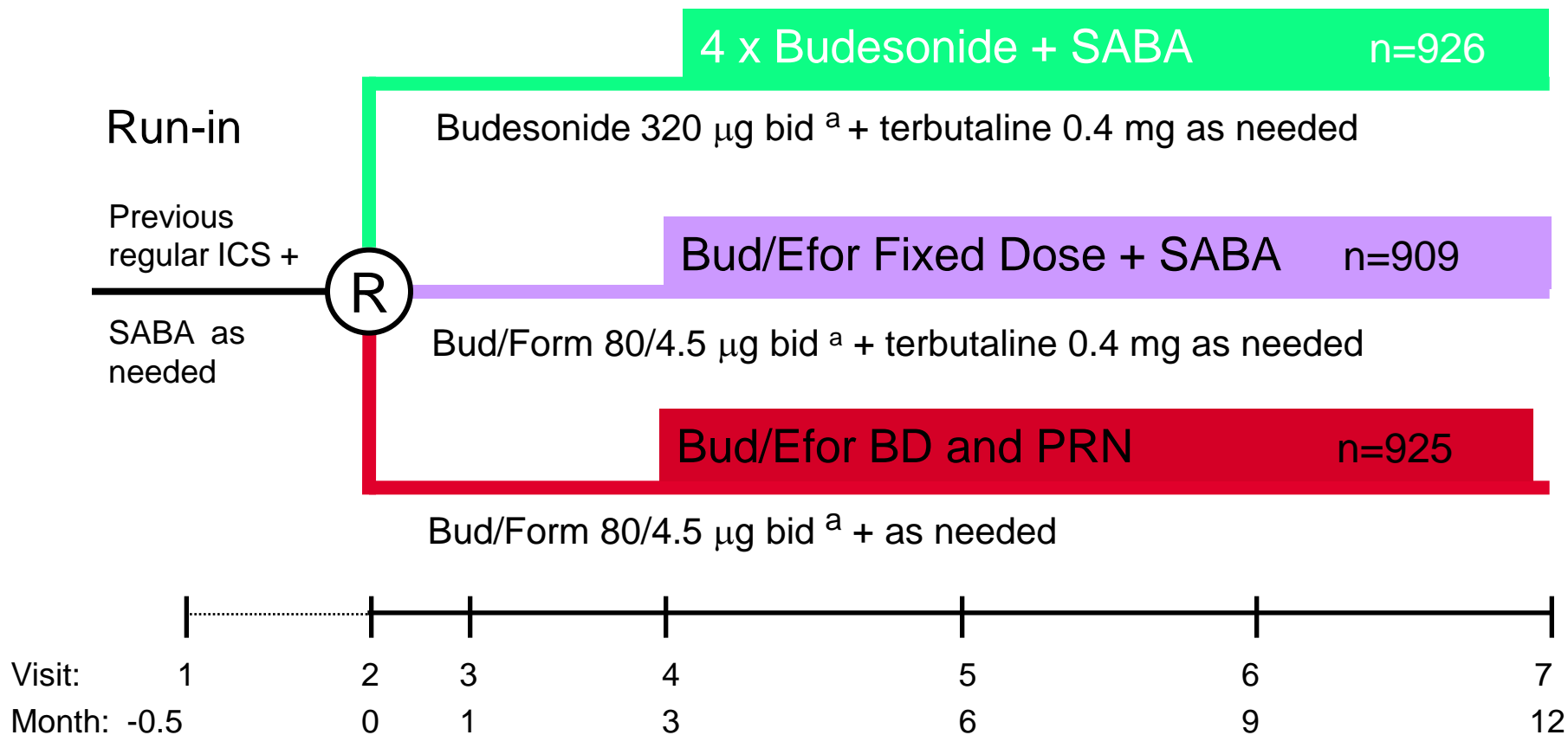
Henry Hyde Salter (1823-1871)



Differentiated asthma from other causes of breathlessness as “paroxysmal dyspnoea of a peculiar character with intervals of healthy respiration between attacks”

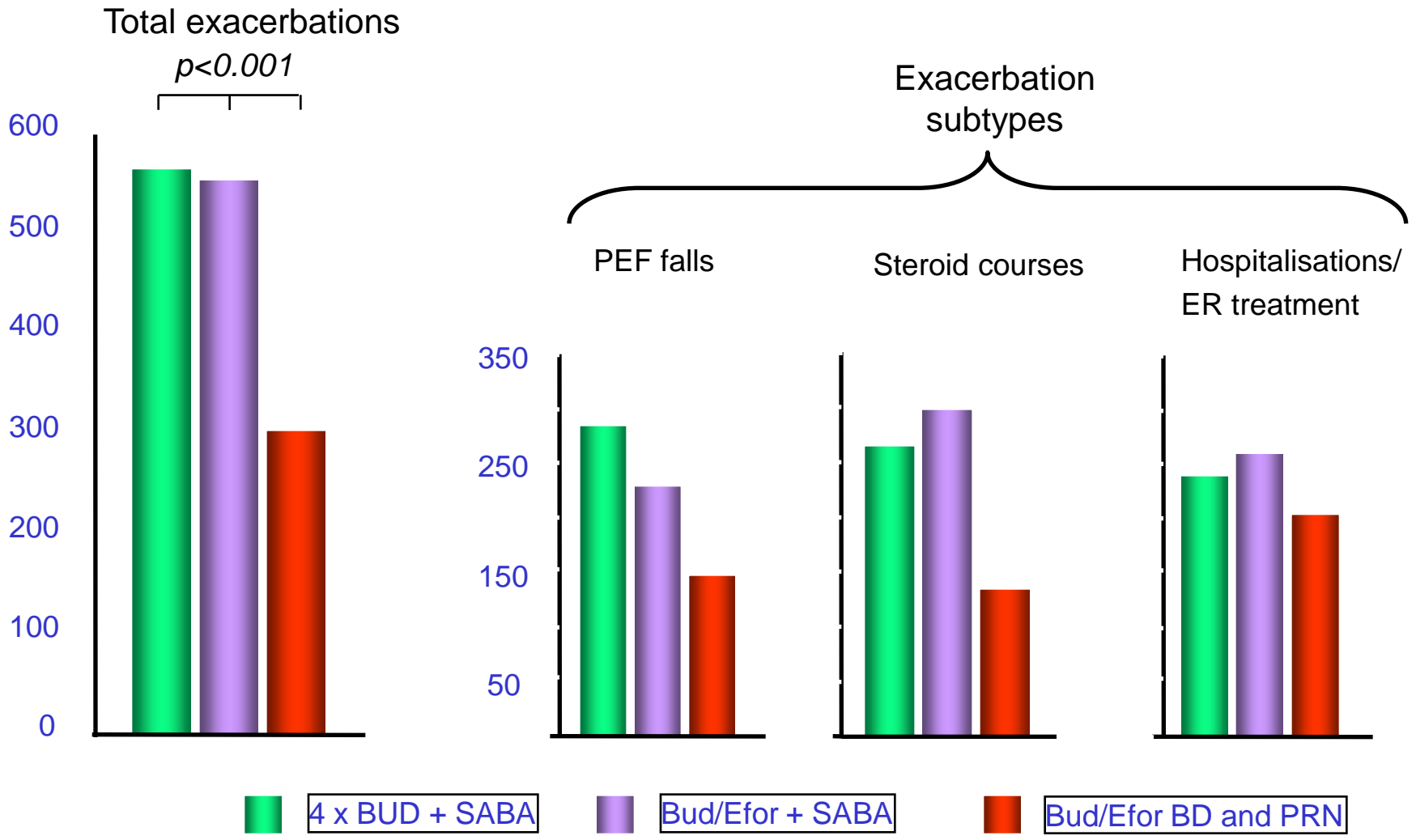
How do combination Steroid /Formoterol Inhalers used regularly and prn compare with High dose steroids alone?

O'Byrne PM et al. Am J Respir Crit Care Med 2005; 171:129-136

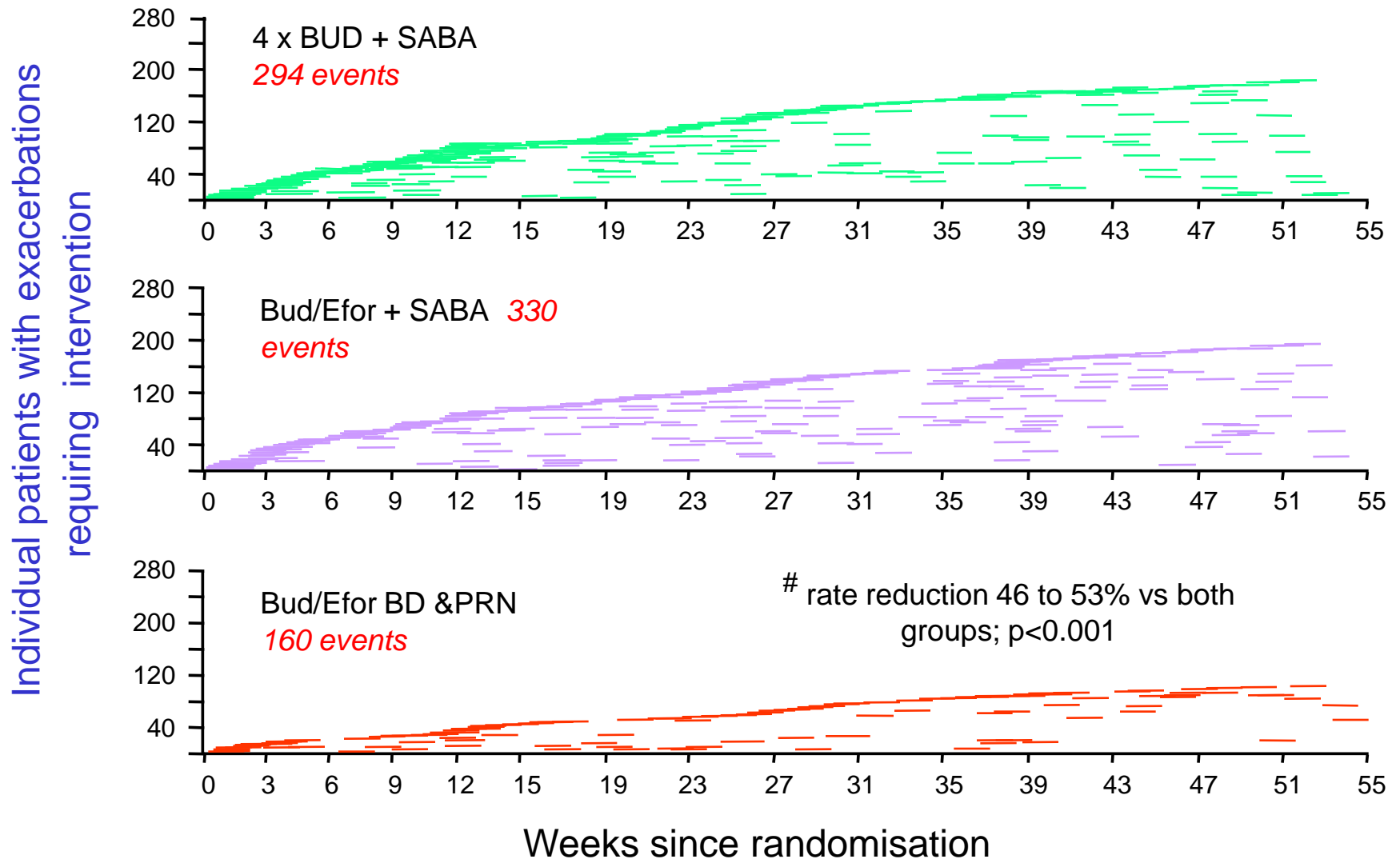


^a Children <12 years received half the daily maintenance dose with a once daily regimen

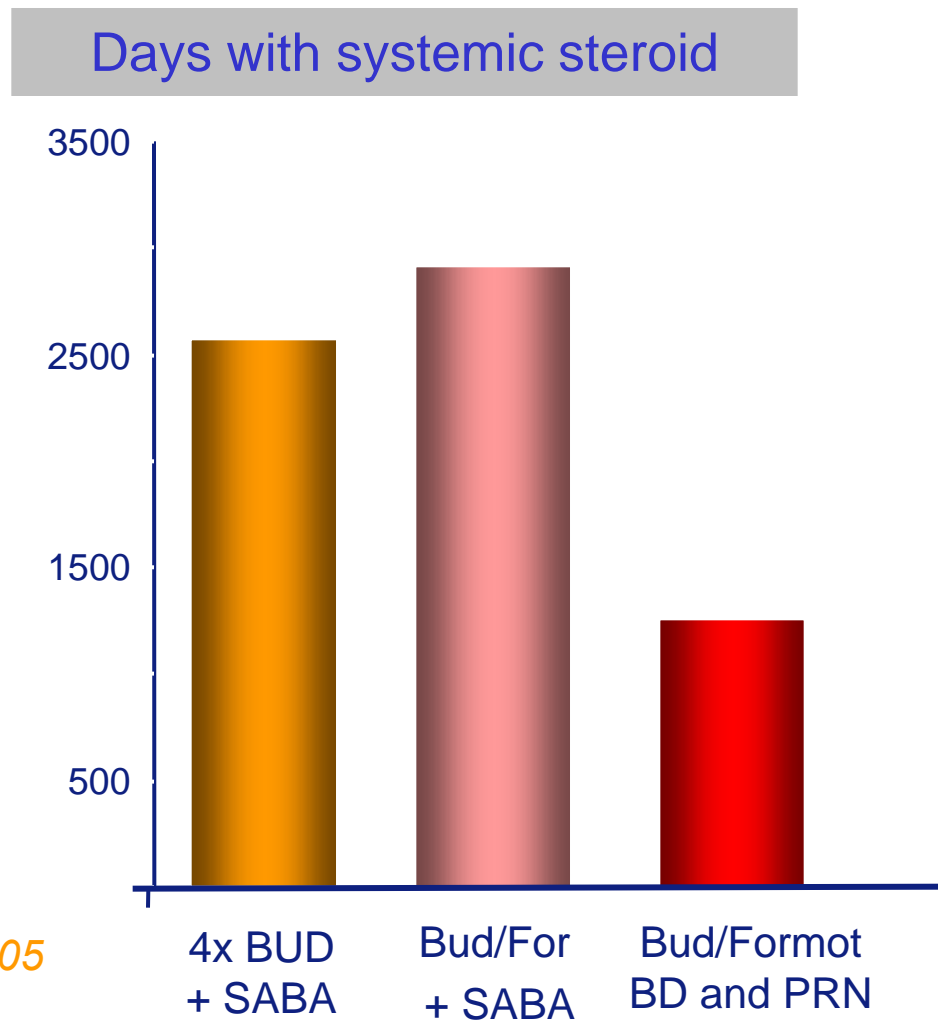
Severe Exacerbations



Total Asthma Exacerbations Requiring Medical Intervention

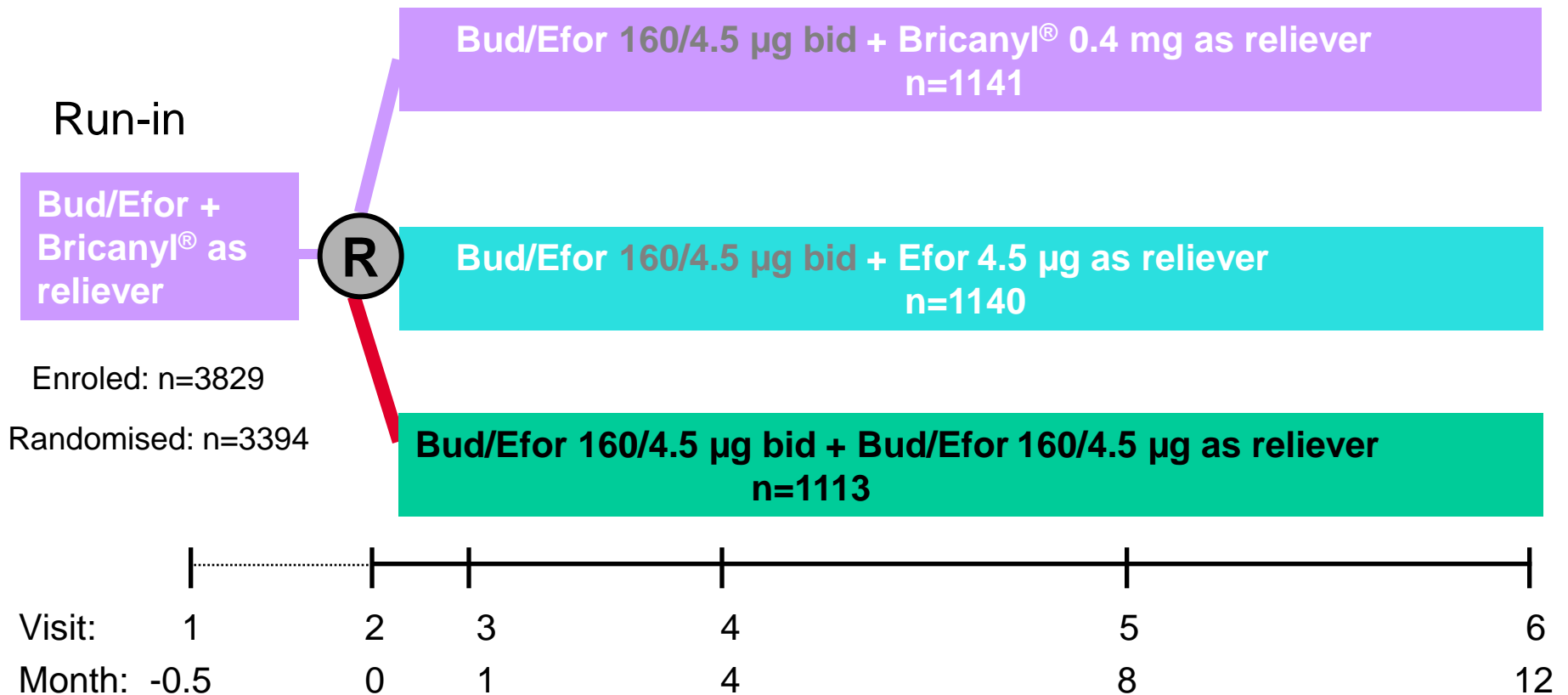


Steroid load during 1 year of treatment



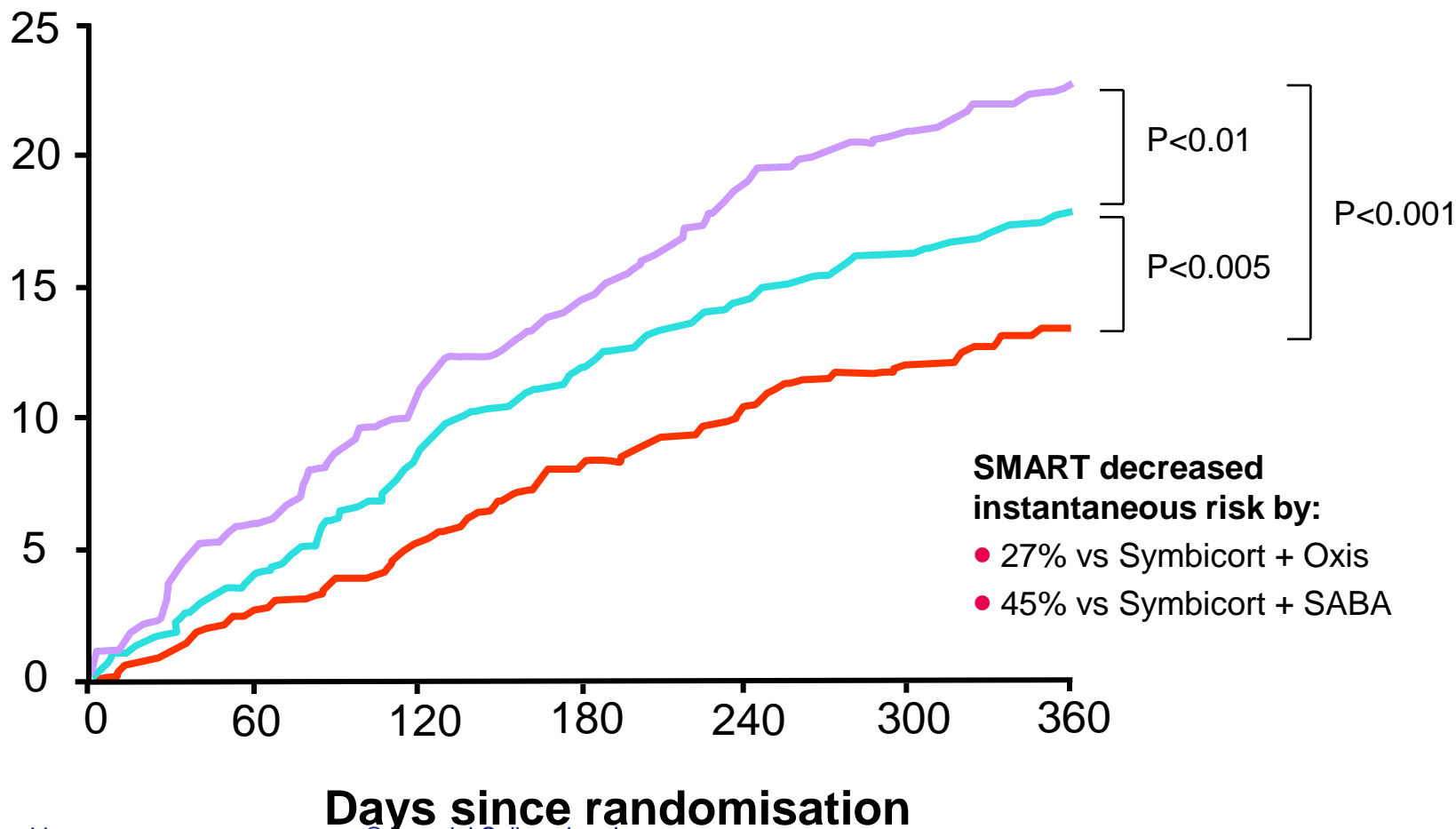
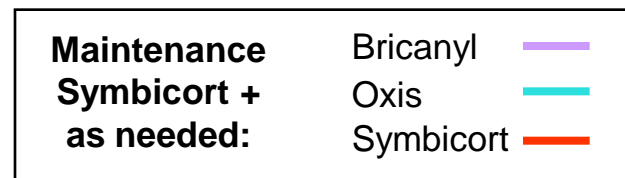
O'Byrne et al., AJRCCM 2005

Is it just a formoterol effect or is the inhaled steroid needed as well?

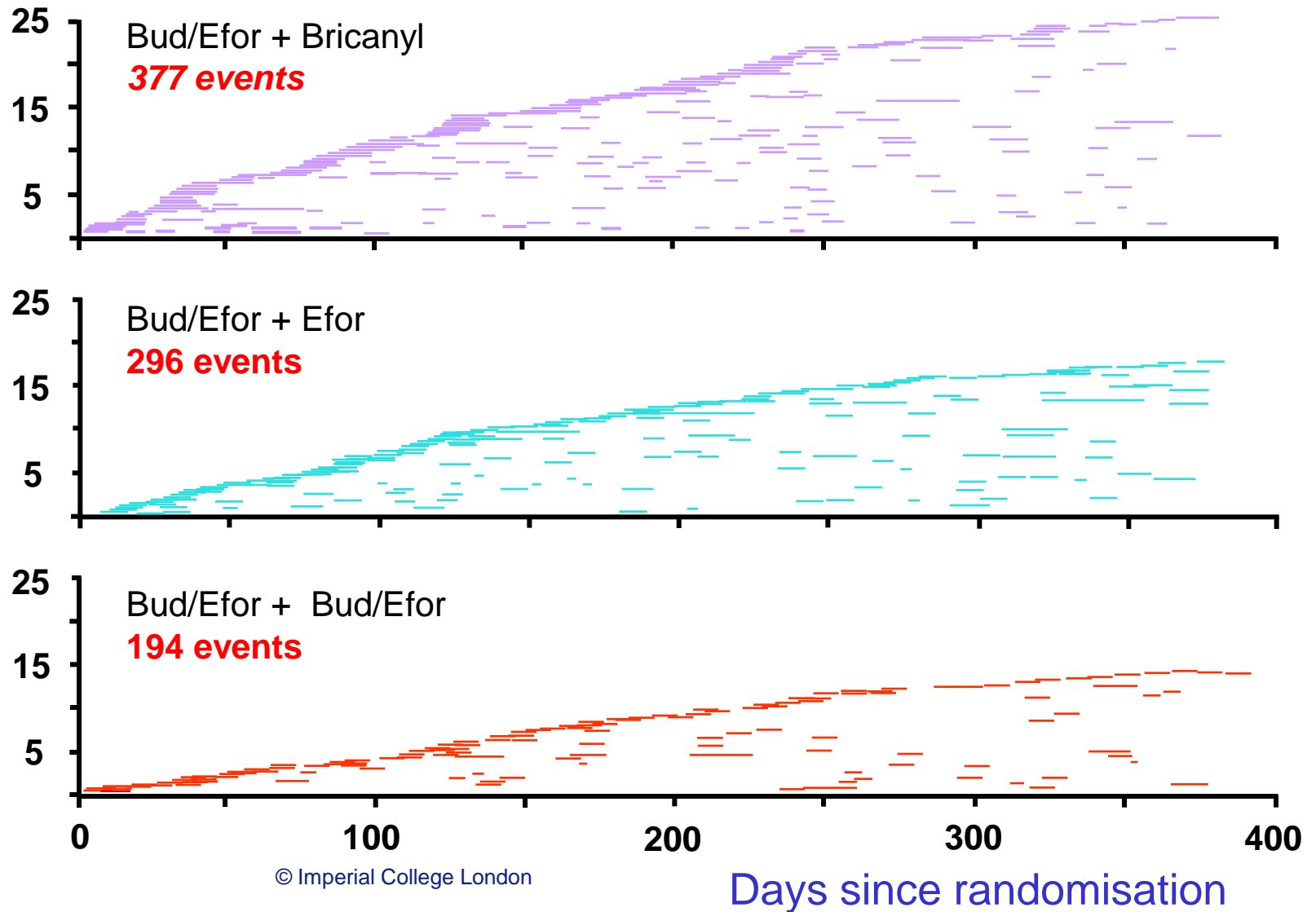


Time to first severe exacerbation

Patients with severe exacerbations (%)



Total severe exacerbations



ICS / Formoterol Combination inhaler as Maintenance and Relief

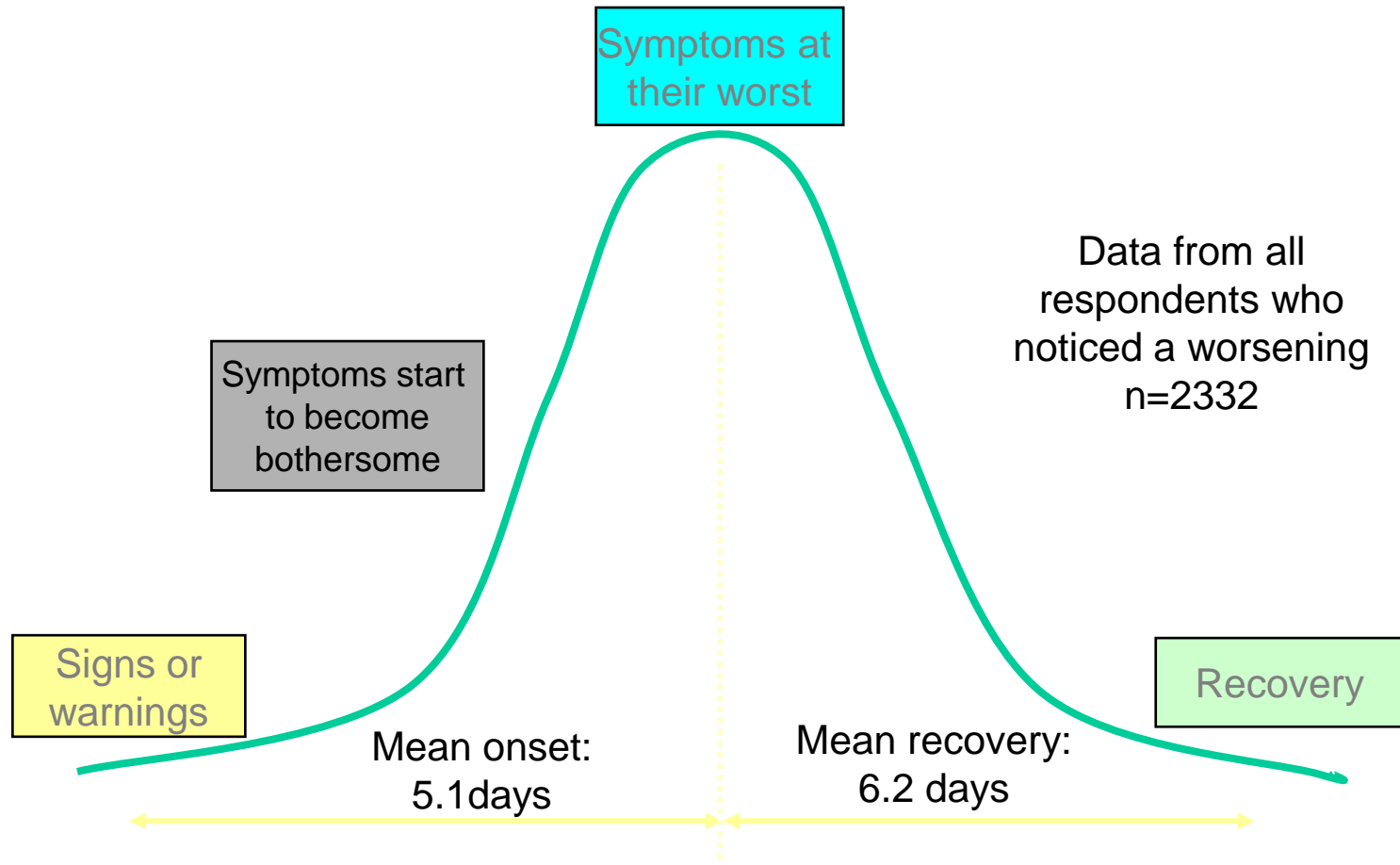
Results in:

- Fewer total and repeat severe exacerbations
- Fewer hospitalisations/ER visits
- Lower systemic corticosteroid use
- Fewer patients with high use of as needed treatment
- Asthma control as good as high dose maintenance

Both an inhaled steroid and eformoterol are needed for the benefit.

How might combination ICS/Formoterol inhalers work “as required” (as well as regularly)?

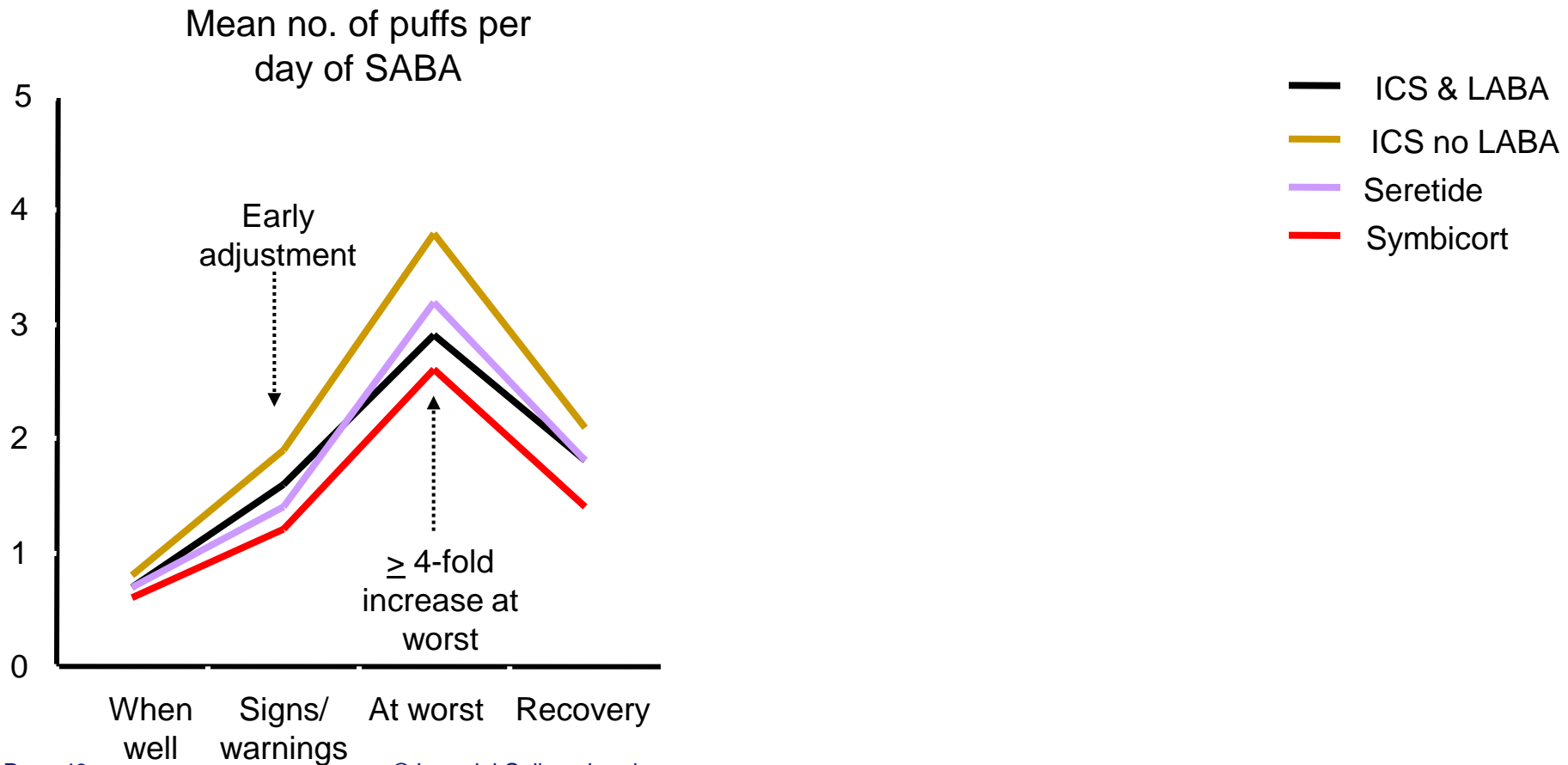
Time from early warning signs to worst symptoms gives a 6-day window of opportunity



Partridge MR, BMC Pulmonary Medicine 2006

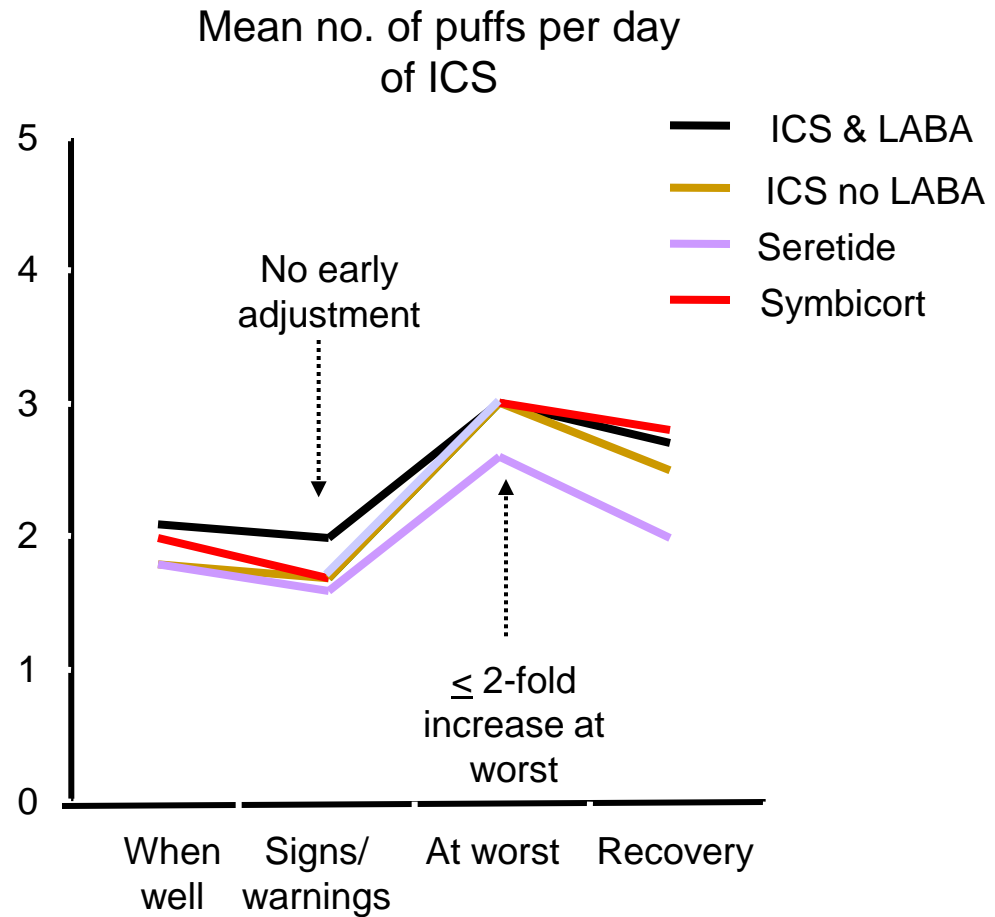
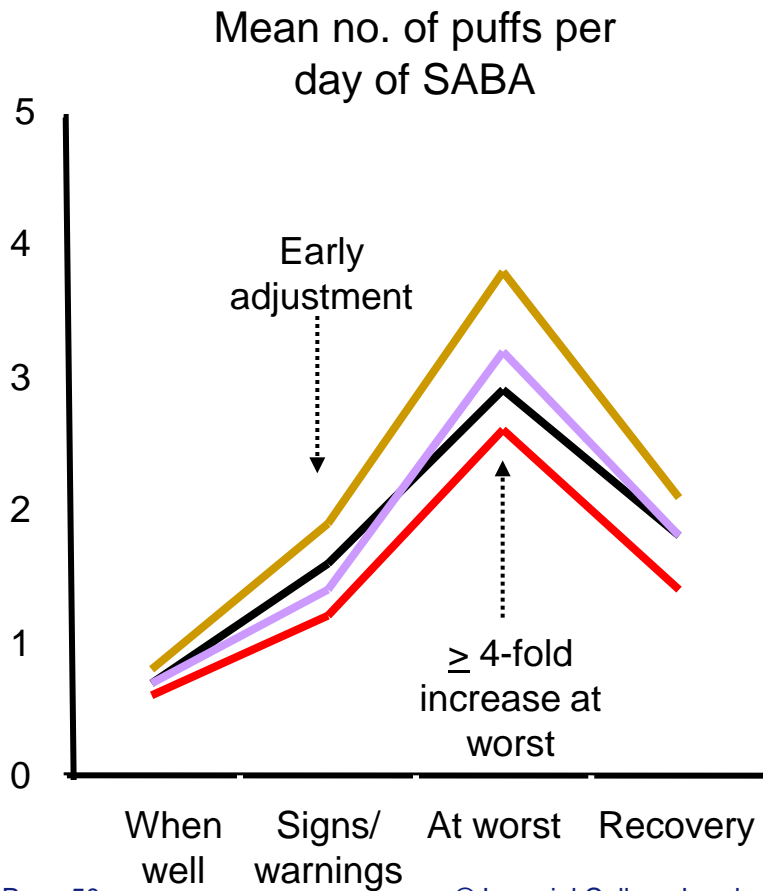
Patients adjust their SABA early and ICS too late at the time of a worsening of symptoms

Partridge MR, BMC Pulmonary Medicine 2006

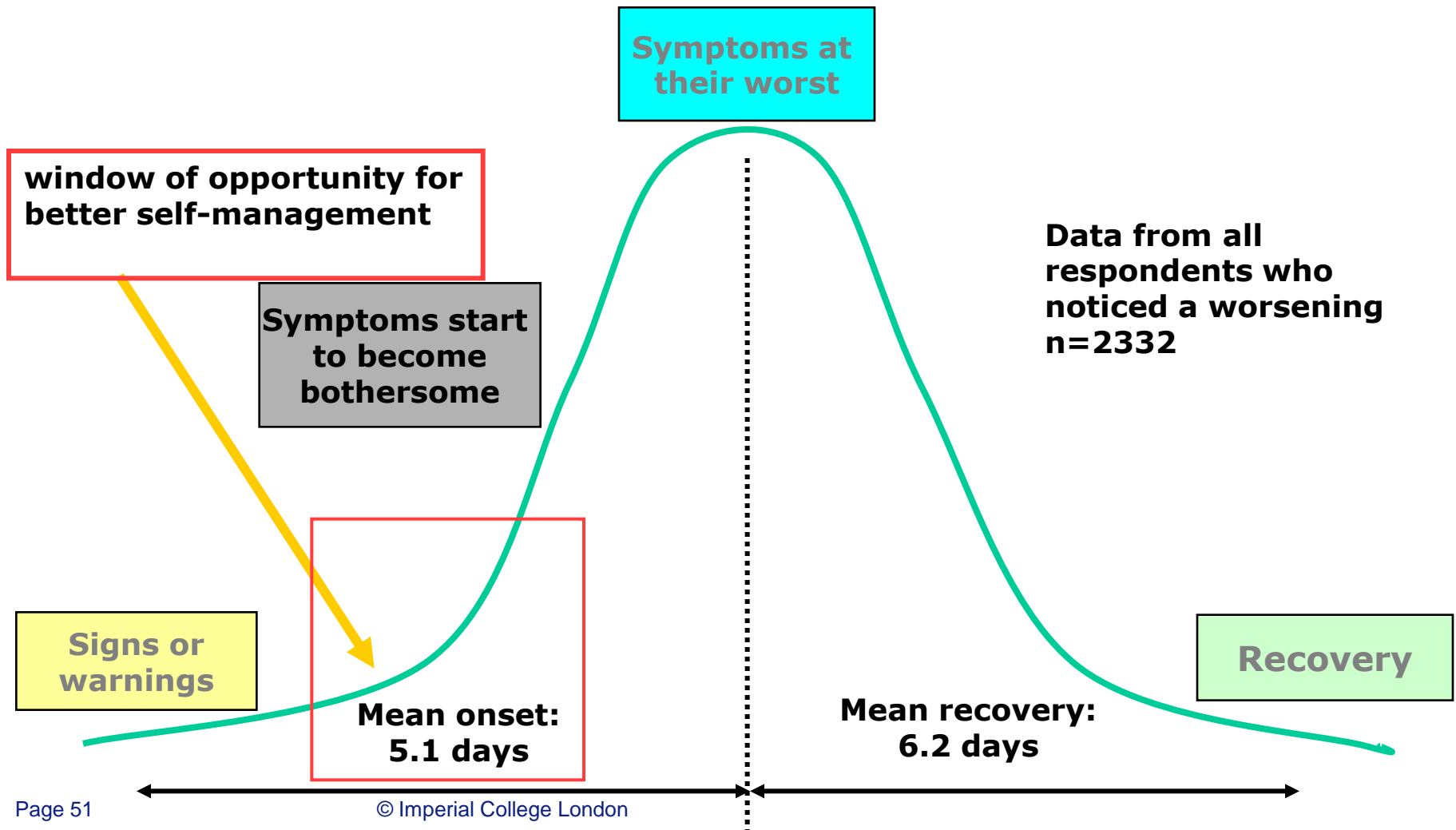


Patients adjust their SABA early and ICS too late at the time of a worsening of symptoms

Partridge MR, BMC Pulmonary Medicine 2006



Time from early signs of worsening up to the worst symptoms around 5-days



Important to remember that varying doses
applies only to formoterol and not to other
LABAs

Formoterol has unique pharmacological properties compared with other long-acting β_2 -agonists

Formoterol

Long duration (>12 hours)

Rapid onset of action

Full receptor agonist

Dose–response

Salmeterol

Long duration (>12 hours)

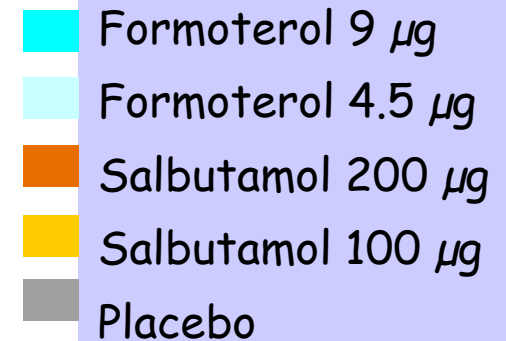
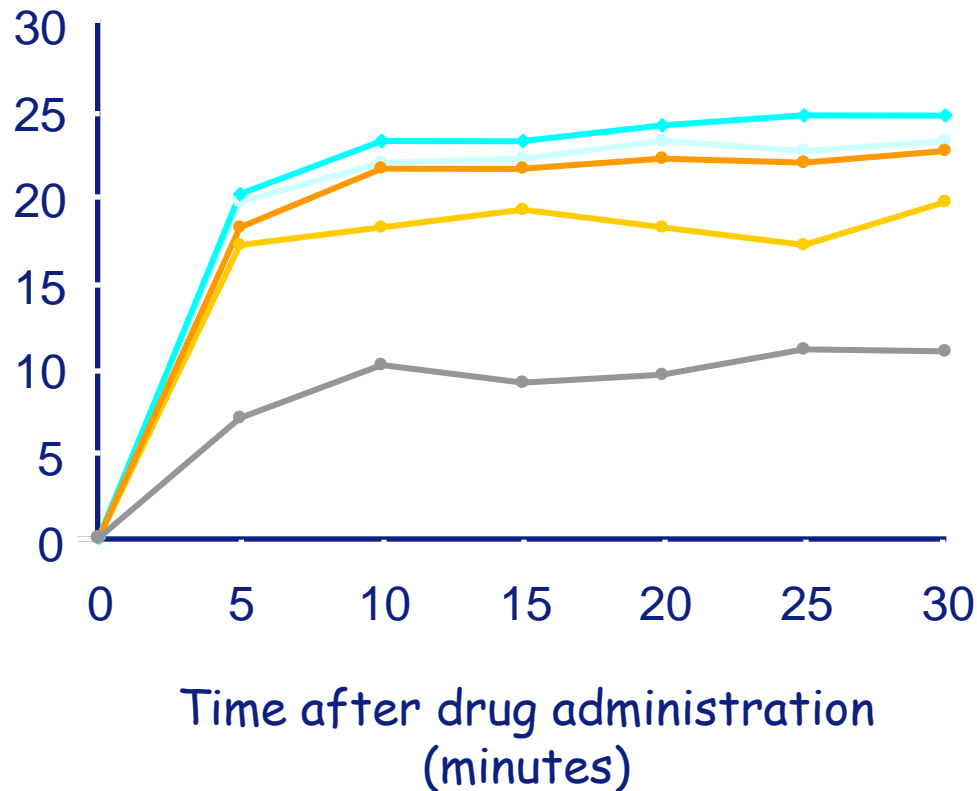
Delayed onset of action

Partial receptor agonist

No dose–response

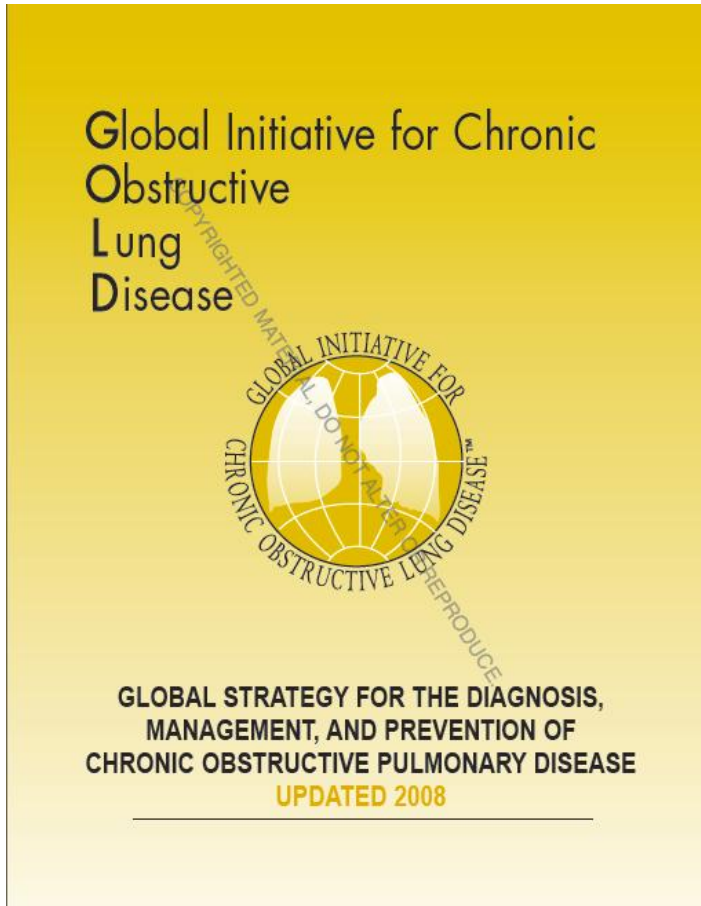
Formoterol Has a Rapid Onset of Action Similar to Salbutamol

Mean FEV₁
(% change from baseline)



FEV₁ at 3 minutes
after inhalation:
p<0.001 for all active
treatments compared
with placebo

Guideline goals for successful COPD management



- **Relieve symptoms**
- **Improve exercise tolerance**
- **Improve health status**
- **Prevent disease progression**
- **Prevent and treat complications**
- **Prevent and treat exacerbations**
- **Reduce mortality**

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- Improve exercise tolerance
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Therapy at Each Stage of COPD

I: Mild

II: Moderate

III: Severe

IV: Very Severe

- $FEV_1/FVC < 70\%$
- $FEV_1 \geq 80\%$ predicted

- $FEV_1/FVC < 70\%$
- $50\% \leq FEV_1 < 80\%$ predicted

- $FEV_1/FVC < 70\%$
- $30\% \leq FEV_1 < 50\%$ predicted

- $FEV_1/FVC < 70\%$
- $FEV_1 < 30\%$ predicted or $FEV_1 < 50\%$ predicted plus chronic respiratory failure

Active reduction of risk factor(s); influenza vaccination
Add short-acting bronchodilator (when needed)

Add regular treatment with one or more long-acting bronchodilators (when needed); **Add** rehabilitation

Add inhaled glucocorticosteroids if repeated exacerbations

Add long term oxygen if chronic respiratory failure.
Consider surgical treatments

The use of short acting inhaled bronchodilators



- GOLD recommends for moderate to very severe COPD, use of regular long acting inhaled bronchodilators (formoterol, salmeterol and tiotropium), rather than short acting bronchodilators.
- Evidence Level A

Formoterol significantly reduces breathlessness in COPD

Breathlessness symptom score

2.4

2.3

2.2

2.1

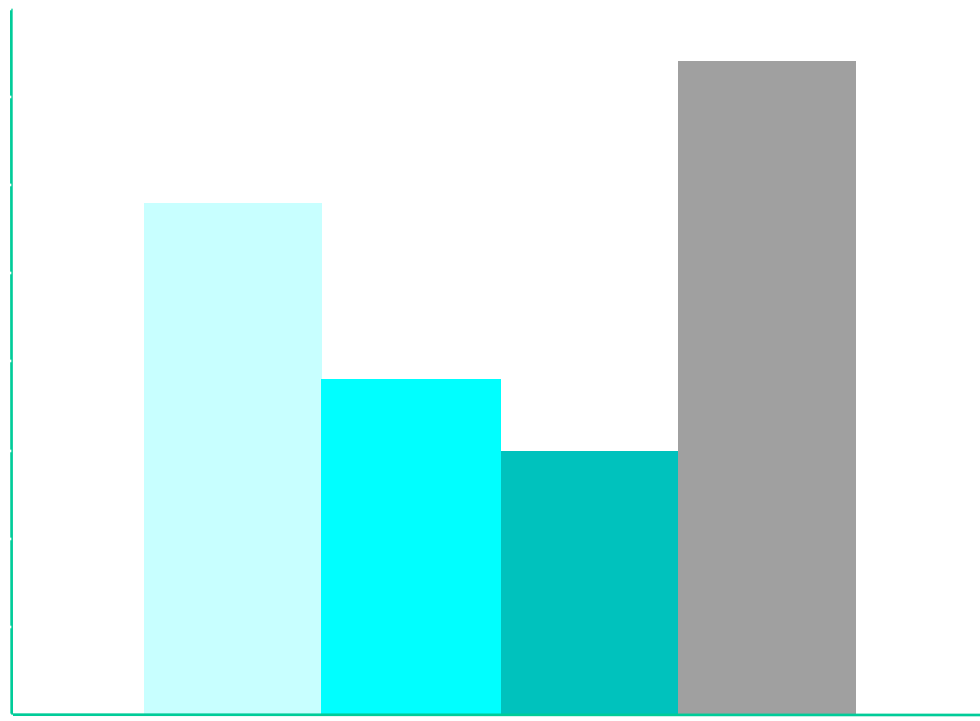
2.0

- Formoterol 4.5 µg bid
- Formoterol 9 µg bid
- Formoterol 18 µg bid
- Placebo

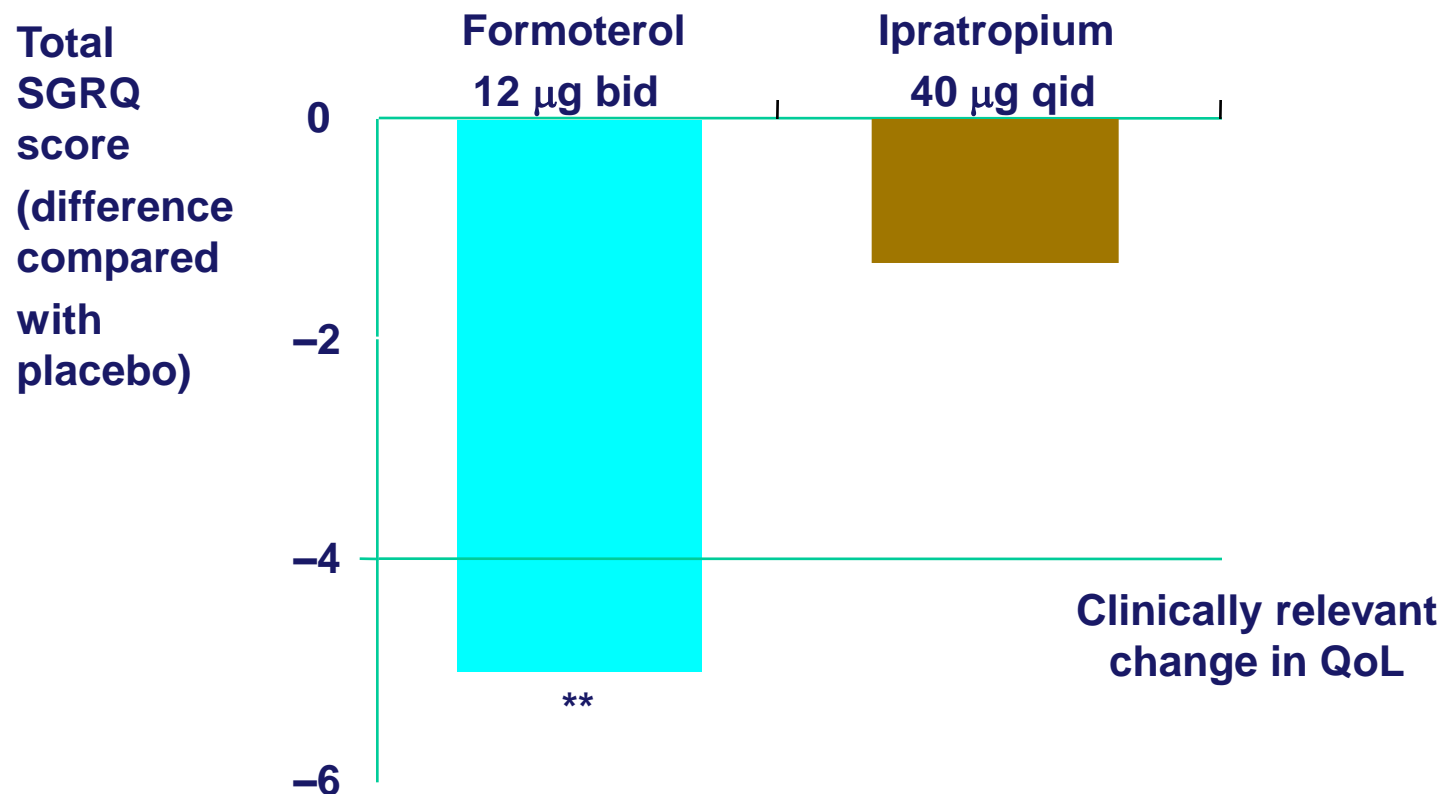
Week 12

**p<0.01, *p<0.05; vs placebo

Adapted from Aalbers et al 2002



Formoterol improves quality of life significantly compared with ipratropium



**p<0.005 vs ipratropium, p<0.001 vs placebo

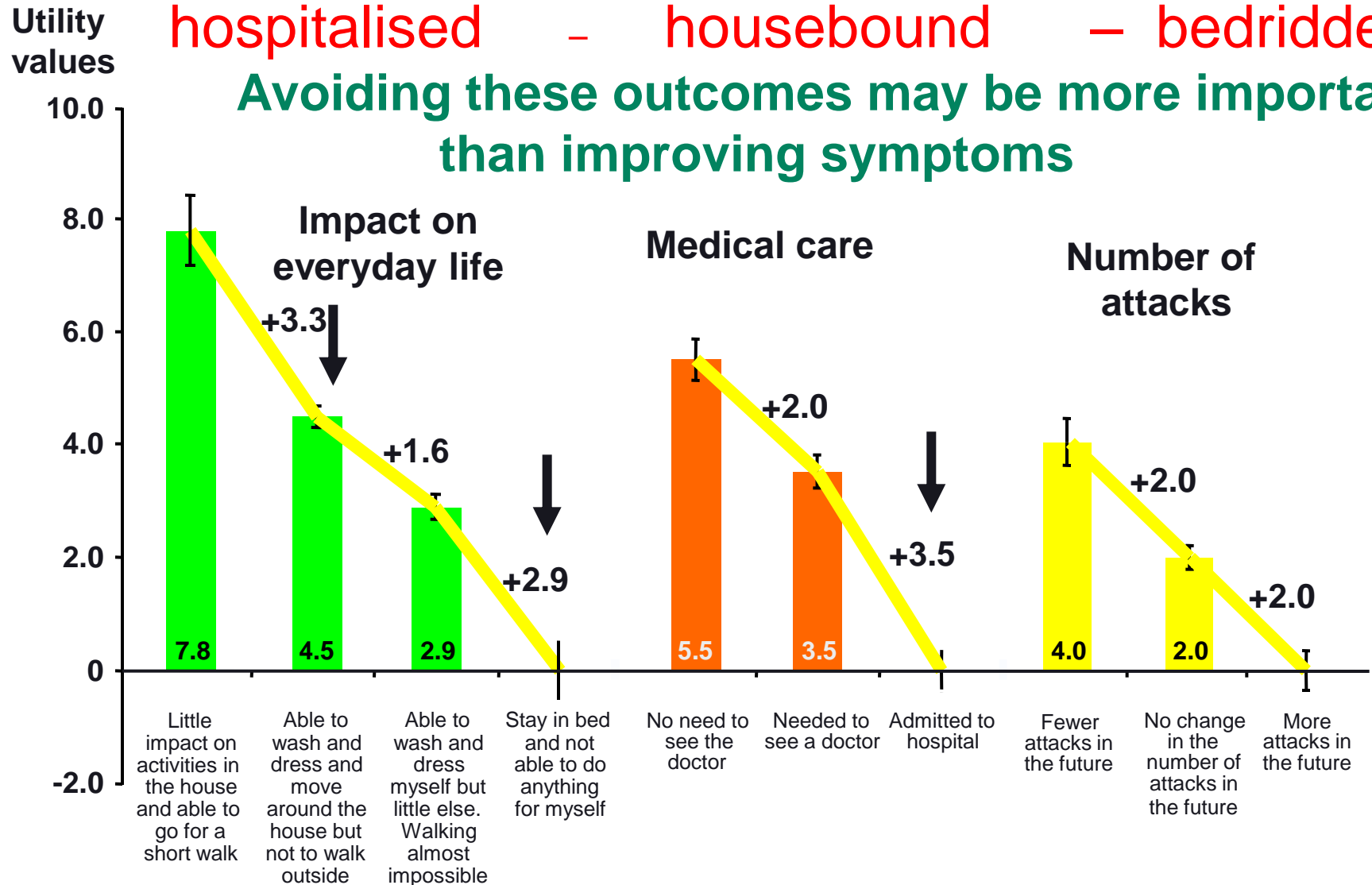
Adapted from Dahl et al 2000

What aspect of their condition do those with COPD most fear?

Discrete choice modelling: Utility shifts

COPD patients are most concerned about being hospitalised – housebound – bedridden

Avoiding these outcomes may be more important than improving symptoms



COPD exacerbations worsen long-term prognosis

- Lung function declines faster in patients with frequent versus infrequent exacerbations

Measure	Infrequent (<2.92/year) ^a	Frequent (>2.92/year) ^a	p-value
No. of patients	63	46	
PEF decline, L/min/year	-0.72	-2.94	<0.001
FEV ₁ decline, mL/year	-32.1	-40.1	<0.05

FEV₁ = forced expiratory volume in 1 second

Donaldson GC, et al. Thorax 2002;57:847–852

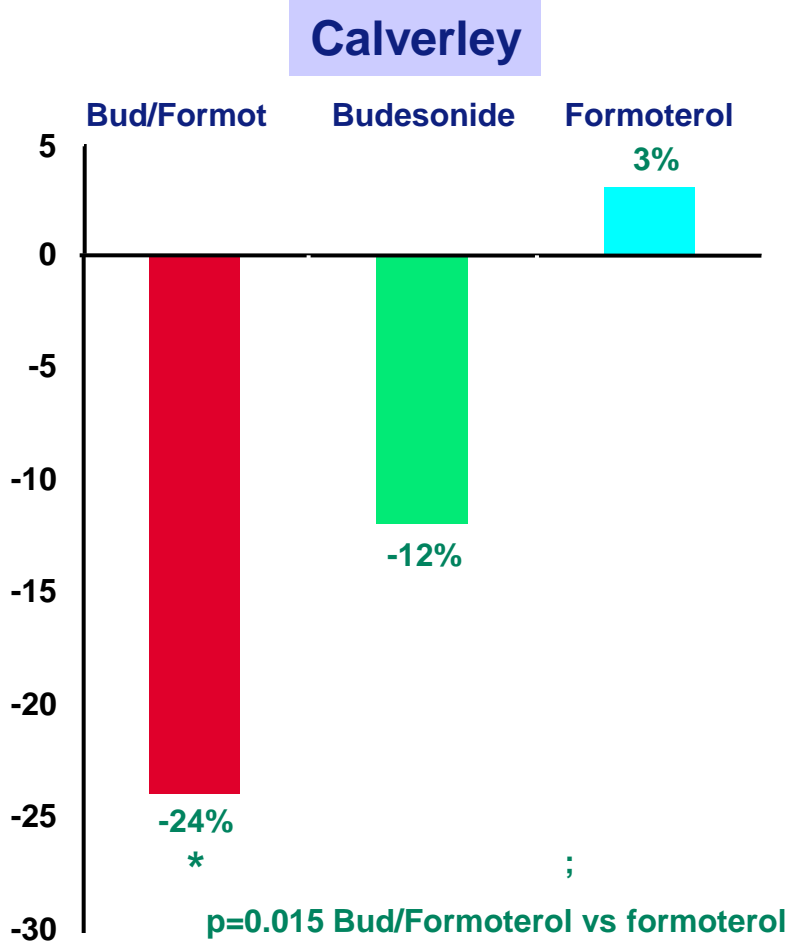
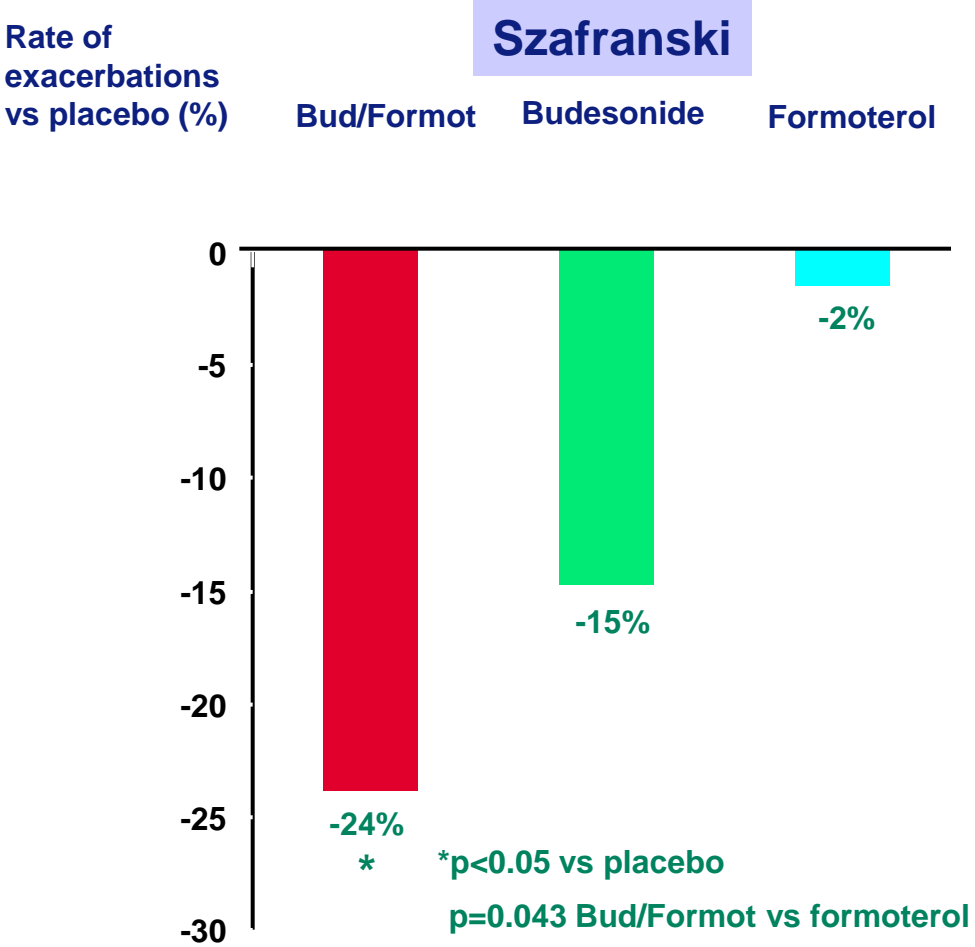
Cost of exacerbations

- 54% of the economic cost of COPD accrues from hospitalisation

Britton M Respir Med 2003; 97: s71-S79

So from a patient viewpoint, from a prognosis viewpoint and from a Government viewpoint, exacerbations are an important issue to address

ICS/Formoterol combinations reduce the rate of exacerbations requiring medical intervention



Szafranski W, et al. Eur Respir J 2003;21:74–81
 Page Calverley PM, et al. Eur Respir J 2003; 22: 912-919

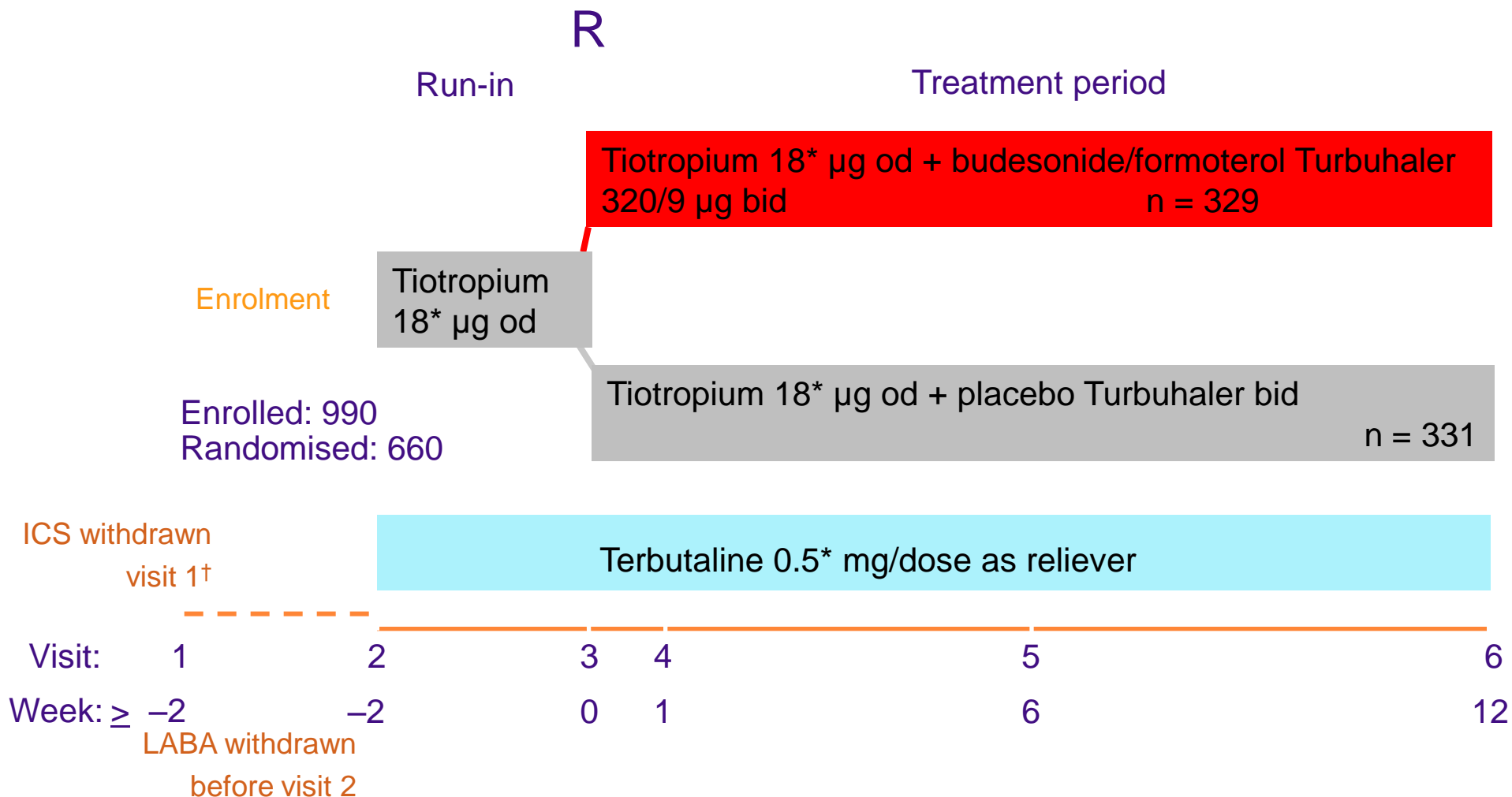
Efficacy and Tolerability of Budesonide/Formoterol Added to Tiotropium in Patients with Chronic Obstructive Pulmonary Disease

Tobias Welte¹, Marc Miravittles², Paul Hernandez³, Göran Eriksson^{4,5}, Stefan Peterson⁵, Tomasz Polanowski⁵, and Romain Kessler⁶

¹Department of Respiratory Medicine, Hannover Medical School, Germany; ²Fundació Clínic, Institut d'Investigacions Biomèdiques August Pi i Sunyer, Barcelona, Spain; ³Respirology Division, Department of Medicine, Dalhousie University, Halifax, Nova Scotia, Canada; ⁴Department of Respiratory Medicine and Allergology, University Hospital, Lund, Sweden; ⁵AstraZeneca Research and Development, Lund, Sweden; and ⁶Department of Pneumology, Hôpitaux Universitaires de Strasbourg, Strasbourg, France

CLIMB: study design

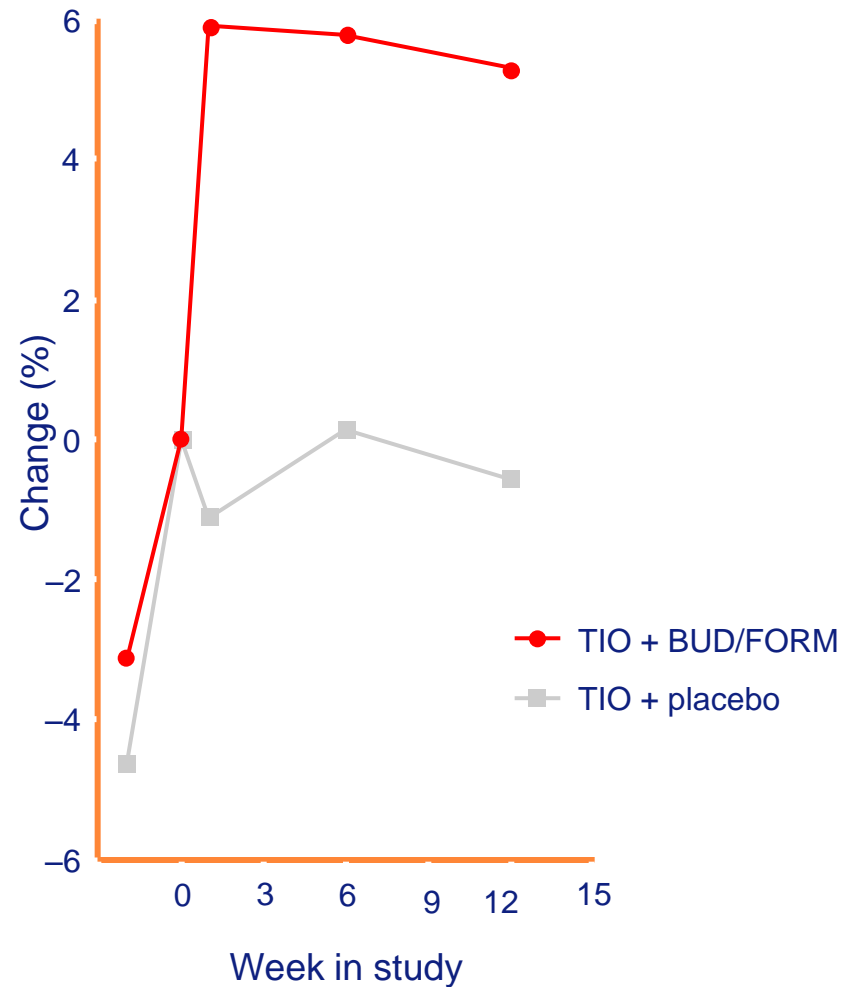
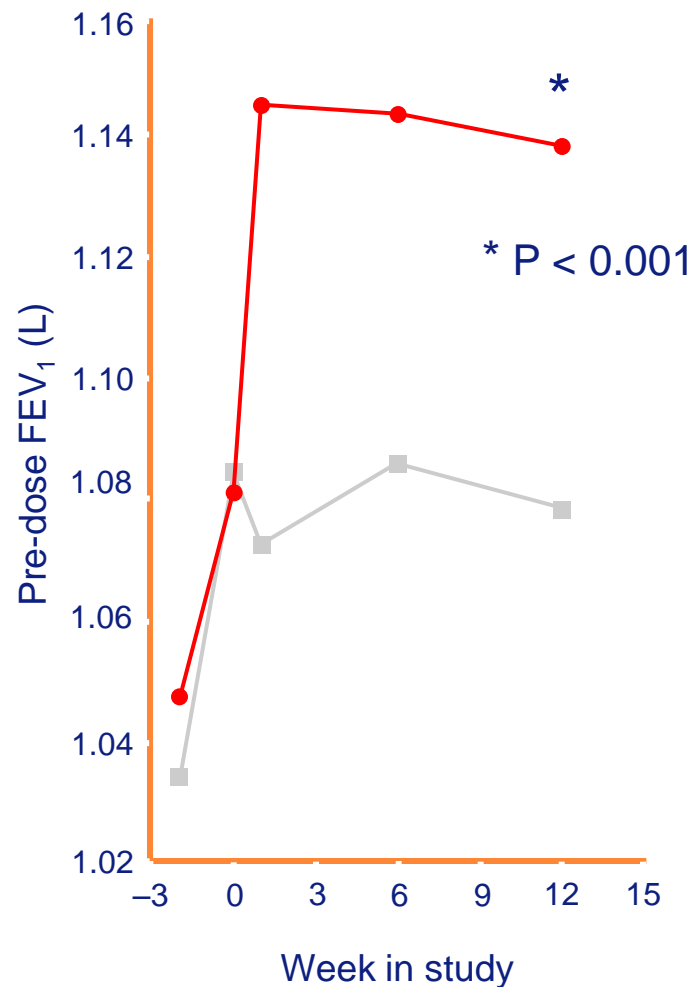
3-month, double-blind, randomised study



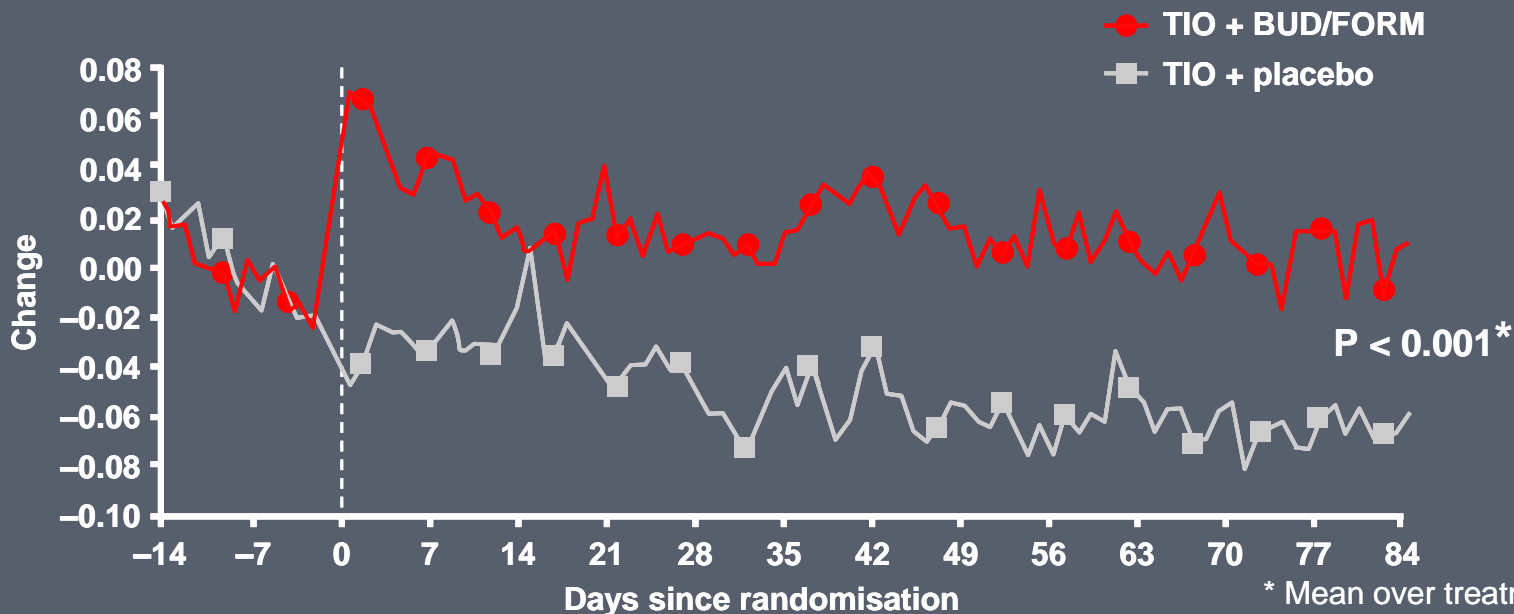
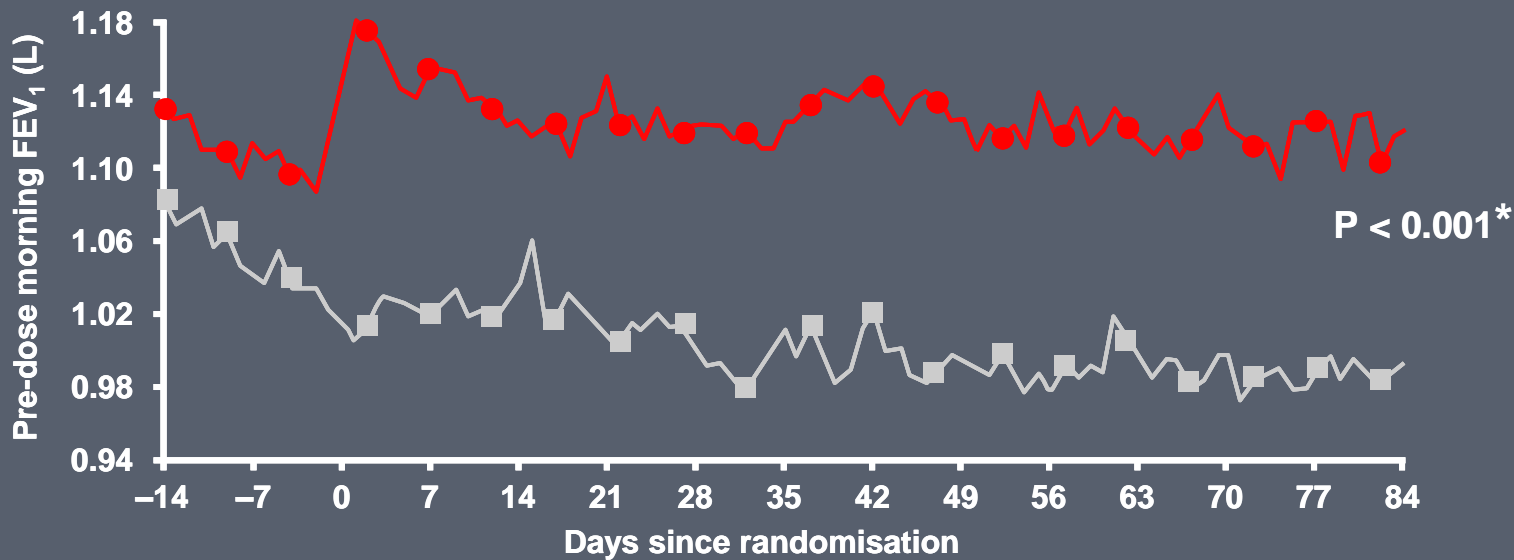
*Doses expressed as metered doses

†Oral and parenteral steroids not used ≥ 4 weeks before randomisation

TIO + BUD/FORM improves change in ratio of pre-dose FEV₁ compared with TIO + placebo

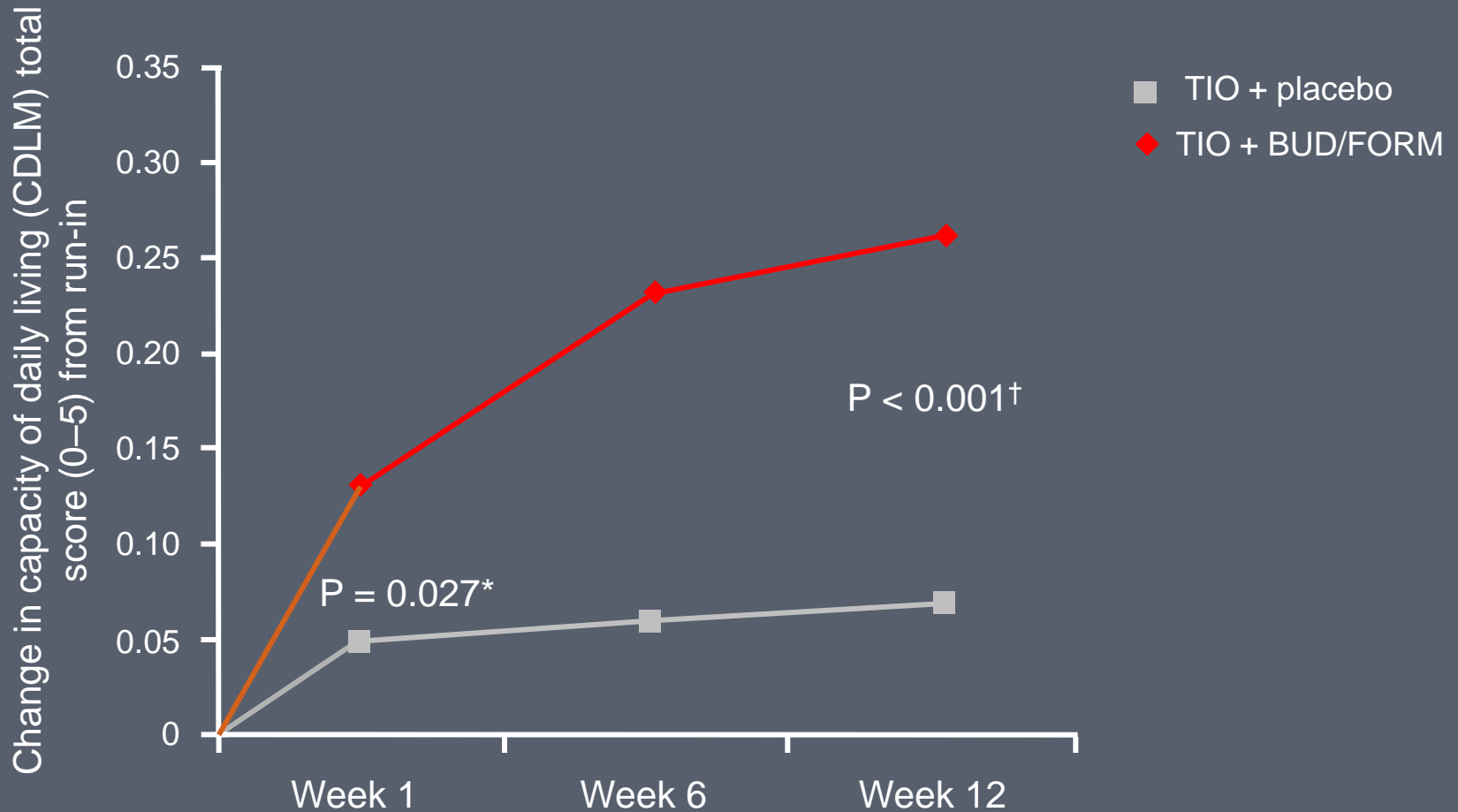


Morning FEV₁ at bedside



* Mean over treatment period

TIO + BUD/FORM improves change in total morning activities score



* Treatment comparison from randomisation to first week of treatment

† Treatment comparison from randomisation to last week of treatment

BUD/FORM = budesonide/formoterol; TIO = tiotropium

Welte T et al. AJRCCM 2009; doi:10.1164/rccm.200904-0492OC.

Thinking beyond the prescription

Let us begin with those with COPD

The patient pathway

Prompt Diagnosis

Optimisation of therapy

Oxygen

Vaccinations

Pulmonary Rehab

Easy access to knowledgeable HCP

Usual care –
Stable disease

Onset of an
exacerbation

Emergency
department/
General
Practitioner

Hospital
Admission

Discharge

Self Management Education

Reserve supplies of
antibiotics and steroids

Case Registers

Pulmonary Rehabilitation

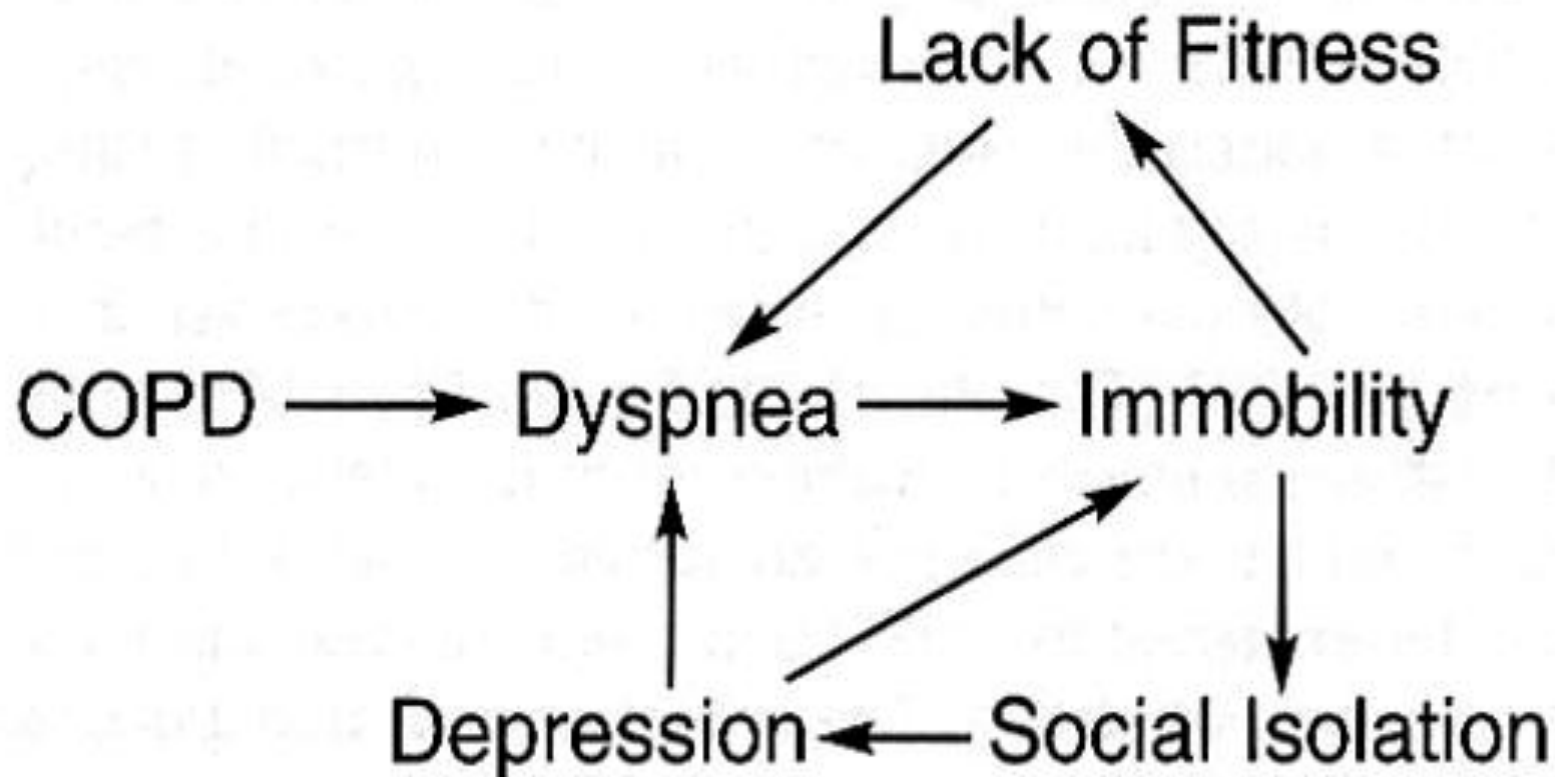
Consists of:

1. Exercise training
2. Nutrition counselling
3. Education,

And there is grade A evidence that it:

- Improves exercise capacity
- Reduces perceived intensity of breathlessness
- Reduces number of hospitalisations and length of stay
- Reduces the anxiety and depression associated with COPD

The cycle of Physical, Social, and Psychosocial Consequences of COPD



**LONG TERM DOMICILIARY OXYGEN THERAPY
IN CHRONIC HYPOXIC COR PULMONALE
COMPLICATING CHRONIC BRONCHITIS AND
EMPHYSEMA**

Report of the Medical Research Council Working Party*

Summary A controlled trial of long term domiciliary oxygen therapy has been carried out in three centres in the U.K. The 87 patients, all under 70 years of age, who took part had chronic bronchitis or emphysema with irreversible airways obstruction, severe arterial hypoxaemia, carbon dioxide retention, and a history of congestive heart failure. The patients were randomised to oxygen therapy (treated) or no oxygen (controls). Oxygen was given by nasal prongs for at least 15 h daily, usually at 2 l/min. The two groups were well matched, both clinically and in terms of lung function and other laboratory findings. 19 of the 42 oxygen treated patients died in the five years of survival follow-up compared with 30 out of 45 controls: in the 66 men in this trial the survival advantage of oxygen did not emerge until 500 days had elapsed. Survival for the 12 female controls was surprisingly poor, 8 of them being dead at 3 years. Mortality was not easy to predict, though a summation of arterial carbon dioxide tension and red cell mass was helpful. Neither time spent in hospital because of exacerbations of respiratory failure nor work attendance were affected by oxygen therapy, but these patients were very ill at the start of the trial and many had already retired on grounds of age or ill-health. Physiological measurements suggested that oxygen did not slow the progress of respiratory failure in those who died early. However, in longer term survivors on oxygen, arterial oxygenation did seem to stop deterioration.

Introduction

THE prognosis for patients in whom chronic bronchitis and emphysema is complicated by hypoxic cor pulmonale and carbon dioxide retention is grave, the three year mortality rate varying from 32 to 100%.¹⁻⁴ Correction of this hypoxaemia

over a long period by domiciliary oxygen therapy reduces pulmonary hypertension⁵⁻¹⁴ and secondary polycythaemia⁹⁻¹⁴ in these patients, but the value of this treatment has not so far been assessed by a controlled clinical trial. In 1973 the Medical Research Council set up in the U.K. a multicentre controlled trial of long term oxygen therapy in such patients in Birmingham, Edinburgh, and Sheffield. This trial was planned to determine if oxygen, given for 15 h a day, over a three year period, could reduce mortality and improve exercise tolerance and working capacity. Changes in the physiological variables were also to be studied.

Patients and Methods

Patients

Men or women under 70 years of age were asked to participate if they had chronic bronchitis or emphysema with irreversible airways obstruction (forced expiratory volume in one second (FEV_{1,0}) < 1.2 litres) and an arterial oxygen tension (PaO₂) between 40 and 60 mm Hg when breathing air at rest. The mean pulmonary arterial pressure was recorded, but it was decided that resting pulmonary arterial hypertension should not be a criterion for entry into the study. Patients were admitted if they had one or more recorded episodes of heart failure with ankle oedema; arterial blood gas, FEV₁, and body weight measurements had to be stable in two repeated measurements at least 3 weeks apart. Patients with fibrotic or infiltrative lung disease, pneumoconiosis (category 2 or more), severe kyphoscoliosis, overt episodes of pulmonary embolism, systemic hypertension (diastolic pressure > 100 mm Hg under 60 years of age or > 110 mm Hg over 65 years of age), proven coronary arterial disease, or other life threatening diseases were excluded.

Patients who agreed to participate in the trial after careful explanation were then allocated to the treatment or control group by means of a table of random numbers, so that within each centre successive groups of 8 patients contained 4 treated patients and 4 controls. Clinical, physiological, and therapeutic details of the 87 patients at entry to the trial are summarised in tables I and II, which show that they had severe persistent airways obstruction, hypoxaemia, CO₂ retention, compensated respiratory acidosis, secondary polycythaemia, and pulmonary hypertension.

Procedure

Treatment other than oxygen was given according to the discretion of the clinician in charge of the patient, and included diuretics, digoxin, and antibiotics (table II). All patients were urged to give up smoking. All patients were seen at a clinic every 2 months throughout the trial, being admitted to hospital for other treatment as necessary, at the discretion of their clinician. The patients were otherwise at home, but visited from time to time by research registrars or technicians attached to the trial to check oxygen usage and, occasionally, to sample arterial blood at home.

*Professor Sir CHARLES STUART-HARRIS (chairman), Prof. J. M. BISHOP, Prof. T. J. H. CLARK, Prof. A. C. DORNHORST, Dr J. E. COTES, Prof. D. C. FLENLEY, Dr P. HOWARD, and Dr P. D. OLDHAM.

Report prepared by Sir CHARLES STUART-HARRIS, Professor FLENLEY, Professor BISHOP, Dr HOWARD, and Dr OLDHAM, with assistance from Dr P. M. A. CALVERLEY. The assistance of the following research fellows is gratefully acknowledged: Dr S. R. Brennan, Dr Calverley, Dr R. M. Jones, Dr R. G. E. Leggen, Dr M. D. Peake, Dr R. A. Stockley, Dr N. F. C. Cain.

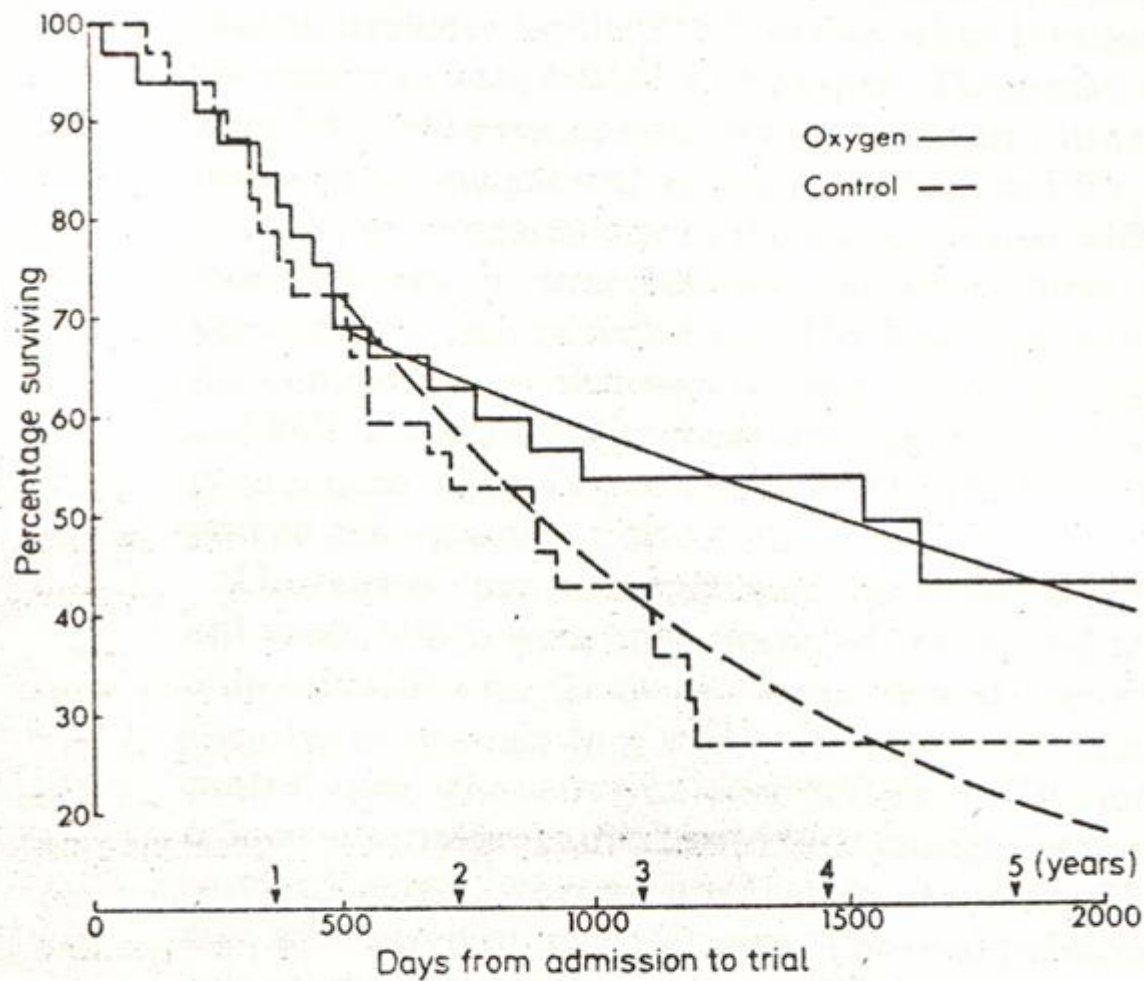
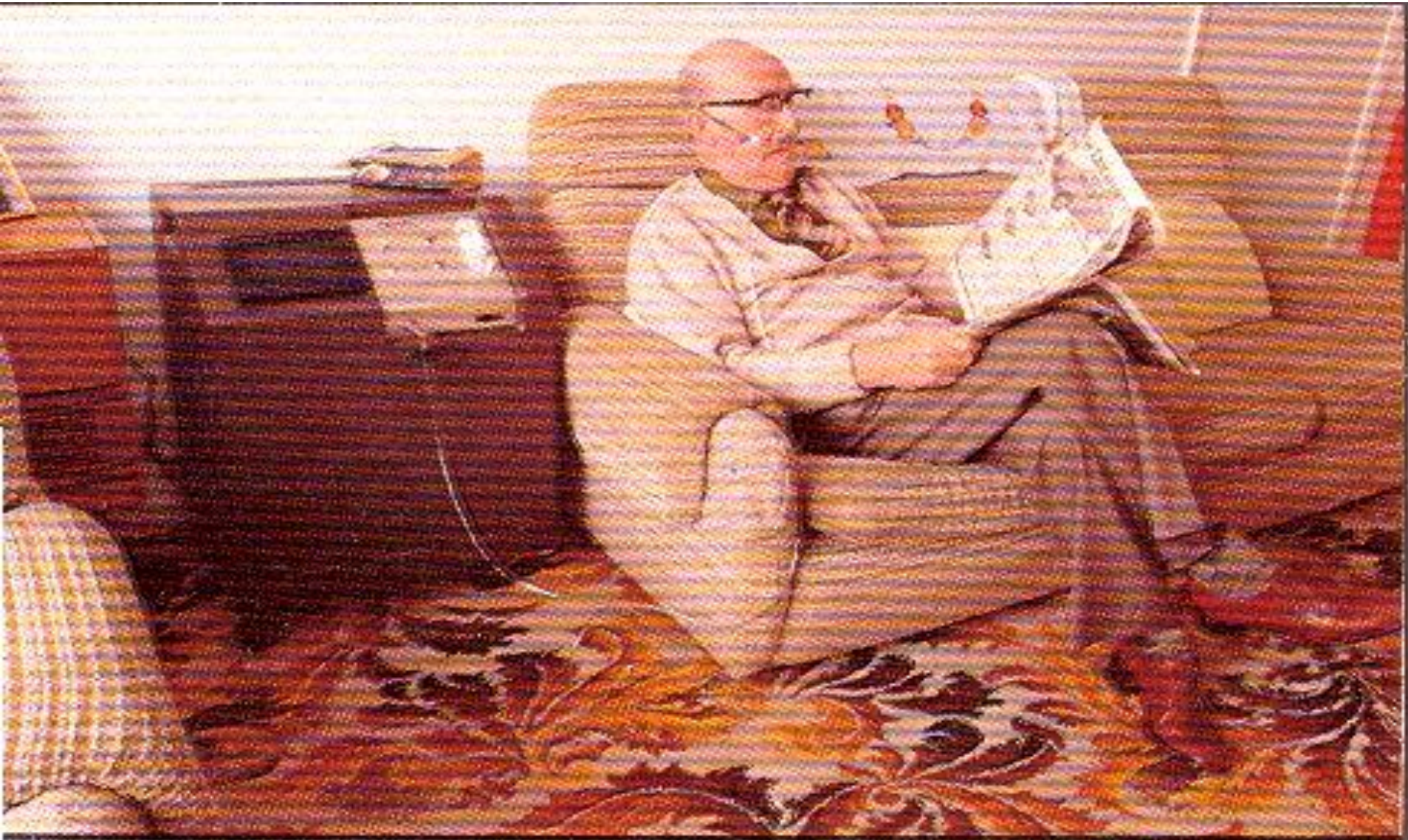


Fig. 1—Mortality in male patients.

Smooth curves indicate expected proportions surviving from 500 days, at constant risk of 11·9% per annum for those on oxygen, 29·4% per annum for the controls.

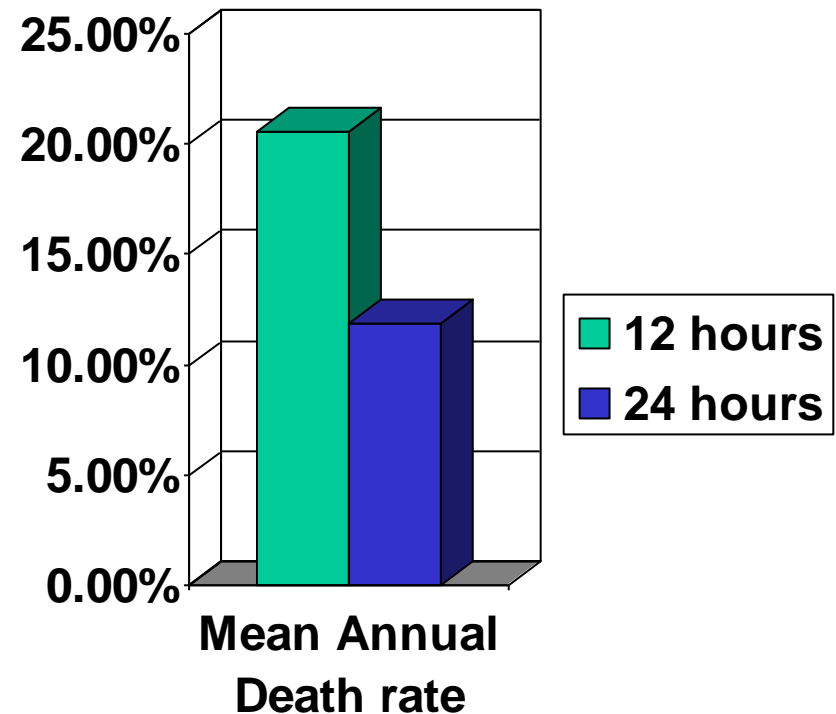
Domiciliary Oxygen from a concentrator



NHLBI NOTT study

(Ann Intern Med 1980;93:391-398)

- No control arm
- Compared 12 hours oxygen per day (102 patients) with 24 hours oxygen (101 patients)

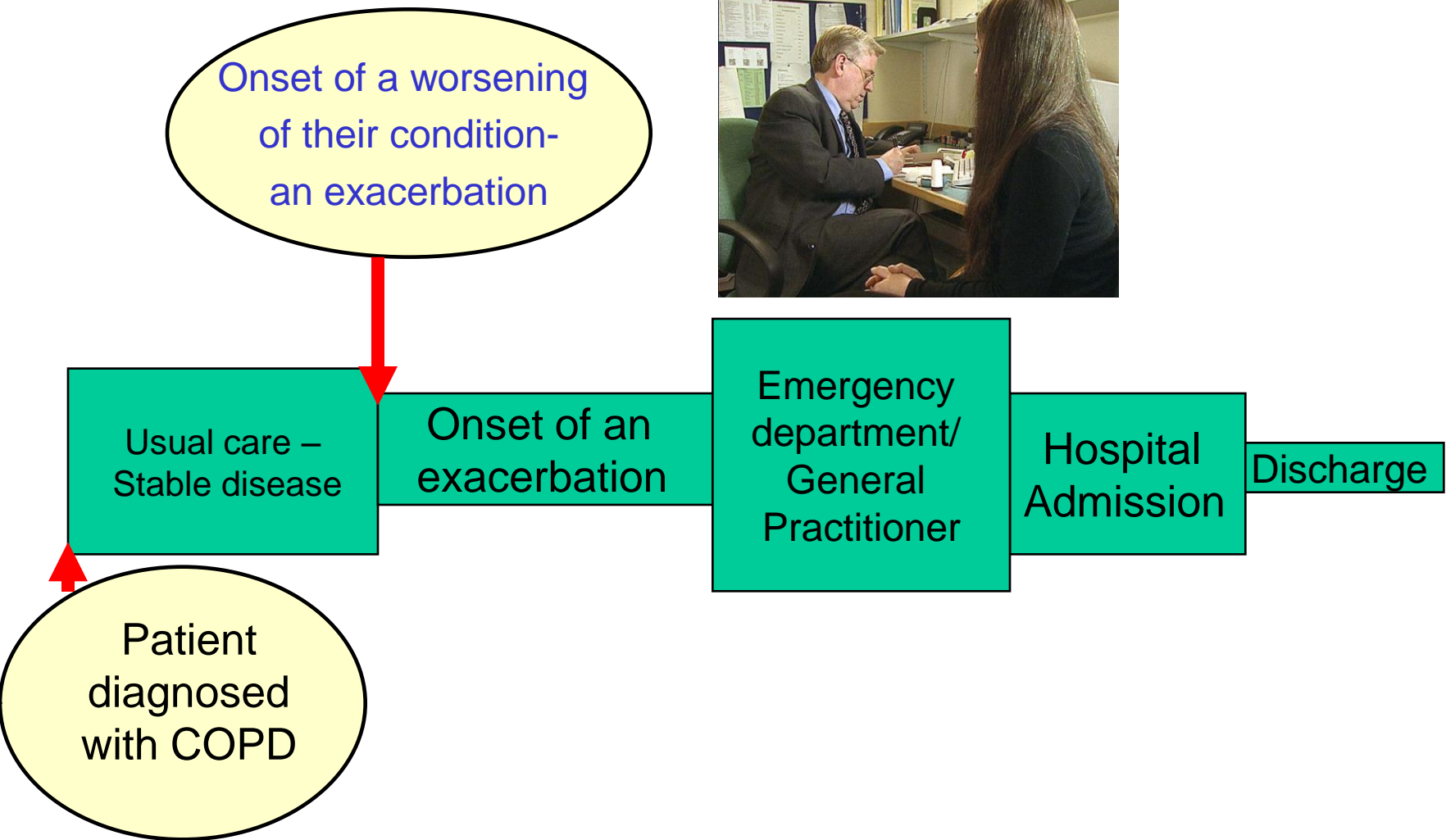


LTOT for COPD : The criteria

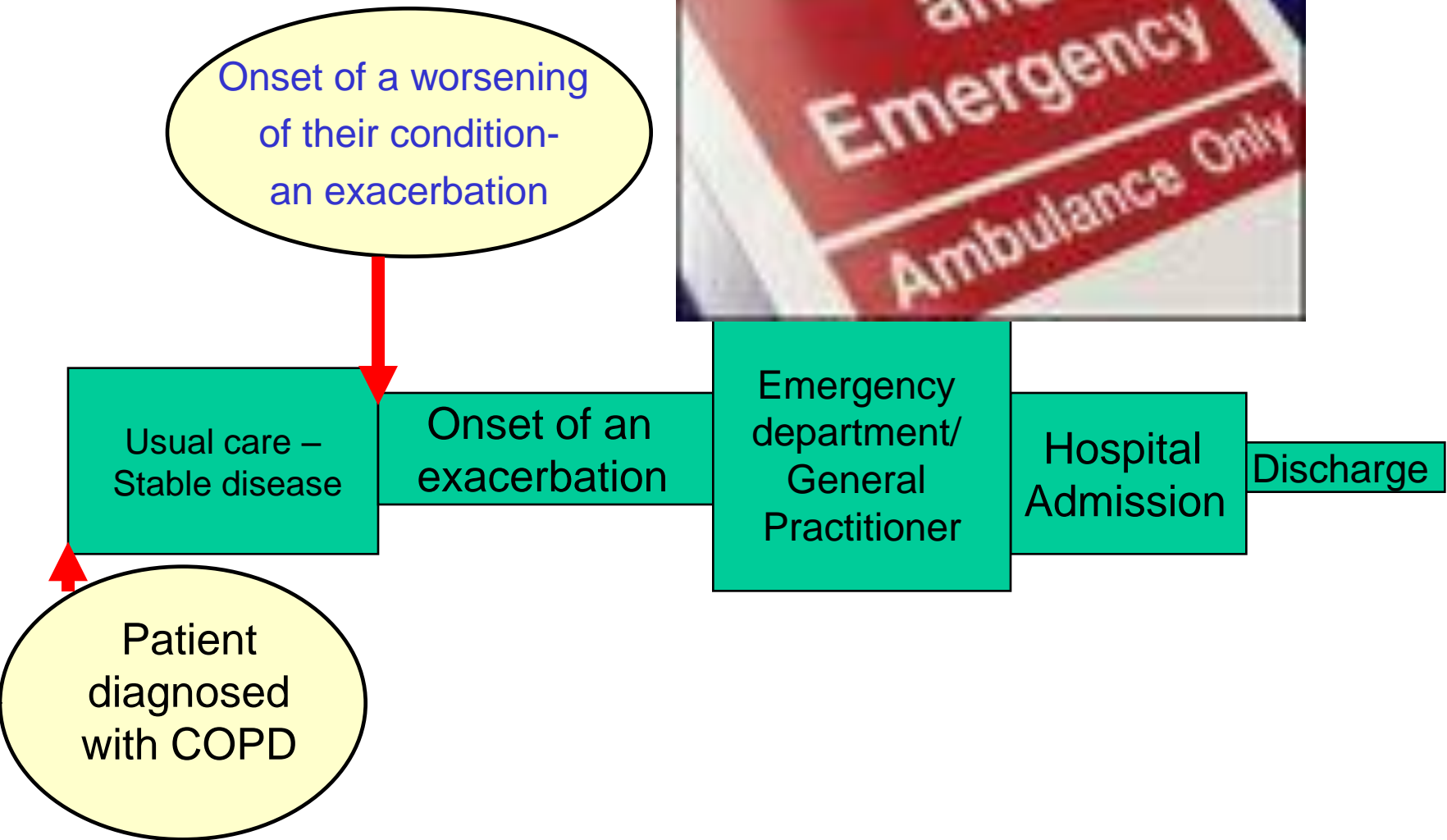
- PaO₂ <7.3 kPa when **stable** breathing air for at least 30 minutes after last supplementary oxygen, or
- PaO₂ between 7.3 and 8.0 when **stable** but plus one of: Secondary polycythaemia
Nocturnal Hypoxaemia (SaO₂ <90% for >30% of the time)
Peripheral oedema or pulmonary hypertension



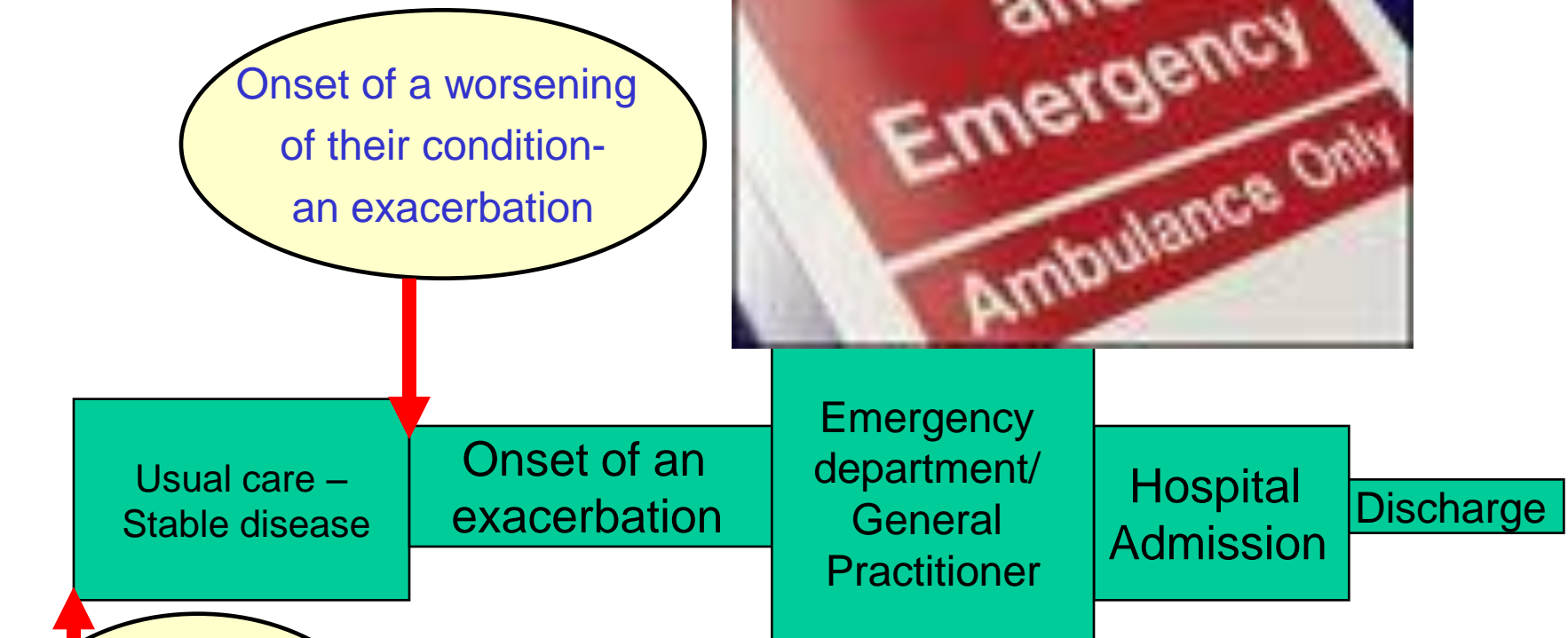
The patient pathway



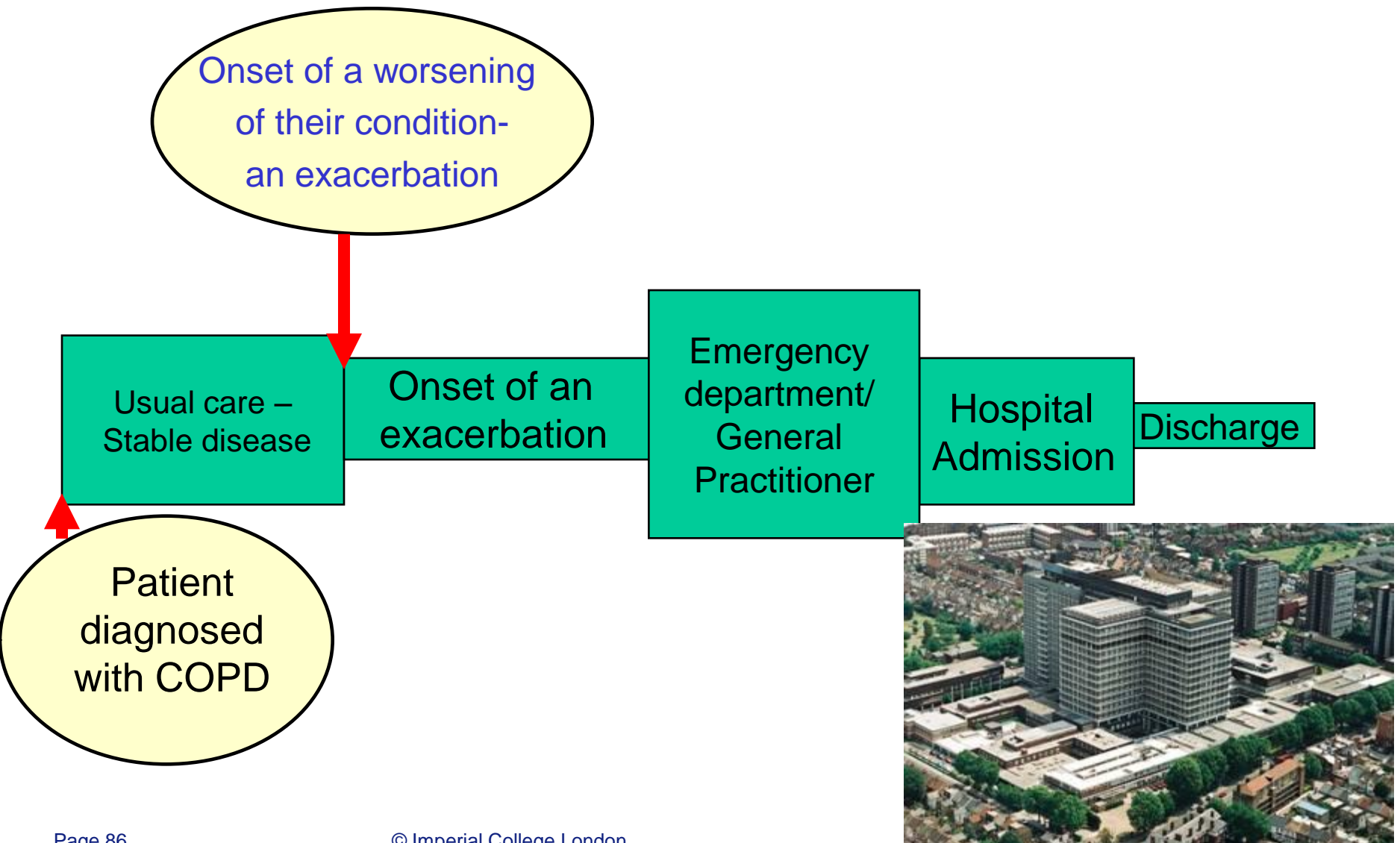
The patient pathway



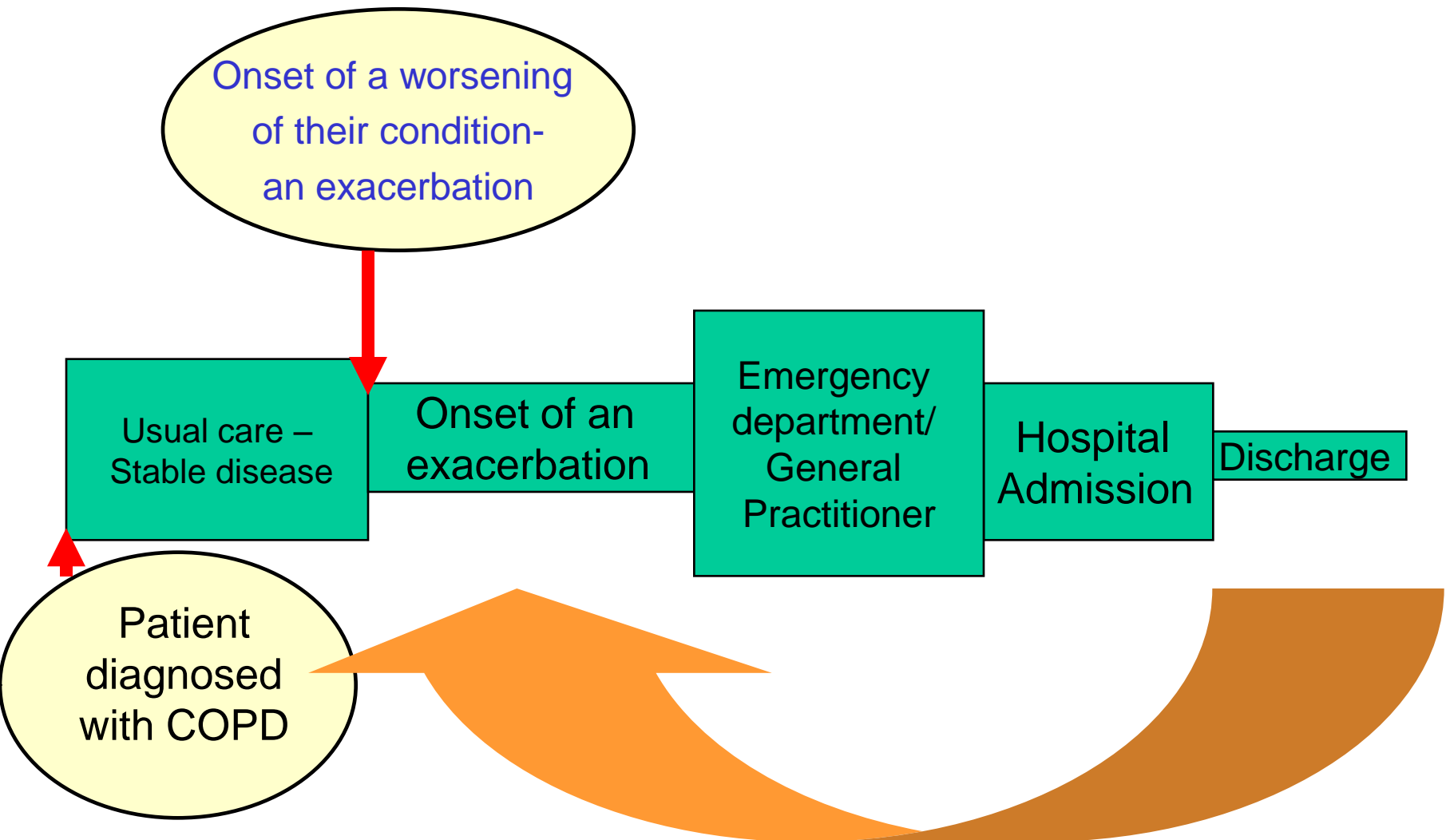
The patient pathway



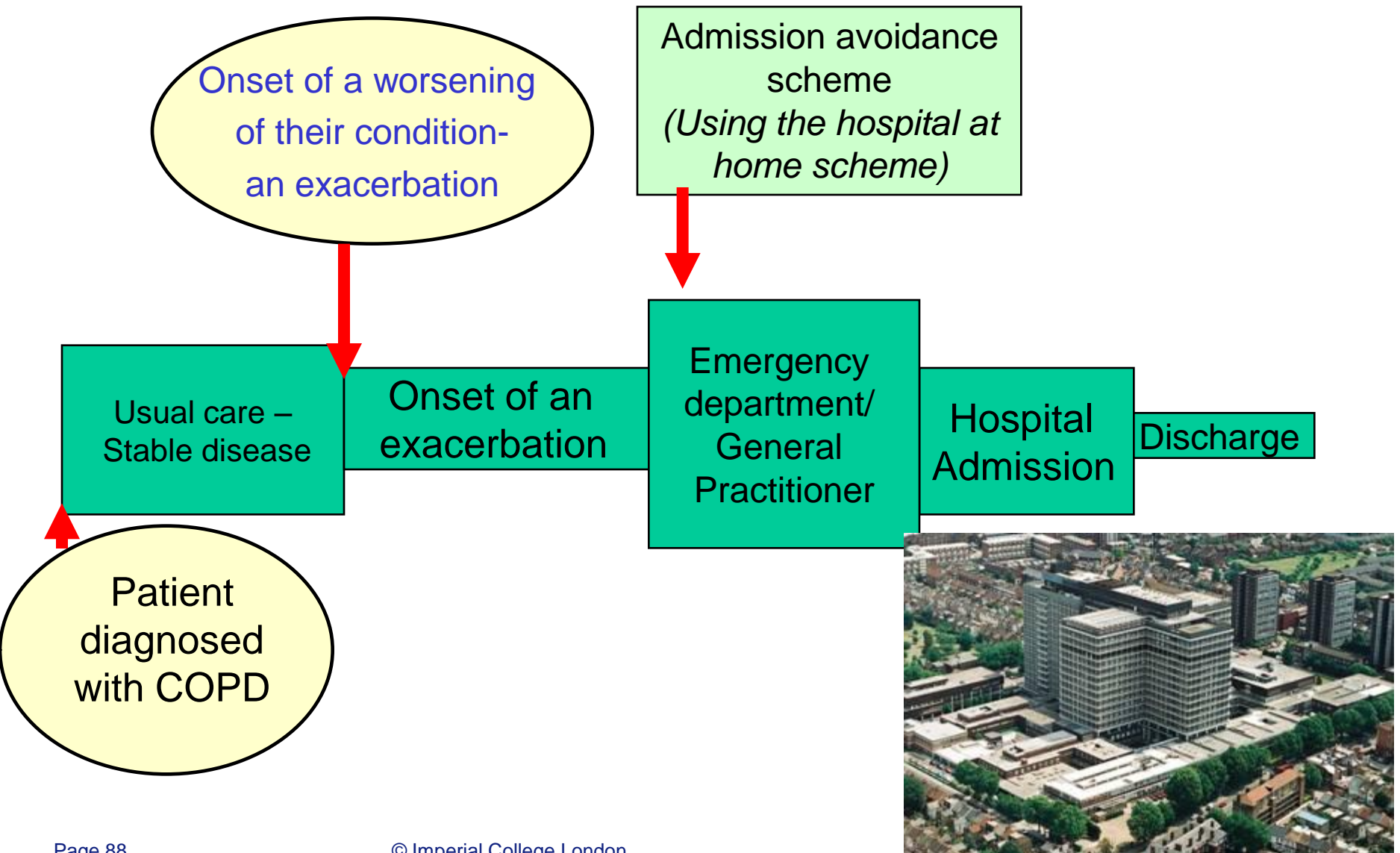
The patient pathway



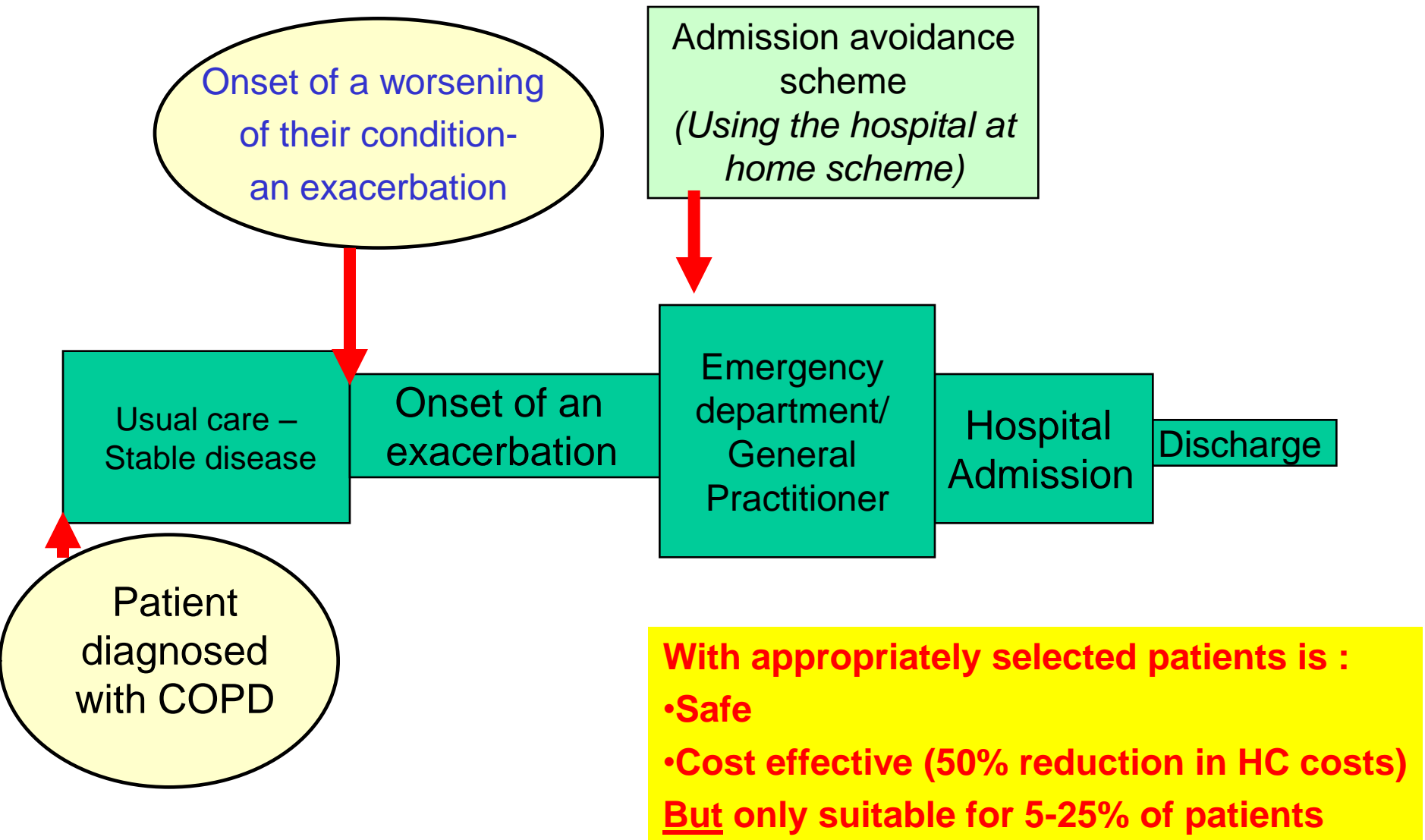
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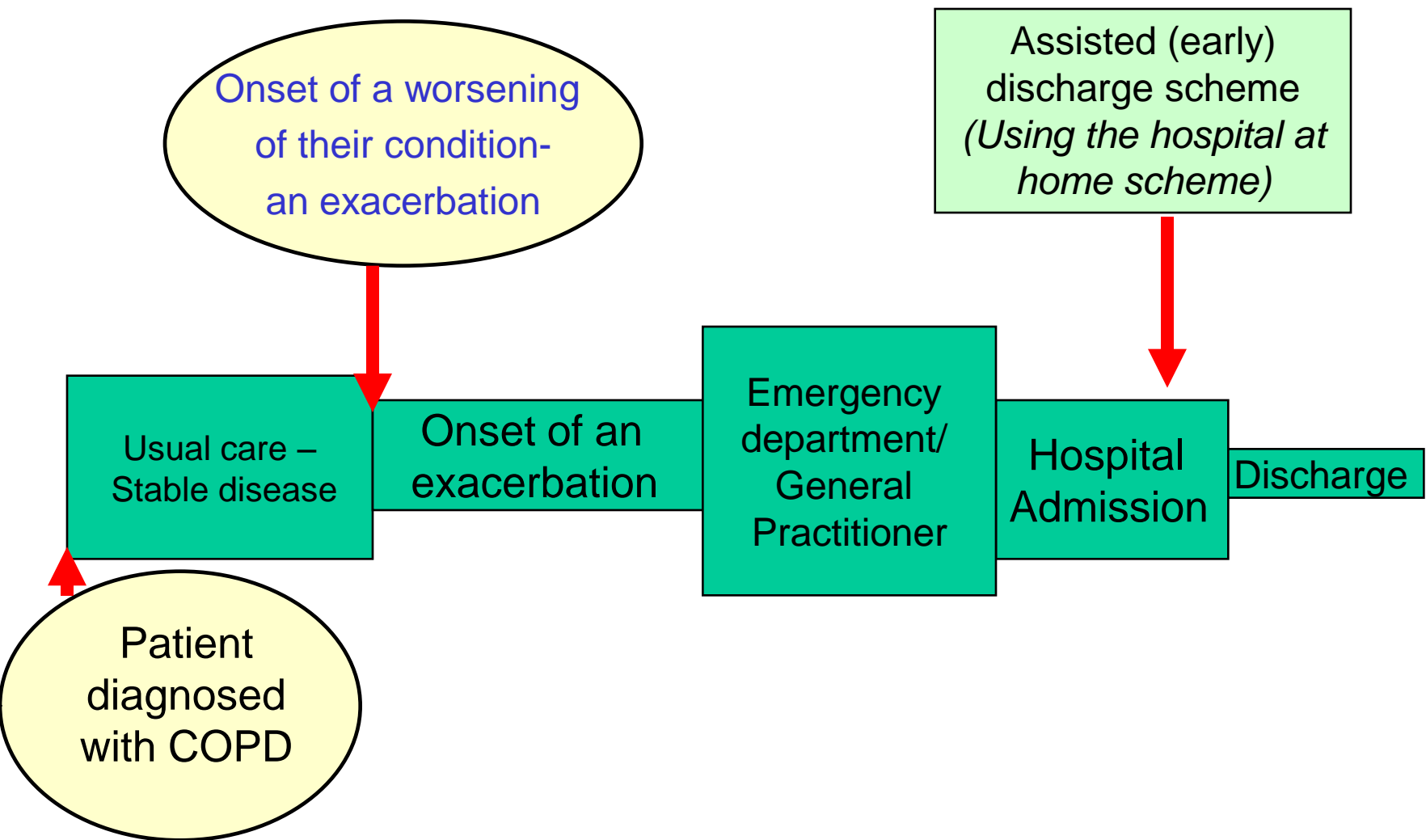
The patient pathway



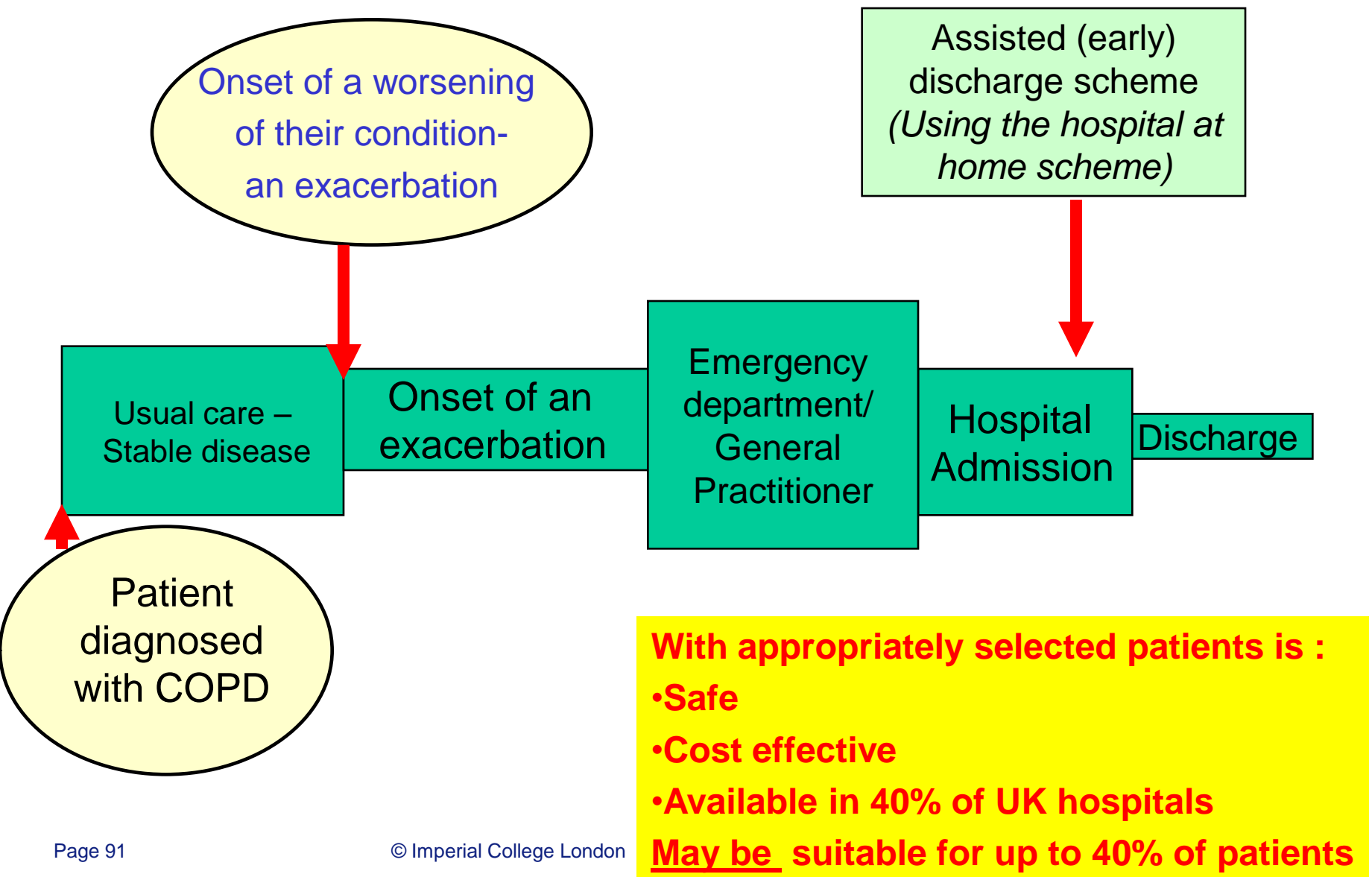
The patient pathway



The patient pathway



The patient pathway

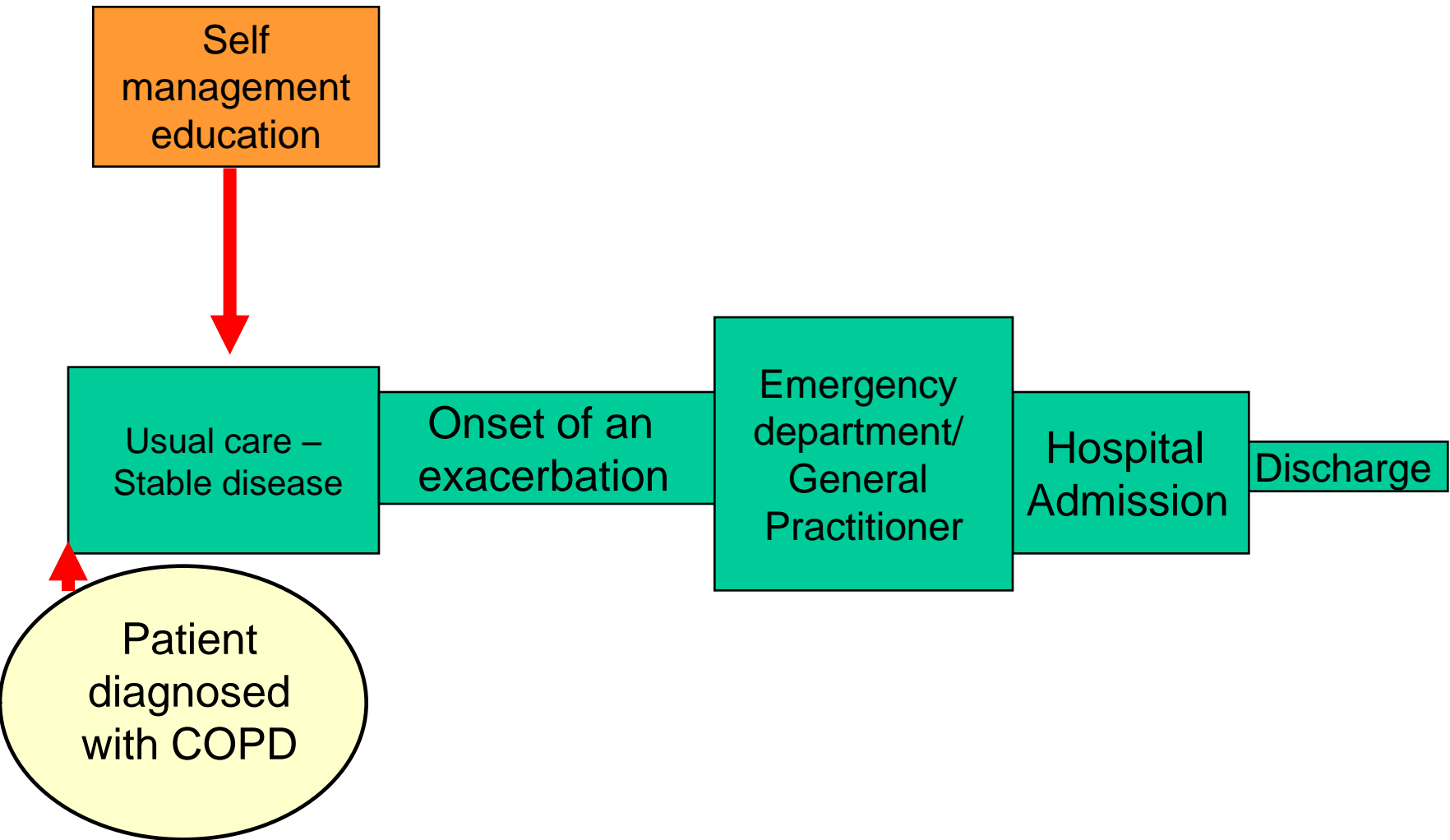


With appropriately selected patients is :

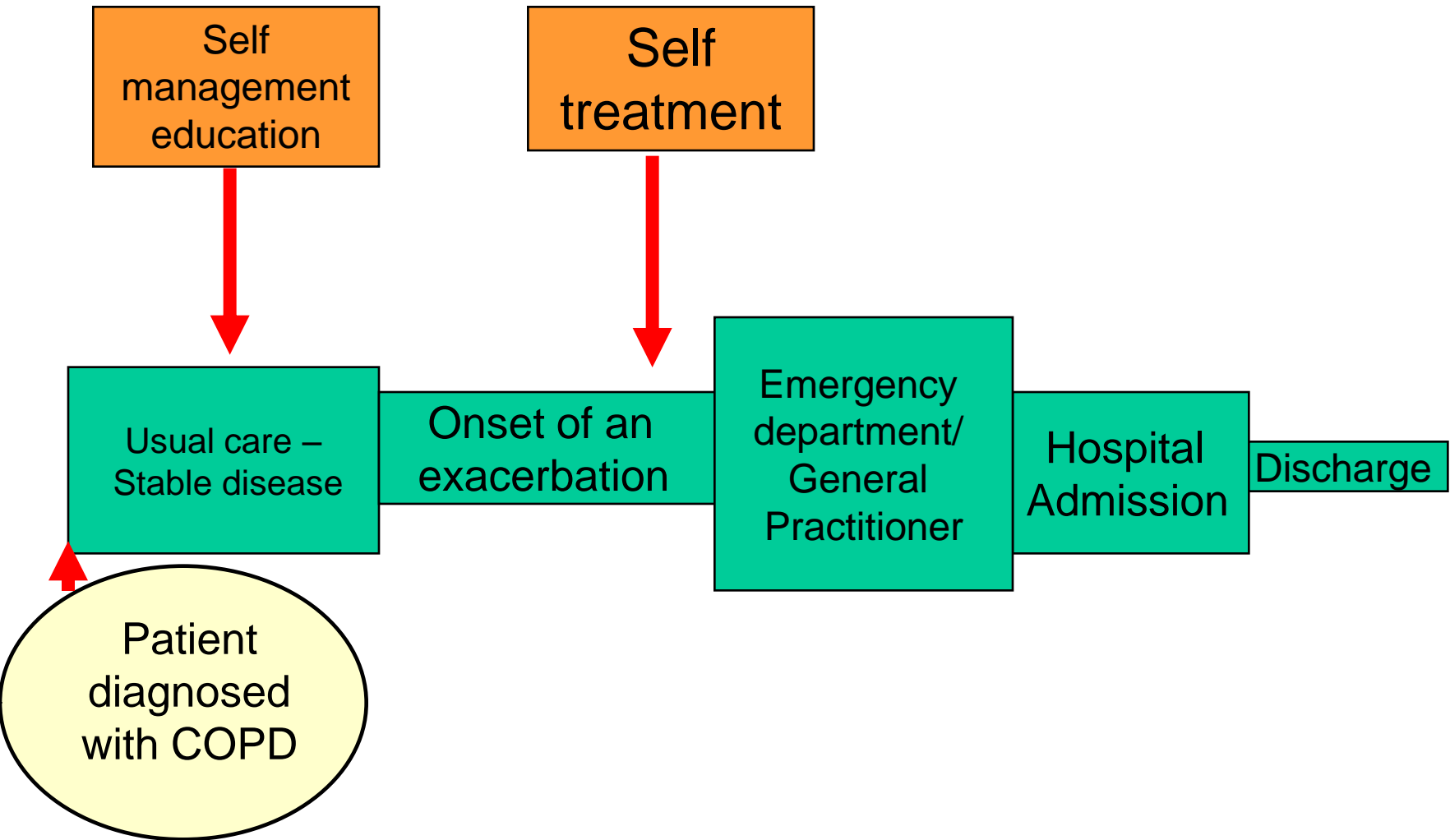
- Safe
- Cost effective
- Available in 40% of UK hospitals

May be suitable for up to 40% of patients

The patient pathway



The patient pathway



Does self Management Education work in COPD?

Does self management work in COPD?

Essentially too little data to yet say that it does

*Monninkhof E, van der Valk P, van der Palen J,
van Herwaarden C, Partridge MR, Zielhuis G
Thorax **2003**;58:394-398*

If results of self management in COPD are negative it might reflect:

- Poor studies
- Wrong intervention
- **Lack of written action plans**
- Ineffective interventions
- Wrong outcomes being monitored

Self management in COPD

Table 2 Characteristics of the self-management education intervention in each study

Reference	Group education	Individual education	Patient brochure	Audiotape	Exercise	Action plan	Smoking cessation	Nutrition
Gallefoss ^{8 15 16}	✓		✓			✓	✓	
Blake ¹⁷		✓	✓	✓				
Cockcroft ¹⁸		✓						
Emery ¹⁹	✓							
Gourley ²⁰ /Solomon ²¹		✓						
Howland ²²	✓				✓		✓	✓
Littlejohns ²³		✓						
Watson ²⁴			✓		✓	✓	✓	✓

Monninkhof E, van der Valk P, van der Palen J, van Herwaarden C, Partridge MR, Zielhuis G Thorax 2003;58:394-398

The need for action plans in COPD

Use of Action plans:

- Increases recognition of a severe exacerbation
- Increases use of antibiotics
- Increases use of oral steroids

But **NOTE only 3 studies included in review**
and results too few to assess effect on
healthcare utilisation

Turnock AC et al Cochrane Database 2005

**Department of Respiratory Medicine
Charing Cross Hospital**



**C.O.P.D. Self-management
Card**

Name:

Address:

Hospital No.:

Chest Specialists Name & Telephone Number:
.....

Respiratory Nurses Name & Telephone Number:
.....

General Practitioners Name & Telephone Number:
.....

How to Help Yourself

Lifestyle changes

1. Stop smoking, avoid smoking and avoid smoky environments.
2. If you are attempting to stop smoking take specialist advice about using nicotine replacement therapies and other smoking cessation medications and access local smoking support services as advised by your doctor or nurse
3. Remember that becoming breathless is not bad for you and regular exercise reduces the symptom of breathlessness, increases your activity levels and makes everyone feel happier!
4. Eat plenty of fruit and vegetables and those with COPD often find it preferable to eat small quantities more often. Gas forming foods such as onions, beans, sauerkraut, broccoli and cabbage are often best avoided. If you are very breathless you may find chewing uncomfortable and where it is possible to mash or liquidise food this may be easier to swallow. If eating makes you feel more breathless and you have oxygen available at home then discuss with your doctor or nurse whether taking additional oxygen using nasal cannulae (specs) may be worth trying.
5. Symptoms of coughing, sputum production and breathlessness are often worse first thing in the morning and this is a time when we often concentrate our physical activities such as exercising to dress, wash, make the bed, prepare breakfast etc. Make sure you have taken your medication as soon as you awake and try and spread out these early morning activities over a longer period to avoid distressing early morning symptoms. If showering is particularly difficult, the placing of a stool within the shower often helps. If you find drying yourself with a towel difficult and do not have a partner to do this for you, then sitting down and using a hair dryer to dry your back may be easier.
6. Ensure that you have annual flu vaccination and have had a vaccination against pneumonia.
7. Listen to weather forecasts and if very cold or polluted weather is expected make sure that you stay inside, drink plenty, take all your regular medications and if your condition does worsen adopt the advice contained within this action plan or seek medical attention.

How to Take Your Treatment and How to Alter Treatment if Your Condition Worsens

Treatment changes

Using your medications:

1. Take yourinhaler (.....) in the dose ofPuffs,times every day
2. Take yourinhaler (.....) in the dose ofPuffs,times every day
3. Take yourinhaler (.....) in the dose ofPuffs,times every day
4. If you feel any more breathless, you may take yourinhaler 2 puffs, every 3 to 4 hours to relieve symptoms and continue taking your regular inhalers as above.
5. If you notice more than two of the following situations then you should start your reserve supply of antibiotics and complete the whole course-
 - Increasingly short of breath
 - Increasing quantities of phlegm/sputum
 - Phlegm or sputum has turned persistently green
6. If despite taking your regular preventative therapies, and taking extra doses of your..... inhaler and despite starting a course of antibiotics you feel more breathless, then start six tablets (5mg strength) prednisolone immediately and repeat this dose every morning for 7 days before stopping the tablets. Let your doctor know after two days that you have started this course of steroid tablets.
7. If your ankles are more swollen than normal you should see your doctor.

If despite all these measures you still feel symptoms are worse then you should ring your doctor on.....

No. of Admissions 1Jan 2000 – 31Aug 2004
2305

No. of patients admitted / 4 years
1247

Alive: **740**

Dead: **507**

Assessment by standard proforma : **574**

Not proforma'd: **166**
(W11, W3, ETC)

Suitable: **297**

Unsuitable: **277**

Suitable & Recruited

Treatment: **61**

Control: **61**

Suitable Declined:

120

Suitable No Response:

55

Sridhar M, Taylor R, Dawson S,
Roberts NJ & Partridge MR
Thorax, 2008; 63: 194-200

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Roberts NJ & Partridge MR
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RESULTS

Intervention Group

Control Group

Exacerbation Treatment Initiator	Baseline N=61	Year 1 N=57	Year 2 N=55	Baseline N=61	Year 1 N=53	Year 2 N=49
Self			192/445 (43.1%)			38/364 P chi² <0.001 (10.4%)
Research Nurse			28/445(6.3%)			0 P chi ² <0.001
GP			140/445 (31.5%)			250/364 P chi² <0.001 (68.7%)
A&E doctor			4/445 (0.9%)			9 /364 (2.5%)
OP Clinic doctor			13/445 (2.9%)			49/364 (5.2%)

Self management behaviour much greater in those in the intervention group than amongst controls who were more likely to have treatment for exacerbations instituted by the GP

RESULTS

Intervention Group

Control Group

	Baseline N=61	Year 1 N=57	Year 2 N=55 (445 Exacerbations)	Baseline N=61	Year 1 N=53	Year 2 N=49
Unscheduled GP visits / contacts in 2 yrs			171			280 P chi² <0.001

Unscheduled need for contact with a GP was statistically much less in the intervention group where patients were more likely to self treat themselves

Sridhar M, Taylor R, Dawson S,
Roberts NJ & Partridge MR
Thorax, 2008; 63: 194-200

RESULTS

Intervention Group

Control Group

	Baseline N=61	Year 1 N=57	Year 2 N=55 (445 Exacerbations)	Baseline N=61	Year 1 N=53	Year 2 N=49
Unscheduled GP visits / contacts in 2 yrs			171			280 P chi ² <0.001

Unscheduled

much
likely to

And the death rate was halved
In the intervention group compared to controls

Sridhar M, Taylor R, Dawson S,
Roberts NJ & Partridge MR
Thorax, 2008; 63: 194-200

So what is the bottom line?

What are the key points in successful management of COPD?

Systematic Review of the Chronic care model in COPD *(Adams AG et al Arch Intern Med 2007;167:551-561)*

Trials that resulted in reduced health care use provided the following:

1. An extensive self management program with an individualized action plan
2. Advanced access to care (“Knowledgeable health care provider”)
3. Guideline based therapy
4. A clinical registry

Similarly for those with asthma we need to ensure

Similarly for those with asthma we need to ensure

- Is involved in decision making regarding their treatment
- Is offered control of their own condition
- Has support and follow up that is convenient

Guidelines for management of asthma in adults: I—chronic persistent asthma

Statement by the British Thoracic Society, Research Unit of the Royal College of Physicians of London, King's Fund Centre, National Asthma Campaign

The morbidity and mortality associated with asthma are unacceptably high in Britain. Many deaths and much unnecessary morbidity have been connected with underuse of inhaled and oral corticosteroids, underuse of objective measurements of severity, and inadequate supervision.^{1,2} As a result of an initiative from the British Thoracic Society, the research unit of the Royal College of Physicians of London, the King's Fund Centre, and the National Asthma Campaign a group of interested physicians and general practitioners was brought together to produce guidelines for the management of asthma in adults. This paper gives guidance on the management of chronic persistent asthma. Guidelines for the management of acute severe asthma will be published in next week's issue.

The recommendations for chronic persistent asthma promote greater use of anti-inflammatory drugs, even in patients with apparently mild asthma; objective monitoring of the progress of asthma using the patients' own measurements of peak expiratory flow; and greater participation of the patient in the management of the condition.

Description of asthma

Asthma is a common and chronic inflammatory condition of the airways whose cause is not completely understood. As a result of inflammation the airways are hyperresponsive and they narrow easily in response to a wide range of stimuli. This may result in coughing, wheezing, chest tightness, and shortness of breath; these symptoms are often worse at night. Narrowing of the airway is usually reversible, but in some patients with chronic asthma the inflammation may lead to irreversible obstruction of air flow.

Characteristic pathological features include the presence in the airway of inflammatory cells, plasma exudation, oedema, hypertrophy of smooth muscle, mucus plugging, and shedding of epithelium. These changes may be present even in patients with mild asthma when they have few symptoms.

Aims of management

The aims of management in adults are to recognise asthma, abolish symptoms, and restore normal or best possible long term function of the airways and reduce the risk of a severe attack. This last aim should be achieved by avoiding recognised causes if they exist and by using the lowest effective doses of convenient treatments with minimal short and long term side

at the step most appropriate for the initial severity of their condition. A short course of oral corticosteroids may be needed at any time and at any step to control their asthma (see box).

Treatment with short course of oral steroids

Short courses of oral steroids may be needed to control exacerbations of asthma at any step. Indications are:

- Symptoms and peak expiratory flow get progressively worse each day
- Peak expiratory flow falls below 60% of patient's best result
- Sleep is disturbed by asthma
- Morning symptoms persist until midday
- Maximum treatment not including oral steroids does not work
- Emergency nebuliser or injected bronchodilators are needed

Give patients prednisolone 30-60 mg daily (60 mg if they are already taking oral steroids) until two days after full recovery, when the drug may be stopped or the dose tapered.

If arranged beforehand short courses of oral steroids may be started on the patient's initiative according to written guidance.

Avoidance

If agents such as allergens, occupational sensitising chemicals, and certain drugs—for example, aspirin and non-steroidal anti-inflammatory drugs—are known to induce asthma in a patient they should be avoided if possible. β Blockers are contraindicated in patients with asthma. Avoidance of day to day triggers such as exercise and cold air generally imposes inappropriate restrictions on lifestyle, and it may be preferable to adjust treatment to cover exposure to these. Smoking should be avoided.

(1) Bronchodilators

A β_2 agonist (such as salbutamol 100-200 μ g or terbutaline 250-500 μ g) should be used as required rather than regularly. Inhalation is the preferred means of administration: the drug is delivered direct to the airway, doses are small, and side effects are minimised. In patients with normal lung function who have only infrequent symptoms and no sleep disturbance this may be the only treatment required.

(2) Inhaled anti-inflammatory agents

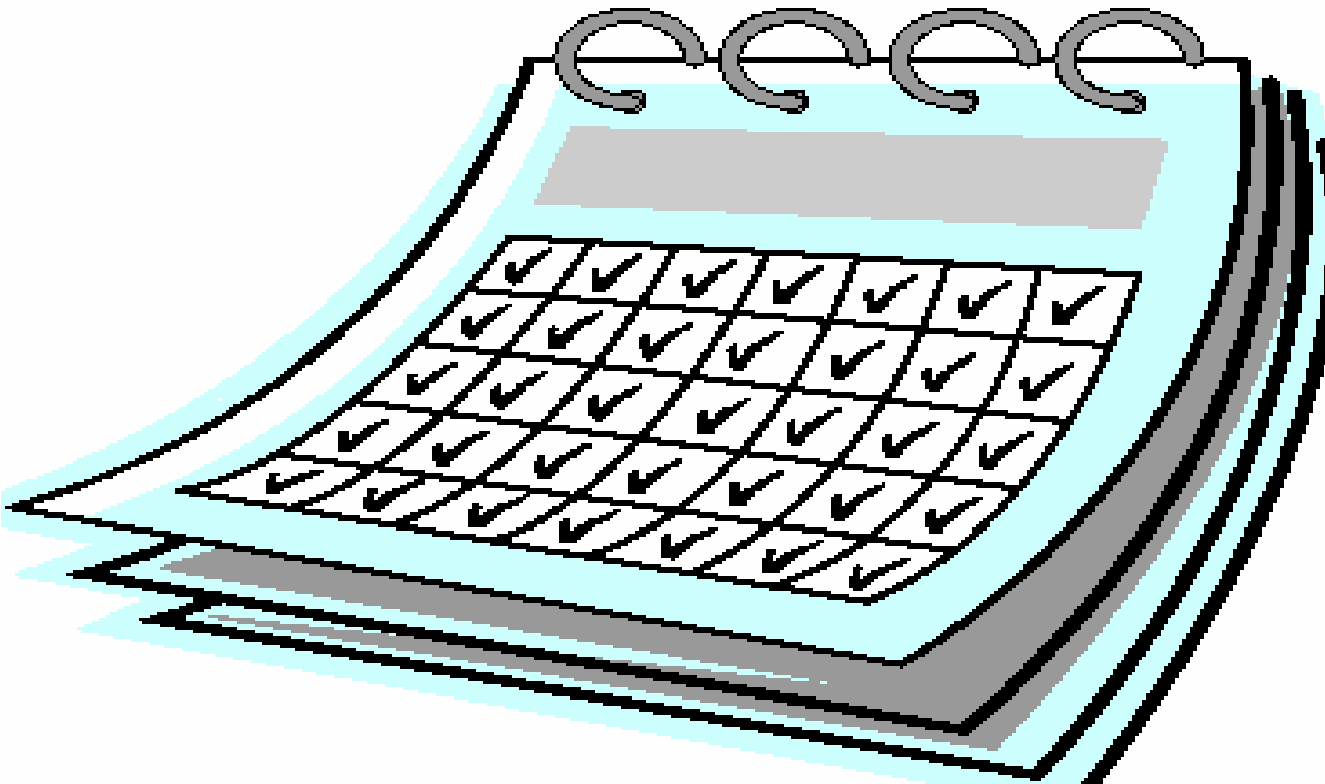
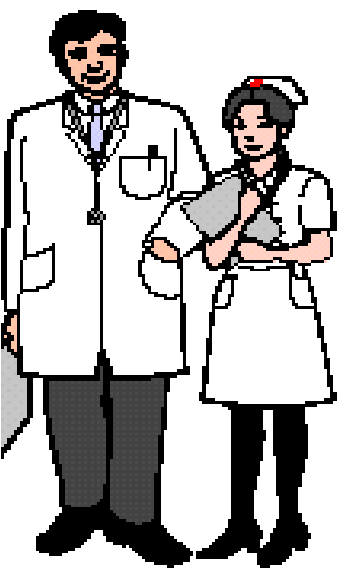
Patients who need to inhale a bronchodilator more than once daily or who have night time symptoms require regular inhaled anti-inflammatory drugs. Options include corticosteroids, sodium cromoglycate (5-20 mg four times daily), and nedocromil sodium

“ As far as possible patients should be trained to manage their own treatment rather than be required to consult their doctor before making changes”

A list of participants is given at the end of

Correspondence: British Thoracic Society, Andrew's Place, NW1 4LB.

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For the
other
364 days,
23 hours
and
30 minutes

.....

Conclusion 1:

- Asthma and COPD share symptoms
- Objective confirmation of diagnosis wherever possible and recording justification of how a diagnosis is made in the notes is important
- The aim in both conditions is both current control and reduction of future risk,
- In Asthma this means adding formoterol to the essential inhaled steroid and where appropriate using it both regularly and as required

Conclusion 2:

- In COPD this means adding inhaled steroids to formoterol to reduce exacerbations and taking the two medicines on waking (and possibly plus tiotropium) to reduce morning symptoms
- In both diseases we should adopt a patient centred approach which necessitates us hearing our patients concerns, and offering convenient support as the informed patient self manages their own condition.

Thank you

<http://www1.imperial.ac.uk/medicine/people/m.partridge/>