

Primary Care Respiratory **UPDATE**



www.pcrs-uk.org/pcru

HIGHLIGHTS ...

The importance of diagnosis

Fractional Exhaled Nitric Oxide Testing

Pull out wall chart: Spirometry

Nurses: Are you ready for revalidation?

Journal Round Up

Policy Round Up





**Around-the-clock COPD symptom control^{1,2}
with morning and evening administration.³**

Significant and sustained bronchodilation from the first dose vs placebo.^{1,2}

Prescribing information

(Please consult the Summary of Product Characteristics (SmPC) before prescribing.)

Eklira[®] Genuair[®]

322 micrograms inhalation powder aclidinium

Presentation: Each delivered dose (the dose leaving the mouthpiece) contains 375 µg aclidinium bromide (equivalent to 322 µg of aclidinium). Each metered dose contains 12.6 mg lactose monohydrate. **Indication:** Eklira Genuair is indicated as a maintenance bronchodilator treatment to relieve symptoms in adult patients with chronic obstructive pulmonary disease (COPD). **Dosage and Administration:** The recommended dose is one inhalation of 322 µg aclidinium twice daily. *Consult SmPC and package leaflet for method of administration.* **Contraindications, Warnings, Precautions:** **Contraindications:** Hypersensitivity to aclidinium bromide, atropine or its derivatives, including ipratropium, oxitropium or tiotropium, or to the excipient lactose monohydrate. **Precautions:** Should not be used to treat asthma or for relief of acute episodes of bronchospasm, i.e. rescue therapy. Paradoxical bronchospasm has been observed with other inhalation therapies. If this occurs, stop medicine and consider other treatment. Re-evaluation of the treatment regimen should be conducted if there is a change in COPD intensity. Use with caution in patients with a myocardial infarction during the previous 6 months, unstable angina, newly diagnosed arrhythmia within the previous 3 months, or hospitalisation within the previous 12 months for heart failure functional classes III and IV as per the "New York Heart Association". Consistent with its anticholinergic activity, dry mouth has

been observed and may in the long term be associated with dental caries. Also, use with caution in patients with symptomatic prostatic hyperplasia or bladder-neck obstruction or with narrow-angle glaucoma. Patients with rare hereditary problems of galactose intolerance, Lapp lactase deficiency or glucose-galactose malabsorption should not take this medicine. **Interactions:** Co-administration with other anticholinergic-containing medicinal products has not been studied and is not recommended. Although no formal *in vivo* drug interaction studies have been performed with Eklira Genuair, it has been used concomitantly with other COPD medicinal products including sympathomimetic bronchodilators, methylxanthines, and oral and inhaled steroids without clinical evidence of drug interactions. **Fertility, Pregnancy and Lactation:** It is considered unlikely that Eklira Genuair administered at the recommended dose will affect fertility in humans. Aclidinium bromide should only be used during pregnancy if the expected benefits outweigh the potential risks. It is unknown whether aclidinium bromide and/or its metabolites are excreted in human milk. The benefit for the breast-feeding child and long-term benefit of therapy for the mother should be considered when making a decision whether to discontinue therapy. **Ability to drive and use machines:** The effects on the ability to drive and use machines are negligible. The occurrence of headache or blurred vision may influence the ability to drive or use machinery. **Adverse Reactions:** **Common:** sinusitis, nasopharyngitis, headache, cough, diarrhoea. **Uncommon:** Blurred vision, tachycardia, dysphonia, dry mouth, rash, pruritus, urinary retention. **Rare:** Hypersensitivity. **Not known:** Angioedema. **Legal Category:** POM **Marketing Authorisation Number(s):** EU/1/12/778/002 - *Carton containing 1 inhaler with 60 unit doses. NHS Cost: £28.60 (excluding VAT)*

Marketing Authorisation Holder:

AstraZeneca AB
SE-151 85 Södertälje
Sweden

Further information is available from:

AstraZeneca UK Ltd.
600 Capability Green
Luton
LU1 3LU, UK
Tel: 0800 783 0033 / 01582 836836
Fax: +44 (0)1582 838 003
Email: Medical.informationuk@astrazeneca.com

Date of Revision: 03/2015

RSP 15 0020

Eklira and Genuair are both registered trademarks.

Adverse events should be reported. Reporting forms and information can be found at www.mhra.gov.uk/yellowcard. Adverse events should also be reported to AstraZeneca on 0800 783 0033.

References: 1. Jones PW, Singh D, Bateman ED, *et al.* Efficacy and safety of twice-daily aclidinium bromide in COPD patients: the ATTAIN study. *Eur Respir J.* 2012; 40(4):830-6. 2. Kerwin EM, D'Urzo AD, Gelb AF, *et al.* Efficacy and safety of a 12-week treatment with twice-daily aclidinium bromide in COPD patients (ACCORD COPD I). *COPD.* 2012;9(2):90-101. 3. Eklira Genuair Summary of Product Characteristics. Barcelona, Spain: Almirall, S.A.

Primary Care Respiratory Society UK National Primary Care Respiratory Conference



Expanding our horizons: delivering high value patient-centred care

15th-17th October, 2015
Whittlebury Hall, Northampton



The premier respiratory conference for primary care - offering essential clinical updates, and helping you work with your patients to transcend traditional boundaries and take a more integrated and holistic approach to care

REGISTER Online NOW
<http://www.pcrs-uk.org/pcrs-uk-annual-conference>

Charity Partners



Education Partners



education for health

Primary Care Respiratory UPDATE

The *Primary Care Respiratory Update* is published quarterly and distributed to members of the Primary Care Respiratory Society UK.

www.pcrs-uk.org/pcru

Editorial Office and Publishers

Primary Care Respiratory Society UK
Unit 2, Warwick House
Kingsbury Road
Curdworth, Warwicks B76 9EE
Tel: +44 (0)1675 477600
Fax: +44 (0)1361 331811
Email: gail@pcrs-uk.org

Advertising and sales

Contact Gail Ryan
Primary Care Respiratory Society UK
Unit 2, Warwick House
Kingsbury Road
Curdworth, Warwicks B76 9EE
Tel: +44 (0)1675 477600
Fax: +44 (0)1361 331811
Email: gail@pcrs-uk.org

Supplements and reprints

From time to time PCRS-UK publishes supplements to the regular journal, which are subject to review by the editorial board.

PCRS-UK also offers licencing opportunities for bulk reproduction of this journal.

For further information, contact Gail Ryan
Primary Care Respiratory Society UK
Unit 2, Warwick House
Kingsbury Road
Curdworth, Warwicks B76 9EE
Tel: +44 (0)1675 477600
Fax: +44 (0)1361 331811
Email: gail@pcrs-uk.org

Printed in the UK by Caric Print Ltd, Bournemouth, Dorset in association with Stephens & George Magazines Ltd. Printed on acid-free paper

Editor

Dr Hilary Pinnock, *Reader, Asthma UK Centre for Applied Research, Allergy and Respiratory Research Group, Centre for Population Health Sciences, University of Edinburgh General Practitioner, Whitstable Medical Practice, Whitstable, Kent*

Editorial board

Ian Culligan, *Specialist Respiratory Physiotherapist, Wirral*
Dr Stephen Gaduzo, *Chair PCRS-UK Executive, Stockport*
Dr Laura Ingle, *PCRS-UK Education Committee, and GP, Oxford*
Dr Basil Penney, *GPwSI in Respiratory Medicine, Darlington*
Anne Rodman, *Independent Respiratory Advanced Nurse Practitioner and Education for Health Regional Trainer, Lichfield*
Dr Iain R Small, *General Practitioner, Peterhead, Co-chair PCRS-UK Quality Award Development Group*
Ruth Thomas, *Senior Community Respiratory Nurse, Milton Keynes*
Steph Wolfe, *Independent Respiratory Nurse Specialist (Primary Care)*

PCRS-UK Chief Executive

Anne Smith

Communications Consultant and Freelance Journalist

Francesca Robinson

Policy Advisor

Bronwen Thompson

PCRS-UK Operations Director

Tricia Bryant

Competing interests are declared to PCRS-UK and this information is kept on file.

The opinions, data and statements that appear in this journal are those of the contributors. The publisher, editor and members of the editorial board do not necessarily share the views expressed herein. Although every effort is made to ensure accuracy and avoid mistakes, no liability on the part of PCRS-UK, the editor or their agents or employees is accepted for the consequences of any inaccurate or misleading information. © 2015 Primary Care Respiratory Society UK. All rights reserved. Apart from fair dealing for the purposes of research or private study, criticism or review, and only as permitted under the Copyright, Designs and Patent Act 1988, this publication may only be produced, stored or transmitted, in any form or by any means, with the prior permission in writing of Primary Care Respiratory Society UK. Enquiries concerning reproduction outside those terms should be submitted to Primary Care Respiratory Society UK via gail@pcrs-uk.org

The Primary Care Respiratory Society UK is a registered charity (Charity No: 1098117) and a company limited by guarantee registered in England (Company No: 4298947). VAT Registration Number: 866 1543 09. Registered offices: PCRS-UK, Unit 2 Warwick House, Kingsbury Road, Sutton Coldfield B76 9EE.
Telephone: +44 (0)1675 477600 Facsimile: +44 (0)121 336 1914
Email: info@pcrs-uk.org Website: <http://www.pcrs-uk.org>

The Primary Care Respiratory Society UK is grateful to its corporate supporters including AstraZeneca UK Ltd, Boehringer Ingelheim Ltd, Chiesi Ltd, GlaxoSmithKline, Napp Pharmaceuticals, Novartis UK, Pfizer Ltd and TEVA UK Ltd for their financial support which supports the core activities of the Charity and allows PCRS-UK to make its services either freely available or at greatly reduced rates to its members.

See http://www.pcrs-uk.org/sites/pcrs-uk.org/files/files/PI_funding.pdf for PCRS-UK statement on pharmaceutical funding.

“ My COPD means my appetite hasn't been very good...

...so I started taking Fortisip Compact Protein. It's very easy to take and I feel like I'm getting better. *Ron; Camden* ”

- Low 125ml volume and easy to take
- The most protein-rich, energy-dense nutritional supplement on the market
- Better compliance^{1*}

Why change to anything else?



*Greater compliance (91%) has been shown with more energy dense supplements (≥ 2 kcal/ml) such as Fortisip Compact Protein when compared to standard oral nutritional supplements.

Reference: 1. Hubbard GP *et al.* *Clin Nutr* 2012;31:293-312.

Nutricia Ltd., White Horse Business Park, Trowbridge, Wilts. BA14 0XQ.

Tel: 01225 751098.

www.nutriciaONS.co.uk

SCG2646-11/14



NUTRICIA
Fortisip[®]
Compact Protein

Right patient, right product, right outcomes

NEW FOR COPD

Duaklir[®]Genuair[®] ▼
aclidinium bromide/formoterol

**BREATHE LIFE
INTO THE MOMENTS
THAT MATTER**



Orange device available in the UK from 1st July 2015

The **ONLY LAMA/LABA** to have demonstrated **IMPROVED** around-the-clock COPD symptom control vs its monotherapies.¹⁻³

PRESCRIBING INFORMATION

(Please consult the Summary of Product Characteristics (SmPC) before prescribing.)

Duaklir[®] Genuair[®] ▼

340 micrograms /12 micrograms inhalation powder aclidinium and formoterol fumarate dihydrate

Presentation: Each delivered dose (the dose leaving the mouthpiece) contains 396 micrograms of aclidinium bromide (equivalent to 340 micrograms of aclidinium) and 11.8 micrograms of formoterol fumarate dihydrate. Each delivered dose contains approximately 11 mg lactose (as monohydrate). **Indication:** Duaklir Genuair is indicated as a maintenance bronchodilator treatment to relieve symptoms in adult patients with chronic obstructive pulmonary disease (COPD). **Dosage and Administration:** The recommended dose is one inhalation of Duaklir Genuair 340 micrograms /12 micrograms twice daily. Consult SmPC and package leaflet for method of administration. **Contraindications, Warnings, Precautions:** **Contraindications:** Hypersensitivity to the active substances or to the excipient lactose monohydrate. **Precautions:** Should not be used to treat asthma or for treatment of acute episodes of bronchospasm, i.e. rescue therapy. In clinical studies, paradoxical bronchospasm was not observed at recommended doses. Paradoxical bronchospasm has been observed with other inhalation therapies. If this occurs, stop medicine and consider other treatment. Use with caution in patients with a myocardial infarction during the previous 6 months, unstable angina, newly diagnosed arrhythmia within the previous 3 months, QTc above 470 msec or hospitalisation within the previous 12 months for heart failure functional classes III and IV as per the "New York Heart Association". β_2 -adrenergic agonists such as formoterol fumarate dihydrate may produce increases in pulse rate and blood pressure, electrocardiogram (ECG) changes such as T wave flattening, ST segment depression and prolongation of the QTc-interval in some patients. If effects occur, treatment may need to be discontinued. Use with caution in patients with severe cardiovascular disorders, convulsive disorders, thyrotoxicosis and phaeochromocytoma. Metabolic effects of hyperglycaemia and hypokalaemia may be observed with high doses of β_2 adrenergic agonists. Consistent with its anticholinergic activity, use with caution in patients with symptomatic prostatic hyperplasia or bladder-neck obstruction or with narrow-angle glaucoma. Dry mouth has been observed and may in the long term be associated with dental caries. Patients with rare hereditary problems of

galactose intolerance, Lapp lactase deficiency or glucose-galactose malabsorption should not take this medicine. **Interactions:** Co-administration with other anticholinergic and/or β_2 -adrenergic agonist containing medicines has not been studied and is not recommended. Although no formal *in vivo* drug interaction studies have been performed with Duaklir Genuair, it has been used concomitantly with other COPD medicinal products including short-acting β_2 -adrenergic bronchodilators, methylxanthines, and oral and inhaled steroids without clinical evidence of drug interactions. Caution is advised in concomitant treatment with methylxanthine derivatives, steroids, or non-potassium-sparing diuretics as this may potentiate the possible hypokalaemic effect of β_2 -adrenergic agonists. β_2 -adrenergic blockers may weaken or antagonise the effect of β_2 -adrenergic agonists. If β_2 -adrenergic blockers are required (including eye drops), cardioselective beta-adrenergic blockers are preferred, and these should be administered with caution. The action of formoterol on the cardiovascular system may be potentiated with medicinal products known to prolong the QTc interval such as monoamine oxidase inhibitors, tricyclic antidepressants, antihistamines or macrolides. Such medicines that prolong QTc interval are known to increase the risk of ventricular arrhythmias and should be administered with caution. **Fertility, Pregnancy and Lactation:** It is considered unlikely that Duaklir Genuair administered at the recommended dose will affect fertility in humans. Duaklir Genuair should only be used during pregnancy if the expected benefits to the mother outweigh the potential risks to the infant. It is unknown whether aclidinium bromide (and/or its metabolites) or formoterol are excreted in human milk. The benefit for the breast-feeding child and long-term benefit of therapy for the mother should be considered when making a decision whether to discontinue therapy. **Ability to drive and use machines:** The effects on the ability to drive and use machines are negligible. The occurrence of headache or blurred vision may influence the ability to drive or use machinery. **Adverse Reactions:** **Common:** Nasopharyngitis, urinary tract infection, sinusitis, tooth abscess, insomnia, anxiety, headache, dizziness, tremor, cough, diarrhoea, nausea, dry mouth, myalgia, muscle spasm, peripheral oedema, blood creatine phosphokinase increased. **Uncommon:** Hypokalaemia, hyperglycaemia, agitation, dysgeusia, blurred vision, tachycardia, ECG QTc prolonged palpitations, dysphonia, throat irritation, rash, pruritus, urinary retention, blood pressure increased. **Rare:** Hypersensitivity, bronchospasm including paradoxical.

Legal Category: POM

Marketing Authorisation Number(s): EU/1/14/964/001 - Carton containing 1 inhaler with 60 unit doses

NHS Cost: £32.50 (excluding VAT)

Marketing Authorisation Holder:

AstraZeneca AB
SE-151 85 Södertälje
Sweden

Further information is available from:

AstraZeneca UK Ltd.

600 Capability Green

Luton

LU1 3LJ, UK

Tel: 0800 783 0033 / 01582 836836

Fax: +44 (0)1582 838 003

Email: Medical.information.uk@astrazeneca.com

Date of Revision: 02/2015

RSP 15 0017

Duaklir and Genuair are both registered trademarks.

Adverse events should be reported. Reporting forms and information can be found at www.mhra.gov.uk/yellowcard. Adverse events should also be reported to AstraZeneca on 0800 783 0033.

References: 1. Duaklir[®] Genuair Summary of Product Characteristics. Södertälje, Sweden: AstraZeneca AB. Revision date April 2015. 2. Singh D, Jones P, Bateman E, et al. Efficacy and safety of aclidinium bromide/formoterol fumarate fixed-dose combinations compared with individual components and placebo in patients with COPD (ACLIFORM-COPD): a multicentre, randomised study. *BMC Pulm Med.* 2014;14:178. 3. Singh D, Chapman KR, Make BJ, et al. LAC30-31: Effect of aclidinium bromide/formoterol fumarate fixed-dose combination (FDC) on night-time and early morning symptoms in COPD. *Eur Respir J.* 2014;44(Suppl 58):A2415.

▼ This medicinal product is subject to additional monitoring. This will allow quick identification of new safety information.

AstraZeneca 

Primary Care Respiratory **UPDATE**



SPECIAL FEATURES

Editor's Round-Up

Hilary Pinnock 7

Chair's perspective: The critical importance of diagnosis. Why we must get it right.

Stephen Gaduzo 8

Nurses: Are you ready for revalidation? PCRS-UK is here to help.

Francesca Robinson 11

REGULAR FEATURES

Policy Round-Up

Bronwen Thompson 15

Getting the Basics Right

Fractional Exhaled Nitric Oxide testing
Fran Robinson 17

Journal Round-Up 21

PCRS-UK News Round-Up 28

British Lung Foundation Professionals 30

Delivering Excellence Locally

Delivering Excellence Locally in West Yorkshire
Fran Robinson, Dr Anuj Handa 32

Delivering Excellence Locally in the East of England
Fran Robinson, Daryl Freeman 33

PCRS-UK Resources and Improvement Tools: Reviewing diagnosis 34

New respiratory leaders inspired by PCRS-UK leadership event 36

PCRS-UK Affiliated Group Leaders Meeting 36

Update your clinical practice: excerpt of educational item from *npj Primary Care Respiratory Medicine* 40

SPECIAL PULL-OUT FEATURE

Your essential guide to spirometry



Data you can rely on.
People you can trust.

Vitalograph COMPACT

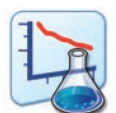
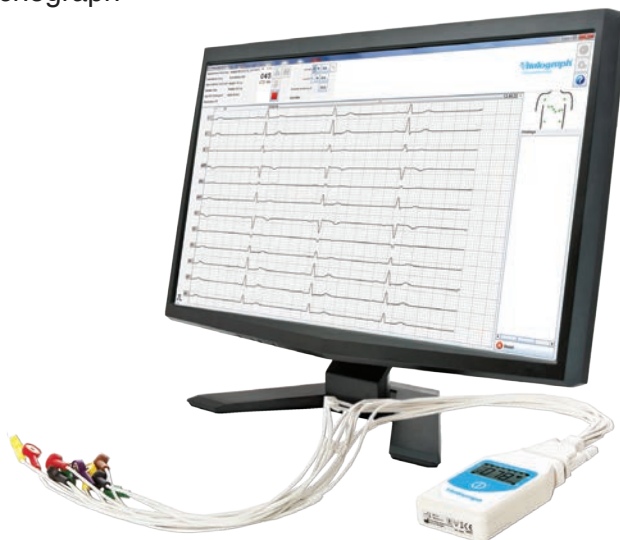
Sophisticated, integrated cardio-respiratory diagnostic workstation

The Vitalograph COMPACT™ Expert is a medical workstation that integrates high quality spirometry, 12-lead ECG, pulse oximetry, COPD assessment, blood pressure measurement, challenge testing and more in a flexible, integrated solution.

The COMPACT Expert saves medical professionals time, effort and money by integrating the management of subjects' data, visits and cardio-respiratory testing in one compact, easy-to-use device.



- High quality spirometry using Fleisch Pneumotachograph
- 12-lead ECG BT
- Bronchial challenge testing
- Pulse oximetry USB
- 6 minute walk test and COPD assessment
- Blood pressure USB
- Weighing scale BT
- Networks multiple COMPACT devices in their own network or to a central database on your existing networks



For more information call 01280 827110 or e-mail sales@vitalograph.co.uk
www.vitalograph.co.uk

Editor's Round-up

Hilary Pinnock, *Editor*



The theme of the September issue of the *Primary Care Respiratory Update* is diagnosis.

We were reminded of the importance of an accurate diagnosis in asthma by the release earlier in the year of the draft guideline from NICE on the diagnosis and monitoring of asthma, and challenged to consider whether our current approach is fit for purpose. It has sparked a debate that we cannot ignore and several of the articles in the *PCRU* pick up the theme. Stephen Gaduzo in his Chair's perspective outlines some of the concerns raised by PCRS-UK members, whilst Carol Stoneham in a practical 'Back to Basics' article shares her very positive experience of actually using Fractional Exhaled Nitric Oxide (FeNO). Whilst discussions continue, our best response as clinicians may be to reflect on the approach we adopt when we suspect that a patient's symptoms are due to asthma, and perhaps checking some of the practical PCRS-UK advice on spirometry highlighted in the centre-fold.

Diagnosis, of course, is central to our role in primary care where we daily see people presenting with undifferentiated symptoms and often very early in the course of a disease process before the classical symptoms and signs have developed. Distinguishing the common self-limiting causes for a symptom from the rare but

serious conditions is what we do – though it is not always an easy task. I attended a roundtable discussion on Interstitial lung disease recently and a colleague from a specialist clinic who saw several new patients a week with the diagnosis was taken aback when I calculated on published prevalence rates that a full time GP can expect to diagnose one new case every 7 years. The paper from the *npj Primary Care Respiratory Medicine* on raising the index of suspicion in primary care is a useful reminder of when we should suspect this group of rare but important conditions.

The imminent launch of revalidation will be high on the agenda of the 420 nurse members of PCRS-UK. A working party of PCRS-UK nurses have developed some practical support with a range of respiratory resources that will prove to colleagues as they prepare reflective portfolios.

As always the *npj Primary Care Respiratory Medicine* is a rich source of interesting and relevant evidence. Side effects of inhaled steroids, helping people with severe COPD, and dealing with the stress induced by incidental finding of pulmonary nodules are just three of the topics addressed in recent papers in our journal. These papers are free to access: do check them out on the *npj Primary Care Respiratory Medicine* website and share them with your colleagues.

Chair's perspective: The critical importance of diagnosis. Why we must get it right.

Stephen Gaduzo, *PCRS-UK Executive Chair*



A draft NICE guideline on the diagnosis and monitoring of asthma¹, published for consultation earlier this year, has sparked a robust debate about the most effective way of making an accurate asthma diagnosis.

It is welcome news that NICE is focusing on this vital area of care because PCRS-UK agrees that there is a need to improve asthma diagnosis. The clearer and more objective a diagnosis, the greater likelihood that it will take the clinician down the right treatment pathway, ensure treatment is as structured as possible and that patients will be given the right information and care. It is also an important factor in reducing variations in the quality of care that we know exist across the UK.

Recent reports have highlighted the need for improvements in asthma care. Significant shortfalls in care resulting in unnecessary deaths were revealed by data collated by The National Review of Asthma Deaths (NRAD).² More recently, a report from Asthma UK, called Patient Safety Failures in Asthma Care,³ revealed that one year on from NRAD there are still issues with safe prescribing in asthma. Getting the diagnosis of asthma right would go a long way to tackling these serious deficiencies and variations in care. In the guideline consultation document NICE cited studies showing that almost a third (30%) of people being treated for asthma no longer show signs of the condition, as a major reason for improving diagnosis.

NICE reinforce the advice that to achieve an accurate diagnosis objective clinical tests should be used as well as taking a full history, including signs and symptoms. In a departure from the existing BTS/SIGN guidance on the management of asthma⁴ it put spirometry and FeNO (fractional exhaled nitric oxide) measurements at the centre of asthma diagnosis. Although the BTS/SIGN guidelines with which we are all familiar, describe and recommend FeNO testing and spirometry as part of

the assessment in some patients, NICE proposed a more prominent place for these tests, and placed less emphasis on taking a full history, performing an examination and undertaking trials of treatment.

As a result, the algorithms it recommended set out an approach to diagnosing asthma that represents a huge shift from current practice, sparking a significant diversity of view among well respected clinicians both for and against the new direction of travel that NICE was recommending.

PCRS-UK has been a key player in this debate, drawing together the views of our members to ensure that we put forward a balanced view. Many PCRS-UK members are worried about the practical difficulties of implementing the new guidance. We presented NICE with the findings of a survey of almost 100 PCRS-UK members who said that what was proposed is neither realistic nor feasible in primary care. Almost eight in ten respondents said the proposed guidance would be "difficult" or "very difficult" to implement. Genuine doubt was also expressed that the guideline would actually improve asthma care in practice.

A key concern that PCRS-UK expressed to NICE was that these new algorithms underplay the importance of the well-established and holistic practice of taking a clinical history, examination, peak flow measurements and trials of treatment. We were not confident about the quality of evidence that lies behind NICE's new algorithms, and recommended to them that this new approach to the diagnosis of asthma should be piloted before being promoted for universal adoption. One particular problem is that FeNO measurements are reduced in people who smoke which may provide false reassurance in smokers.

NICE's enthusiasm for widespread FeNO testing is not currently reflected either in primary or secondary care. Few hospitals, including my own local

clinic in Stockport, offer it, suggesting that there is no widespread consensus yet as to its value.

However we know of some clinicians, such as Carol Stonham, a nurse practitioner and PCRS-UK's nurse lead, who have adopted FeNO testing in general practice and are very positive about it. In our article Back to Basics FeNO testing (see page 17) Carol explains why she finds the test a useful addition to the diagnostic process. There are also experts like Mike Thomas, Professor of Primary Care Research at the University of Southampton, who has conducted research into the use of FeNO testing and is a strong advocate for its place in general practice.

There are outstanding questions about the practical implementation of FeNO testing. Who will pay the capital cost of acquiring the new FeNO machines and the not insignificant on-going cost of consumables? Will practices be reimbursed for the longer appointments required to do the tests? If practices are unable or unwilling to make the investments required to provide the service, will they have to refer their patients to diagnostic hubs or secondary care to make a diagnosis of asthma? Have all the associated costs of such a dramatic change in practice been factored in to NICE's plan?

So what about the place of spirometry? The draft NICE guideline proposed that diagnosis is confirmed by spirometry in all patients, including children from the age of 5. While we agree with NICE that spirometry does have a valid role, particularly where there is diagnostic uncertainty, some of the pitfalls need to be recognised. For example: even when asthma is present, spirometry (and FeNO) may be normal. Accurate spirometry in young children, as recommended by the draft NICE guidance, is difficult to achieve in both primary and secondary care. Pressure on spirometry services would be sure to rise dramatically if the suggested new guidance is implemented.

The quality of spirometry in primary (and indeed secondary) care can be very variable. Many practices have spirometers, however

the healthcare professionals performing the tests and interpreting the results are not always adequately trained or fully competent in doing so. The quality of spirometry has to be the priority to address.

PCRS-UK has long argued for standards of spirometry to be improved and for practitioners to be accredited. Previous chairs, our research and policy leads and I, as well as representatives of other relevant organisations, have been involved in, sometimes quite heated, negotiations with NHS England to draw up guidance on quality assured spirometry. All the experts eventually reached agreement on the approach to training and accreditation that would be desirable for performing and interpreting spirometry. That was over a year ago and we are still waiting for these much needed standards to be published. We shall be using the more widespread use of spirometry proposed by NICE as ammunition to press NHS England to publish the spirometry guidance as a matter of urgency.

A final concern we have is that the proposed NICE guidance would present primary care with conflicting guidelines. We are urging NICE to collaborate with BTS/SIGN to ensure primary care has consistent, clear, guidance. If we end up with rival guidelines this will simply confuse clinicians. The danger is they may ignore the guidance altogether because it will seem to be too complex or difficult to follow which can only jeopardise efficient care for patients.

We have expressed all these concerns to NICE during the consultation phase of the draft guideline. We met Professor Mark Baker, director of clinical practice at NICE, and had a full and frank exchange of views with him. For me, the heart of this debate is about the need for primary care to find the most effective way of making an accurate diagnosis of asthma. The draft guidance proposed by NICE may be the right way to go in the longer term, but many of us feel that what they were proposing is too big a step at this stage. In short, PCRS-UK has forcibly argued that we need to put the brakes on and think carefully about how any such new approaches could be implemented.

NICE has announced that the final guideline will be delayed in order to allow additional time to work with commissioners and healthcare professionals in asthma care to make sure the recommendations can be introduced effectively and efficiently - suggesting that NICE has heard the concerns expressed by PCRS-UK and others.

In the meantime, the new approach to the diagnosis of asthma, proposed by NICE, has raised an important debate and I would urge you all to review your processes for the diagnosis and monitoring of asthma to ensure you do not have patients continuing to receive medication when they do not need it, or indeed patients who have their asthma diagnosis overlooked.

PCRS-UK has a wealth of online, evidence based resources, written by experts that can guide you in providing the highest standard of respiratory care for your patients. For example, we have recently updated our Quick Guide to Asthma, which takes into account the thinking that was proposed in the NICE draft guidance as well as the established approach advocated in the BTS SIGN guideline. Why not refresh your diagnostic skills by downloading our Diagnosis of Asthma in Adults and Diagnosis of Asthma in Children opinion sheets or take a look at our practice improvement tools.

PCRS-UK will continue to represent your views, as grassroots clinicians, in this debate and will continue to keep you up to date through e-alerts and social media when any new announcements are made about the eventual publication of the new guidance.

References

1. Asthma: diagnosis and monitoring of asthma in adults, children and young people. NICE. January 2015 <http://www.nice.org.uk/guidance/indevelopment/gid-cgwave0640>
2. Why asthma still kills. The National Review of Asthma Deaths. Royal College of Physicians. May 2014 <https://www.rcplondon.ac.uk/projects/national-review-asthma-deaths>
3. Patient safety failures in asthma care: the scale of unsafe prescribing in the UK. Asthma UK 2015. <https://www.asthma.org.uk/patient-safety>
4. BTS/SIGN Asthma guideline 2014 <https://www.brit-thoracic.org.uk/guidelines-and-quality-standards/asthma-guideline/>

ONE WAY

to treat persistent Air Leaks

Introducing a new, minimally invasive one way valve for treating air leaks.

Zephyr - the safe, effective way to block air flow

The Zephyr® Endobronchial Valve is designed to block air from passing through the leaking channel in the lung and into the pleural cavity. This one way valve reduces or eliminates airflow through the leak, allowing the tissue to heal and potentially allowing the patient to return to normal breathing.

- Self-expanding retainer covered in silicone
- One-way valve designed to allow trapped air and fluids to vent during exhalation
- Designed to be removable
- Single procedure for clinical response
- Can be successfully implanted during a variety of ventilation protocols



Actual size

For details of UK treatment centers go to: www.pulmonx.com
email uk-info@pulmonx.com
or call **07545 294 780**



pulmonX®

Revalidation

Nurses: Are you ready for revalidation? PCRS-UK is here to help

Francesca Robinson, *PCRS-UK Communications Consultant*, outlines new PCRS-UK materials to support nurse revalidation developed by PCRS-UK working party of nurses **Sally Harris**, **Ren Lawlor**, **Debbie Roots**, **Carol Stonham**, **Steph Wolfe** and **Tricia Bryant**



The Nursing and Midwifery Council (NMC) is shortly expected to give the go-ahead to revalidation for nurses and PCRS-UK is here to help you prepare with a wealth of resources that can be used to ensure you are providing a high standard of patient-centred respiratory care.

A report on the findings of the 19 sites that have been piloting the process will be published this month. The NMC Council will be making a final decision about proceeding with revalidation in October, with the first nurses and midwives expected to go through the process in April 2016.

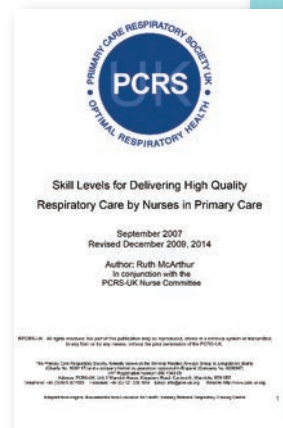
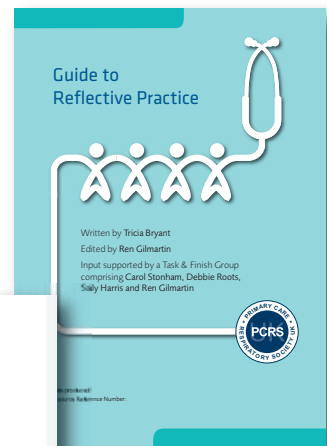
More than 2,100 participants from a range of health and social care settings took part in the pilots earlier this year and many felt that revalidation was a positive experience. Early indicators suggest that nurses and midwives found revalidation straightforward and that reflection against the Code provided an opportunity for individuals to think about their professional behaviour.

PCRS-UK is launching some new tools at the PCRS-UK conference in October that are designed to help primary care nurses understand how to undertake reflective practice and ensure their respiratory knowledge and skills are up to date.

These are two of the essential requirements of revalidation, which requires all nurses to demonstrate that they can practise safely in order to stay on the NMC register.

Nurses should already be starting to think about the revalidation requirements, which are more onerous than the previous PREP system under which nurses were only held to account if they were subject to a NMC spot check.

The new PCRS-UK tools – a Guide to Reflective Practice, a self-rating respiratory skills checklist and an updated skills evaluation document – will support nurses to reflect, in a structured way, what their current knowledge base is in relation to respiratory disease management. They are designed to help nurses to identify learning gaps and explore the minimum standards required for the respiratory work they do.



These tools can be used to prepare for revalidation using respiratory disease as a model. This could then be translated across other disease areas.

Reflective practice in respiratory care - a new guide for nurses

The new PCRS-UK Guide to Reflective Practice aims to demystify the formal process of reflection, with which many nurses are unfamiliar. This is a process nurses are required to undertake under the recently revised NMC Code of Practice¹, which now governs all areas of respiratory and other practice.

As part of the revalidation requirements, nurses will need to collate and provide a minimum of five reflective accounts of how feedback they have received has improved or affirmed their practice and have discussed these as part of their confirmation ("third party" checking).

The new PCRS-UK Guide explains that reflection on respiratory care is a way of considering and examining your own thoughts, actions and reactions, and sometimes those of others, to a given situation or event in order to gain a better understanding of yourself and to identify more effective ways of responding in future. The process can allow you to improve your critical thinking, change your approaches to patient care, promote self-awareness and improve your communication skills.

However, to reflect effectively one must be prepared to uncover one's own perceptions and to be objective about how these perceptions and subsequent judgements may have affected one's chosen actions.

Many nurses find this level of personal insight unnerving but the Guide says that reflection should be looked at in a positive way, as a process by which we can learn about ourselves, our colleagues and our environments in such a way that the result is an improvement in future care.

The PCRS-UK Guide explains in practical ways how nurses can reflect on their respiratory practice through: keeping an on-going personal record; reflecting on an area of practice; reflecting on the patient experience, reflecting on the clinic environment, reflecting

on peer feedback and on learning and sharing knowledge with the team. It suggests that nurses can use PCRS-UK practice worksheets on respiratory disease to reflect on their respiratory practice.

The Guide sets out as an example how the practice improvement worksheet 'Post-acute care bundle in COPD'² could be used to reflect on a specific area of practice. It recommends reading through and reviewing the resources provided as links in the worksheet then seeking the support of the practice team to identify a patient attending for an unscheduled COPD visit. The nurse could then arrange to review the patient within 48 hours of being seen and assess the patient as described in the worksheet.

It sets out the type of questions a nurse needs to ask in order to complete the reflective process.

These are:

- How did you feel/what were your thoughts about providing a care bundle for the patient?
- Did you learn anything new from the worksheet or its associated resources?
- What did you discover when you reviewed the patient?
- Were the outcomes of the consultation as you would expect or did you discover anything new about your own knowledge, the process for reviewing people with COPD, any organisation of care issues or anything new about the patient's own knowledge and management of their condition?
- What have you learned by implementing this care bundle?
- How will what you have learned influence your practice in future?

This tool and reflective process could also be used across other areas of their practice areas.

Are your respiratory skills up to date?

In order to meet the requirements of revalidation nurses will need to prove that their skills are up to date. This is to ensure they are practising safely and put patient expectations at the heart of practice. These are two standards required by the NMC Code and also a central focus of PCRS-UK's mission to promote high standards of patient-centred care.

To support nurses as they prepare for revalidation PCRS-UK has updated its skills document to help practitioners review their respiratory expertise. The document sets out skill levels for delivering high quality respiratory care by nurses in primary care. It sets out in detail the various skills required for minimal, medium and maximum involvement in respiratory care.

Nurses can use the document to identify any gaps in their skills and knowledge. It will provide them with the evidence they need to request time and resources from their managers to attend any training courses they might need to fill those gaps. They will be able to argue that this training is essential for ensuring they are practising safely in line with the requirements of the new NMC Code.

Rate your respiratory skills

In order to help nurses rate their respiratory skills PCRS-UK has produced a new skills checklist. It covers various skills relating to asthma, COPD, inhaler devices, smoking and smoking cessation, allergic problems, symptoms in respiratory disease assessment, management and diagnosis, co-morbidities, communication and record keeping.

Nurses can rate themselves from "I feel very confident about this" to "I need to learn a lot more about this" to identify where there are gaps in their knowledge and skills.

Undergoing this self-rating process will enable nurses to evaluate in which areas of practice they might need either to refresh, update or extend their knowledge.

How PCRS-UK resources can help you prepare for revalidation:

- **Opinion sheets³:** These address key topics ranging from managing allergic rhinitis and asthma to a guide to using spirometry, and are an invaluable resource for updating and refreshing respiratory skills. Working through them will earn CPD points. For revalidation nurses will be required to complete 40 hours CPD over a three-year period.
- **Practice improvement worksheets⁴:** These enable nurses to assess where improvements could be made in their practice which would lead to better outcomes for patients and more efficient use of clinical time. Using these worksheets will help nurses to demonstrate the four principles of the new NMC Code: prioritising people; practising effectively; preserving safety and promoting professionalism and trust.
- **Up to date guidance on COPD and asthma.** Reading these guides will help to ensure you are following the latest, most up to date guidance. This will ensure you are practising safely and to a high standard.
- **PCRS-UK local affiliated respiratory nurse groups.** Joining a local nurse group will enable you to share good practice, keep yourself up to date and meet like-minded colleagues who are enthusiastic about providing high standards of respiratory care. These groups also provide opportunities for peer review, one of the requirements of revalidation.
- **PCRS-UK annual conference⁵.** Attending the PCRS-UK annual conference on 16-17 October at Whittlebury Hall, Northampton, will not only earn you CPD points but will ensure you hear about the latest clinical updates and cutting edge re-

At a glance... Revalidation requirements

Nurses will need to ensure they are familiar with the NMC's new code of conduct¹, which governs standards of practice.

The Code places patient expectations at the heart of professional practice and reflects changes in healthcare over the last seven years, including the Francis report into care failings at Mid Staffordshire Foundation Trust.

The Code is built around four principles:

- prioritising people
- practising effectively
- preserving safety
- promoting professionalism and trust

The requirements of revalidation are that nurses:

- Complete 450 minimum hours of practice and 40 hours CPD over a three year period
- Obtain a minimum of five pieces of feedback over a year period from a range of sources
- Record at least five reflections on this feedback, the Code and/or learning activities undertaken, and have a professional development discussion with another NMC registrant, covering these reflections
- Obtain confirmation from a third party that they have met the requirements for revalidation.

The NMC recommends that nurses keep the evidence that they have met these requirements in a portfolio. It suggests that it may be helpful to structure the portfolio according to the themes in the Code.

search. This year there is a range of back-to-basics skills workshops where you can update your hands-on clinical skills.

- **PCRS-UK membership.** Maintaining your membership of PCRS-UK each year will ensure you are kept up to date with the latest respiratory resources, which you can access on the PCRS-UK website, and hear about the latest developments in respiratory policy and guidance through e-alerts.

References

1. The Code for nurses and midwives www.nmc.org.uk/standards/code
2. Post-acute COPD care bundle improvement worksheet <https://www.pcrs-uk.org/resource/Improvement-tools/post-acute-copd-care-bundle-improvement-worksheet>
3. PCRS-UK opinion sheets. <https://www.pcrs-uk.org/opinion-sheets>
4. PCRS-UK improvement worksheets <https://www.pcrs-uk.org/worksheets>
5. PCRS-UK annual conference <https://www.pcrs-uk.org/pcrs-uk-annual-conference>



British Thoracic Society

Winter Meeting 2015

WEDNESDAY 2 TO FRIDAY 4 DECEMBER

QEI Centre, London



Book now for Europe's biggest single Society Respiratory meeting. Participate in a wide variety of symposia and abstract sessions, featuring speakers from around the UK, Europe and USA:

- Mesothelioma: potential new treatment options and ongoing controversies
- Joint BTS/BALR symposium: epigenetics and lung disease
- New directions in TB diagnostics and therapy
- CF: current concepts and controversies
- Fifty years of COPD: where next on assessment and treatment?
- Joint BTS/BPRS symposium: infection in chronic suppurative airways disease
- Respiratory viral infections: learning from the past, treating the present, predicting the future
- BTS/BLF/BALR early career investigators prize symposium
- Joint BTS/BTOG symposium
- Joint BTS/BPRS symposium: global threats in paediatric lung disease
- Protease mediated tissue remodelling in chronic lung diseases
- Plenary scientific symposium
- COPD MAP – collaborative research providing insights into COPD
- Combating eosinophilic inflammation in asthma
- What's new in rare lung diseases?
- Antibiotic resistance
- Breathing life into MND: electricity, air and care
- ILD: what's on the horizon?
- Sensational developments in lung disease
- Occupational lung disease: the general chest clinic and beyond
- Interventional bronchoscopy for benign disease
- Pulmonary embolism: from acute to chronic

Also prestigious Guest Lectures: the Snell Memorial Lecture (Professor Paul Fine), The BTS Lecture (Professor Alvar Agusti) and the Moran Campbell Lecture (Professor Mike Morgan). Plus BTS Journal Clubs on Sleep, Asthma and CT Screening.

To see the full programme and to book your place, visit our website at: www.brit-thoracic.org.uk

Early bird discounts for bookings received before 12 October 2015

WORKING FOR HEALTHIER LUNGS



Policy Round-Up

Bronwen Thompson, *PCRS-UK Policy Advisor*

A summary of the latest developments in the UK health services, including any major new reports, guidelines and other documents relevant to primary care respiratory medicine

NICE busy with respiratory guidance

It seems to be a particularly busy period for respiratory work at NICE. Having done very little work on asthma for some time, NICE has been progressing two asthma guidelines in their work programme. Our Chair's perspective piece outlines the debates that the draft asthma diagnosis and monitoring guideline has prompted – and the article on FeNO elaborates on the potential role of this objective test. We await with interest final publication of this delayed guideline later this year.

The final scope of the Asthma management guideline has now been published so we know it will focus on pharmacological management of chronic asthma, review of pharmacological therapy and some aspects of non-pharmacological management of asthma (adherence, risk stratification, supported self-management and breathing exercises). Areas excluded from the guideline include comparison of inhaler devices, management of acute attacks, and biologics. We are pleased that the guideline development group includes 4 primary care professionals. The publication date for this guideline is expected to be June 2017.

Other respiratory work at NICE includes publication of a quality standard (QS) on tobacco harm reduction, and the development of quality standards for COPD and pneumonia for publication in January 2016. The tobacco harm reduction QS acknowledges that even if smokers don't quit completely, their health will benefit from reducing the amount of tobacco they use – whether this is by reducing the number of cigarettes they smoke, or whether they stop smoking but continue to use nicotine replacements. So the focus is on reducing the damage that tobacco can do rather than on quitting smoking. (see overleaf – To quit or not to quit)

A pneumonia quality standard is in its final stages. The draft QS which has been out to consultation proposed specific actions to be taken in primary care, based on a severity assessment, as well as guidance to secondary care for appropriate management in hospital. We are providing input to the consultation and look forward to seeing the final guidance.

COPD was one of the first conditions to have a quality standard (2011) and this is now being revised and updated. The draft QS revealed that they propose reducing the number of quality statements from 13 to 9, in order to focus attention on the areas of COPD care that are most in need of improvement. We contributed to the consultation in August following input from the members of our policy network aligned to COPD. For primary care the draft QS included the importance of post-bronchodilator spirometry, regular assessment of inhaler technique, assessment for eligibility for long term oxygen, and referral for pulmonary rehabilitation.

Asthma UK report on prescribing issues from National Review of Asthma Deaths

In June, Asthma UK published a report suggesting that, one year on from publication of the National Review of Asthma Deaths many asthma patients were still receiving medication deemed unsafe, and out of line with national guidelines. They analysed the notes of asthma patients in a number of practices across the country and extrapolated from the data gathered on these to the UK asthma population. They suggested that 22,000 people with asthma may be on long acting beta agonists (LABAs) or long acting muscarinic antagonists (LAMAs) without inhaled steroids (ICS) and as many as 100,000 people may be receiving over 12 short acting bronchodilators (SABAs) a year. PCRS-UK put these figures into perspective - 22,840 people prescribed a LABA or LAMA without ICS, and 106,742 people prescribed more than 12 short-acting bronchodilators a year represent only 0.4% and 2% respectively of the total 5.4 million people receiving treatment for asthma. The positive message for primary care is that the vast majority of people with asthma are therefore receiving appropriate medication.



Nevertheless, patient safety is paramount, and however small the numbers, any level of unsafe prescribing is a serious problem. PCRS-UK therefore continues to urge all practices to review their use of asthma medication by interrogating prescribing records on GP computer systems to ensure it is in line with the BTS/SIGN Asthma Guideline and safety guidance from the Medicines & Healthcare products Regulatory Agency (MHRA). The PCRS-UK practice improvement worksheet produced in response to the findings in NRAD outlines simple steps primary care health professionals can take to review and improve asthma care in their practices and provides appropriate supporting resources. We made this available to non-members following the report publication so that all primary care professionals could use it to improve care.

The Asthma UK report 'Patient safety failures in asthma: the scale of unsafe prescribing in the UK' <http://www.asthma.org.uk/patient-safety>

National asthma and COPD audits – an update

PCRS-UK was delighted when COPD was added to the list of national audits and we expected it to be only the second condition after diabetes to be included in primary care data in its national audit. However, the furore over care.data in 2014 heightened sensitivities over the collection of patient data from practices, so that the decision was finally taken in August that no primary care data – except that available through routine collection by QOF – will be collected as part of the national COPD audit. This was clearly disappointing news for the committee that had been working for several years to get this off the ground, and for patients and primary care professionals. Instead the COPD audit will focus only on organisation of care and outcomes in secondary care, and pulmonary rehabilitation. The exception is that as care.data does not apply in Wales, the primary care audit will proceed in Wales this autumn. (The National Audit programme does not cover Scotland and Northern Ireland.)

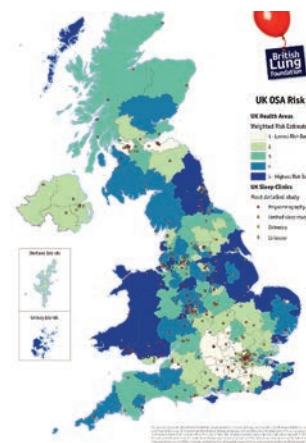
So while we were pleased when it was announced in May that asthma has been added to the national audit programme – once again, it looks as though the high level controversy over data confidentiality will prevent primary care data being collected. This is clearly disappointing when the vast majority of asthma care takes place in primary care and we are currently looking at whether there is anything PCRS-UK can do further to influence the situation. We hope that in due course the care.data issues will be resolved and data collection for audit in primary care in England can be resumed.

Focus on obstructive sleep apnoea

PCRS-UK have endorsed an excellent resource from the British Lung Foundation which is designed to bring together guidance from a range of sources to inform the planning and commissioning of services for obstructive sleep apnoea (OSA) – the OSA Commis-

sioning Toolkit. The BLF has been running an awareness campaign about OSA for several years and have brought the learning from the campaign together with clinical guidance to deliver a toolkit that will provide guidance to clinicians and managers who are developing a service for OSA. As well as 'Tips for GPs', there is a predicted risk map, which demonstrates which parts of the country are deemed to have the greatest population at risk of OSA. A handy calculator enables the prevalence of OSA in a region to be estimated by inputting local data, and the health economics around OSA are also included in the toolkit. We encourage you to look at how this toolkit could help with designing services locally in collaboration with commissioning colleagues.

Access the report at <https://www.blf.org.uk/Page/OSA-toolkit>



In brief – To quit or not to quit.... or try vaping?

There has been a lot of noise recently regarding tobacco smoking and vaping with e-cigarettes. The 'Smoking still kills' report, published by Action on Smoking and Health (ASH) in June, proposes new targets for a renewed national strategy to accelerate the decline in smoking prevalence over the next decade. This was followed by a review of the evidence on e-cigarettes from Public Health England in August, which reinforced the significantly reduced harm from using e-cigarettes compared to smoking tobacco. This was echoed by the NICE quality standard on reducing harm from tobacco, which promotes the use of any licensed nicotine replacement products for damage limitation. Debates still rage about whether e-cigarettes should be banned in public places alongside tobacco-containing cigarettes – Wales says yes, England says no – and medical organisations are divided. What is clear is that e-cigarettes are proving very popular – sales of other nicotine replacement products such as patches are falling – and sales of e-cigarettes have outstripped dental floss and canned soup. Watch this space!

Keep an eye out for our revised materials on stopping smoking as we will be keeping them up to date as EU legislation on tobacco and e-cigarettes comes into force in May 2016.

GETTING THE BASICS RIGHT

Fractional Exhaled Nitric Oxide testing



Fran Robinson talks to Carol Stonham a nurse practitioner, PCRS-UK nurse lead and vice chair of the PCRS-UK Education Committee, who has been carrying out FeNO testing for asthma for a number of years in her practice, the Minchinhampton Surgery, Gloucestershire. Carol says the test is so useful that now she would find diagnosing and monitoring asthma much more difficult without it.

What is a FeNO test?

Patients with allergic airway inflammation generally have higher than normal levels of nitric oxide (NO) in their exhaled breath. By measuring the concentration of NO in exhaled breath (fractional exhaled nitric oxide or FeNO), clinicians can identify inflammation in the lungs.

Why you need to know about FeNO testing

NICE consulted on draft guidance for the diagnosis and monitoring of asthma¹ in January this year in which FeNO tests were recommended to help confirm the diagnosis of asthma in adults and children.

The draft guidance suggested that asthma should no longer be diagnosed on clinical symptoms or spirometry alone, or by a trial of steroids. It recommended spirometry as a first line of investigation and that a FeNO test should be offered if a diagnosis of asthma is being considered in adults and young people over the age of five.

The draft guidance also suggested that clinicians should use FeNO testing to help manage asthma in patients who continue to have symptoms despite being prescribed inhaled corticosteroids. The draft guidance has sparked considerable debate (see Chair's perspective page 8). Very few people use FeNO test currently, so we spoke to one person who has, Carol Stonham, to find out about its potential value in practice

Three reasons to take a FeNO measurement:

1: To assist in diagnosis

The FeNO test can help to diagnose airway inflammation and also determine when respiratory symptoms are not due to asthma.

Carol says: *"The problem with spirometry is that if patients are asymptomatic on the day they come in for the test, which measures obstruction in the airways, the result will be normal. The likelihood is they will still have some underlying inflammation and this is where the FeNO test is useful because it measures inflammation."*

"Taking a good, accurate clinical history is always the basis of suspecting an asthma diagnosis. However quite often the history is not straightforward – you think the patient has probably got asthma but it's not definite. In my experience it's those intermediate probability cases – about 60% of patients – where a FeNO test is really useful."

"I see it as being part of a jigsaw puzzle, it is not used on its own but is used in addition to other diagnostic criteria".

2: To aid treatment

The FeNO measurement can help the clinician to determine the likelihood of steroid responsiveness and can guide stepwise changes in anti-inflammatory medication: step-down dosing, step-up dosing, or discontinuation of treatment.

If a patient on a low dose inhaled steroid becomes symptomatic and compliance and inhaler technique have been checked, a FeNO test can be used to indicate whether they need more anti-inflammatory therapy.

Likewise a FeNO measurement can help the clinician assess whether to step down treatment. Carol says: *"We don't step patients down as often as we should – we are notoriously bad at doing that. There are many reasons why we don't: we don't have the confidence"*

to do it; we don't know when to do it; we are not really sure how to do it; patients don't want to do it because they feel well - and those are all given, but not valid reasons. But high doses of medication are expensive and come with side effects for patients so we should step patients down when we can.

"If you have a patient whose asthma is under control: they have had at least three months when they have been really well, have not been using their rescue inhaler, have had no symptoms and are functioning well i.e. they are not reducing their activity and are not awake at night, then this could be a good time step them down.

"The beauty of the FeNO test is that it can help to reassure them that stepping down treatment is beneficial. You can do a test and give the patient a measurement then call them back a few weeks later and do another test to show them that the level of inflammation is not rising."

3: To identify poor adherence to treatment

The FeNO test can also help to determine whether patients are adhering to their prescribed corticosteroid treatment.

Carol explains: *"If a patient's FeNO level is rising and they are on a reasonable dose of steroid it's a good way of opening a conversation about inhaler technique or to ask them about how they are taking their medication. If they are not taking medication as prescribed, they will usually be happy to chat about it, especially when the raised FeNO level is suggesting something is amiss. Usually a four second break in the conversation is enough space for a patient to feel they need to 'confess' . . .*

"So if you have got a patient who isn't on board with their care you can use the FeNO test measurements to show that treatment can make a difference over a short period of time. Their FeNO measurement can drop from 68 to 32 in a matter of a couple of weeks, the patient starts feeling better and that helps them to grasp the concept that their treatment works. Patients also like it because they have a figure that they can see is improving."

How do you carry out a FeNO test?

A FeNO device is a hand held machine which requires a 10 second blow at 60 Litres a minute. It works like an alcohol breathalyser giving the results on the spot.

The patient first breathes out into the air to ensure any atmospheric nitric oxide (NO) is eliminated. They then seal their lips around the mouthpiece of the machine, take a full breath in and blow back out again into the machine. This then gives a clean measurement of NO in the exhaled breath. The patient is guided to breathe at the right flow rate by a computerised graphic. If the flow rate is not at the right level the machine will not give a measurement.

Carol says: *"There is a shorter test you can do for smaller children but I usually use the 10 second test for most patients over six years old."*

How much training is needed to use a FeNO device?

A short training is needed which is provided by the manufacturer either by coming in to the surgery or via online webinars. The practitioner needs to be shown how to use the machine and to be taught what the different FeNO measurements mean. For example tonsillitis or smoking can result in higher readings amongst other things.

What does it cost?

One of the FeNO machine manufacturers says the costs of the machine plus the consumables work out at £6 per test over 5 years or £450 per quarter. These prices are obviously dependent on the number of test done and are exclusive of VAT but provide a rough ballpark of the costs involved .

Carol believes that the FeNO test saves money by reducing unscheduled emergency GP appointments, A&E visits and hospitalisation and by ensuring clinicians put patients on the right medication first time and by improving compliance. Stepping down to appropriate doses of inhaled steroids also saves money.

References

1. Asthma: diagnosis and monitoring of asthma in adults, children and young people. NICE. January 2015 <http://www.nice.org.uk/guidance/indevelopment/gid-cgwave0640>



FeNO by NIOX®

GET STARTED WITH FeNO

NICE recommends
FeNO testing

Be sure
about
asthma

NICE recommends FeNO* testing:

- ◆ For a faster and more accurate asthma diagnosis^{1,2}
 - ◆ To help manage symptomatic asthma patients¹
 - ◆ To help detect non-adherence to treatment¹
- ...and because it is highly cost effective.^{1,2}

*Fractional Exhaled Nitric Oxide

1. NICE diagnostics guidance Measuring FeNO in Asthma [DG12] 2014: <http://www.nice.org.uk/guidance/dg12>

2. NICE Draft guidance Asthma Diagnosis and Monitoring 2015: <http://www.nice.org.uk/guidance/gid-cgwave0640/documents/asthma-diagnosis-and-monitoring-draft-guideline2>

NIOX products are sold and distributed
in the UK and Ireland by:



NIOX®

CHANGING THE FACE
OF ASTHMA CONTROL

Phone: +44 845 605 5521 Email: info@hc21.eu

The **only pMDI** licensed in COPD for patients who require an ICS/LABA combination inhaler¹

(FEV₁ < 50% predicted)



FOSTAIR[®]
pMDI Beclometasone
+ formoterol



Chiesi



FOSTAIR pMDI 100/6 (pressurised metered dose inhaler)
(Beclometasone dipropionate/formoterol fumarate dihydrate)
Please refer to the full Summary of Product Characteristics (SmPC)
before prescribing

Prescribing information

Presentation Each metered dose contains 100 micrograms (mcg) of beclometasone dipropionate (BDP) and 6mcg of formoterol fumarate dihydrate. **Indications** *Asthma:* Regular treatment of asthma in patients who are not adequately controlled on inhaled corticosteroids (ICS) and 'as needed' rapid acting beta₂-agonist or patients who are adequately controlled on both ICS and long-acting beta₂-agonists (LABA), where the use of an ICS/LABA combination is appropriate. *COPD:* Symptomatic treatment of patients with severe COPD (FEV₁ <50% predicted normal) and a history of repeated exacerbations, who have significant symptoms despite regular therapy with long-acting bronchodilators. **Dosage and administration** For inhalation in adult patients (≥18 years). BDP in Fostair is characterised by an extrafine particle size distribution which results in a more potent effect than formulations of BDP with a non-extrafine particle size distribution (100mcg of BDP extrafine in Fostair are equivalent to 250mcg of BDP in a non-extrafine formulation). *Asthma:* Fostair may be used as a maintenance therapy (with a separate rapid-acting bronchodilator as needed) or as a maintenance and reliever therapy (taken as a regular maintenance treatment and as needed in response to asthma symptoms). *Maintenance therapy:* 1-2 inhalations twice daily. The maximum daily dose is 4 inhalations. Patients should receive the lowest dose that effectively controls their symptoms. *Maintenance and reliever therapy:* 1 inhalation twice daily plus 1 additional inhalation as needed in response to symptoms. If symptoms persist after a few minutes, an additional inhalation is recommended. The maximum daily dose is 8 inhalations. *COPD:* 2 inhalations twice daily. Can be used with the AeroChamber Plus[®] spacer device. **Contraindications** Hypersensitivity to the active substances or to any of the excipients (HFA-134a, ethanol anhydrous, hydrochloric acid). **Warnings and precautions** Use with caution in patients with cardiac arrhythmias, aortic stenosis, hypertrophic obstructive cardiomyopathy, severe heart disease, occlusive vascular diseases, arterial hypertension,

aneurysm, thyrotoxicosis, diabetes mellitus, pheochromocytoma and untreated hypokalaemia. Caution should also be used when treating patients with known prolongation of the QTc interval (QTc > 0.44 seconds). Formoterol itself may induce QTc prolongation. Potentially serious hypokalaemia may result from beta₂-agonist therapy which may be potentiated by concomitant treatments and increase the risk of arrhythmias. Formoterol may cause a rise in blood glucose levels. As Fostair contains a corticosteroid, it should be administered with caution in patients with pulmonary tuberculosis or fungal/viral airway infections. Fostair treatment should not be stopped abruptly. Treatment should not be initiated during exacerbations or deteriorating asthma. Fostair treatment should be discontinued immediately if the patient experiences a paradoxical bronchospasm. **Systemic effects:** Systemic effects of ICS may occur, particularly at high doses for long periods, but are less likely than with oral steroids. These include Cushing's syndrome, Cushingoid features, adrenal suppression, decrease in bone mineral density, cataract and glaucoma and more rarely, a range of psychological or behavioural effects including psychomotor hyperactivity, sleep disorders, anxiety, depression and aggression. Prolonged treatment with high doses of inhaled corticosteroids may result in adrenal suppression and/or acute adrenal crisis. **Interactions** Beta-blockers should be avoided in asthmatic patients. Concomitant administration of other beta-adrenergic drugs may potentiate the adverse reactions of Fostair. Concomitant treatment with quinidine, disopyramide, procainamide, phenothiazines, antihistamines, monoamine oxidase inhibitors and tricyclic antidepressants can prolong the QTc interval and increase the risk of ventricular arrhythmias. L-dopa, L-thyroxine, oxytocin and alcohol can impair cardiac tolerance towards beta₂-sympathomimetics. Hypertensive reactions may occur following co-administration with monoamine oxidase inhibitors. Concomitant treatment with beta₂-agonists and xanthine derivatives, steroids or diuretics may potentiate hypokalaemia effects. Hypokalaemia may increase the likelihood of arrhythmias in patients receiving digitalis glycosides. **Fertility, pregnancy and lactation** Fostair should only be used during pregnancy or breast-feeding if the expected benefits outweigh the potential risks. **Effects on driving and operating machinery** Fostair is unlikely to have any effect on the ability to drive or operate machinery.

Side effects *Common:* pharyngitis, oral candidiasis, headache, dysphonia. *Uncommon:* influenza, oral fungal infection, oropharyngeal candidiasis, oesophageal candidiasis, vulvovaginal candidiasis, gastroenteritis, sinusitis, rhinitis, pneumonia, granulocytopenia, allergic dermatitis, hypokalaemia, hyperglycaemia, restlessness, tremor, dizziness, otosalginitis, palpitations, prolongation of QTc interval, electrocardiogram change, tachycardia, tachyarrhythmia, atrial fibrillation, hyperaemia, flushing, cough, productive cough, throat irritation, asthmatic crisis, diarrhoea, dry mouth, dyspepsia, dysphagia, burning sensation of the lips, nausea, dysgeusia, pruritus, rash, hyperhidrosis, urticaria, muscle spasms, myalgia, C-reactive protein increased, platelet count increased, free fatty acids increased, blood insulin increased, blood ketone body increased, blood cortisol decrease. *Rare:* ventricular extrasystoles, angina pectoris, paradoxical bronchospasm, angioedema, nephritis, blood pressure increased, blood pressure decreased. *Very rare:* thrombocytopenia, hypersensitivity reactions, including erythema, lips face, eyes and pharyngeal oedema, adrenal suppression, glaucoma, cataract, dyspnoea, exacerbation of asthma, growth retardation in children and adolescents, peripheral oedema, bone density decreased. *Unknown frequency:* psychomotor hyperactivity, sleep disorders, anxiety, depression, aggression, behavioural changes (predominantly in children) (Refer to SmPC for full list of side effects). **Legal category** POM **Packs and prices** £29.32 1x120 actuations. **Marketing authorisation number** PL 08829/0156. **Marketing authorisation holder** Chiesi Limited, 333 Styal Road, Manchester, M22 5LG **Date of preparation** June 2015

Adverse events should be reported. Reporting forms and information can be found at www.mhra.gov.uk/yellowcard. Adverse events should also be reported to Chiesi Limited. (address as above) Tel: 0161 488 5555.

Reference 1. MIMS. March-May 2015.

CHFOS2014.0934b. Date of preparation: July 2015.

Journal Round-Up

npj Primary Care Respiratory Medicine Key Summaries

npj | Primary Care
Respiratory Medicine

A selection of short summaries of original research articles published in *npj Primary Care Respiratory Medicine*. The articles featured have been selected by the Primary Care Respiratory Update editorial board as being the most relevant and useful to primary care respiratory clinical practice in the UK. You can download freely any articles of interest from the website <http://www.nature.com/npjpcrm/>

npj Primary Care Respiratory Medicine is the only fully indexed scientific journal devoted to the management of respiratory diseases in primary care. It is an international, online, open access journal and is part of the Nature Partner Journal series.

If you would like to be informed when a new paper is published by *npj Primary Care Respiratory Medicine* simply join the npj Primary Care Respiratory Medicine e-alert list to receive notification direct to your inbox. Visit www.nature.com/npjpcrm/ and click the link on the right titled E-alert.

** EDITOR'S CHOICE **

Patient-reported side effects, concerns and adherence to corticosteroid treatment for asthma, and comparison with physician estimates of side-effect prevalence: a UK-wide, cross-sectional study

Vanessa Cooper, Leanne Metcalf, Jenny Versnel, Jane Upton, Samantha Walker & Rob Horne. *npj Primary Care Respiratory Medicine* 25, Article number: 15026 (2015) doi:10.1038/npjpcrm.2015.26 Published online 09 July 2015

Asthma: Corticosteroids from patient and professional perspectives
Asthma patients in a UK-wide study reported side effects of corticosteroid treatment at a higher frequency than estimates made by clinicians. Professor Rob Horne of University College London led a questionnaire-based cross-sectional study in collaboration with Asthma UK and Education for Health. Questionnaires were completed by 1,524 patients taking inhaled corticosteroids, 233 taking oral corticosteroids and 244 health professionals. An unexpectedly large proportion of

people with asthma experienced side effects and had concerns about their treatment. For example, 46% of patients taking inhaled corticosteroids reported sore throat while health professionals estimated this would be 10%. Concerns about corticosteroids compromised adherence to treatment. Male participants and those who were older reported fewer side effects and had fewer concerns about their treatment. These findings have implications for the design of interventions to optimize asthma control.

HELPing older people with very severe chronic obstructive pulmonary disease (HELP-COPD): mixed-method feasibility pilot randomised controlled trial of a novel intervention

Susan Buckingham, Marilyn Kendall, Susie Ferguson, William MacNee, Aziz Sheikh, Patrick White, Allison Worth, Kirsty Boyd, Scott A Murray & Hilary Pinnock. *npj Primary Care Respiratory Medicine* 25, Article number: 15020 (2015) doi:10.1038/npjpcrm.2015.20 Published online 16 April 2015

Severe lung disease: Exploring initiatives for better care

A trial to improve care for patients with severe lung disease concludes that holistic treatment throughout the disease may be beneficial. Severe chronic obstructive pulmonary disease (COPD) has a massive physical and psychological impact on patients' lives, prompting calls for palliative care provision. In a pilot trial of elderly patients with very severe COPD, Hilary Pinnock and co-workers at the University of Edinburgh, Scotland, investigated the effectiveness of supportive home visits by a specialist respiratory nurse, who assessed patients' wellbeing and needs, and agreed upon a plan of

action with them. Qualitative interviews with patients and professionals indicated that the holistic assessment was well received and appreciated. However, few actions resulted from the nurse's intervention, partly due to overlaps with existing care. The team believes integrating holistic assessment into routine COPD treatment may be more appropriate.

Effectiveness of community-based integrated care in frail COPD patients: a randomised controlled trial

Carme Hernández, Albert Alonso, Judith Garcia-Aymerich, Ignasi Serra, Dolors Martí, Robert Rodríguez-Roisin, Georgia Narsavage, Maria Carmen Gomez, Josep Roca & NEXES consortium. *npj Primary Care Respiratory Medicine* 25, Article number: 15022 (2015) doi:10.1038/npjpcrm.2015.22 Published online 09 April 2015

Chronic obstructive pulmonary disease: Investigating integrated care
Community-based integrated care (IC) for chronic obstructive pulmonary disease (COPD) improves outcomes but does not reduce hospitalization. Spanish researchers led by Carme Hernández at the

University of Barcelona conducted a randomized controlled trial to explore the effectiveness of a community-based IC service in preventing hospitalizations and emergency department visits. The service consisted of assessment and education, individualized care plans, call center access and coordination between different levels of care. Seventy-six stable frail COPD patients received the IC intervention for 12 months, while 84 received usual care. Hospital admissions, emergency department visits and mortality were then monitored for six years after return to normal care. The intervention improved clinical outcomes and decreased emergency department visits, but did not reduce hospital admissions. Patient risk analysis and workforce preparation were identified as key requirements for developing IC services.

Initial step-up treatment changes in asthmatic children already prescribed inhaled corticosteroids: a historical cohort study

Steve W Turner, Kathryn Richardson, Annie Burden, Mike Thomas, Clare Murray & David Price. *npj Primary Care Respiratory Medicine* 25, Article number: 15041 (2015) doi:10.1038/npjpcrm.2015.41 Published online 11 June 2015

Asthma: Stepping up treatment in children

Clinicians in the UK frequently opt to deviate from the guidelines for increasing treatment choices in children with asthma. When standard doses of inhaled corticosteroids (ICS) fail to control symptoms in children older than four years, guidelines recommend adding long-acting beta agonist (LABA), with other treatment options considered if symptoms persist. Stephen Turner at the University of Aberdeen and co-workers across the UK conducted a historical cohort study of treatment "step-up" choices in 10,793 children aged between 5 and 12. An increased dose of ICS was prescribed for 58% of the children. LABA was introduced for only 32%, combined in some cases with either a change of inhaler type or prescribing leukotriene receptor antagonist. Deviation from treatment guidelines appears to be an active decision based on patient characteristics including obesity, antibiotic therapy and rhinitis.

Prevalence of asthma-COPD overlap syndrome among primary care asthmatics with a smoking history: a cross-sectional study

Toni Kiljander, Timo Helin, Kari Venho, Antero Jaakkola & Lauri Lehtimäki. *npj Primary Care Respiratory Medicine* 25, Article number: 15047 (2015) doi:10.1038/npjpcrm.2015.47 Published online 16 July 2015

Asthma: Overlap with chronic obstructive pulmonary disease

Chronic obstructive pulmonary disease (COPD) affects many older asthmatic patients with a history of smoking but no previous COPD diagnosis. Finnish researchers led by Toni Kiljander of Terveystalo Hospital, Turku, investigated the prevalence of asthma-COPD overlap syndrome (ACOS) in a cross-sectional study of 190 primary care asthma patients. Spirometry studies and questionnaires revealed that 27.4% of the patients had ACOS. The best predictors of ACOS were age of 60 years or more and at least 20 pack-years of smoking history, each pack-year equating to 20 cigarettes per day for one year. Although ACOS appears to be clinically highly significant, little is known about its treatment as affected patients are typically excluded from therapy trials for asthma or COPD. More attention should be given to the possibility of ACOS in elderly asthmatics with a smoking history.

Patient perceptions of severe COPD and transitions towards death: a qualitative study identifying milestones and developing key opportunities

Amanda Landers, Rachel Wiseman, Suzanne Pitama & Lutz Beekert. *npj Primary Care Respiratory Medicine* 25, Article number: 15043 (2015) doi:10.1038/npjpcrm.2015.43 Published online 09 July 2015

Chronic lung disease: Identifying need for the palliative approach Patient understanding and acknowledgement of advancing lung disease may be improved by identifying transition points in health deterioration. Chronic obstructive pulmonary disease (COPD) is a progressive condition leading to disabling shortness of breath, severe lifestyle disruption and death. However, many patients describe COPD as a way of life rather than a life-threatening illness, making identification of palliative care needs difficult. Amanda Landers at Nurse Maude Hospice, Christchurch, with scientists from across the region interviewed 15 patients to explore their perceptions of severe COPD. The team identified six common transition points described by the patients. The transitions include mourning the loss of recreational hobbies to the need for long-term oxygen therapy and worries about self-care. Discussing these pivotal transitions may help stimulate patients and health service professionals to change focus and plan for the future.

'I still don't know diddly': a longitudinal qualitative study of patients' knowledge and distress while undergoing evaluation of incidental pulmonary nodules

Donald R Sullivan, Sara E Golden, Linda Ganzini, Lissi Hansen & Christopher G Slatore. *npj Primary Care Respiratory Medicine* 25, Article number: 15028 (2015) doi:10.1038/npjpcrm.2015.28 Published online 16 April 2015

Lung screening: Improvements in patient communication essential Renewed efforts are needed to alleviate patient stress and keep them informed following the diagnosis of small growths on the lungs. The discovery of so-called 'pulmonary nodules' during routine chest screening can cause considerable distress to patients. The nodules, though often benign, require monitoring via follow-up scans, leading to a period of uncertainty which potentially affects patients' well-being. In a longitudinal study involving interviews with 17 patients with newly diagnosed nodules, Donald Sullivan at Oregon Health and Science University, USA, and co-workers found that stress was often exacerbated by ineffective communication from medical staff. This led to patient misunderstandings about lung cancer risk, for example, which were compounded by the complex information in letters they received. Patients much preferred to speak to their primary care providers. The researchers call for improved staff training and education resources.

Exploring the impact of chronic obstructive pulmonary disease (COPD) on diabetes control in diabetes patients: a prospective observational study in general practice

Hilde D Luijckx, Wim JC de Grauw, Jacobus HJ Bor, Chris van Weel, Antoine LM Lagro-Janssen, Marion CJ Biermans & Tjard R Schermer. *npj Primary Care Respiratory Medicine* 25, Article number: 15032 (2015) doi:10.1038/npjpcrm.2015.32 Published online 23 April 2015

Chronic obstructive pulmonary disease: Effects on control of diabetes Further research is required to understand how the presence of lung disease may affect the treatment of type II diabetes in patients. People suffering from chronic obstructive pulmonary disease

(COPD) often have a co-existing disease, making both conditions more difficult to treat. To determine the impact of COPD on control of diabetes, Hilde Luijckx and co-workers at Radboud University Medical Center in The Netherlands conducted a longitudinal study of 610 primary care patients with diabetes, 63 of whom also had COPD. There was no significant association between glycated haemoglobin levels and COPD. However, the researchers found that systolic blood pressure was significantly associated with COPD patients' socioeconomic status (SES), gradually increasing in middle and high SES groups, and decreasing in low SES groups. High body mass index increased systolic blood pressure in diabetes patients without COPD.

Preference, satisfaction and critical errors with Genuair and Breezhaler inhalers in patients with COPD: a randomised, cross-over, multicentre study

Sergi Pascual, Jan Feimer, Anthony De Soyza, Jaume Sauleda Roig, John Haughney, Laura Padullés, Beatriz Seoane, Ludmyla Rebeda, Anna Ribera & Henry Chrystyn. *npj Primary Care Respiratory Medicine* 25, Article number: 15018 (2015) doi:10.1038/npjpcrm.2015.18 Published online 30 April 2015

Chronic obstructive pulmonary disease: Patients prefer Genuair inhaler over Breezhaler

More patients preferred the multi-dose Genuair than the single-dose Breezhaler inhaler in a randomised, cross-over, multicentre study. An international research team led by Henry Chrystyn of the University of Huddersfield, UK, enrolled 128 patients with chronic obstructive pulmonary disease and moderate-to-severe airflow obstruction in a two-week study involving daily inhaler use. The inhalers contained placebo, while patients continued with their

normal medication. Of the 110 patients who indicated a preference, 72.7% preferred the Genuair inhaler over the Breezhaler. Patients were more willing to continue using Genuair than Breezhaler. These differences were statistically significant. There was no difference in the number of patients who made critical errors of technique with each inhaler, and the overall incidence of errors was low. The authors recommend further research to compare Genuair with other inhalers over a longer time period.

Predictors of poor-quality spirometry in two cohorts of older adults in Russia and Belgium: a cross-sectional study

Eralda Turkeshi, Dmitry Zelenukha, Bert Vaes, Elena Andreeva, Elena Frolova & Jean-Marie Degryse. *npj Primary Care Respiratory Medicine* 25, Article number: 15048 (2015) doi:10.1038/npjpcrm.2015.48 Published online 23 July 2015

Respiratory disease: Effective lung function tests in elderly patients
A test used to assess how well a person's lungs are working should be used more often in elderly patients, Belgian researchers say. Eralda Turkeshi from the Catholic University of Louvain in Brussels and colleagues note that previous studies had found that elderly patients with cognitive impairment had difficulty performing a lung function blow test called 'spirometry'. However, their study of two groups of elderly patients in Belgium and Russia found that tests in over three-quarters of these patients met acceptable clinical standards after adjusting for gender, age and level of education. These tests should be used in older adults without restrictions based on mental or physical functioning, they concluded. Spirometry is a valuable tool for the diagnosis and management of respiratory diseases as well as for assessing overall health, they suggested.

Best of the rest

These reviews were prepared by Dr Basil Penney and published by Doctors.net.uk Journal Watch. They have been selected and edited for inclusion into *Primary Care Respiratory Update* by editor Dr Hilary Pinnock.



The Doctors.net.uk Journal Watch service covers other specialities as well as respiratory medicine. Doctors.net.uk is the largest network of GMC-registered doctors in the UK. To find out more about membership visit <http://www.doctors.net.uk>

Abbreviations used in these reviews are:		CVS	Cardiovascular System	LLN	Lower Limit of Normal
\$	Dollars	EBA	Economic Burden of Asthma	Mg/mL	milligrams per millilitre
ACOS	Asthma-COPD overlap syndrome	ERS	European Respiratory Society	NHS	National Health Service
ACT	Asthma Control Test	FENO	Fractional Exhaled Nitric Oxide	NICE	National Institute for Health and Care Excellence
AHR	Airway Hyper-responsiveness	FEV1	Forced expiratory volume in 1 second	OR	Odds ratio
ATS	American Thoracic Society	FVC	Forced vital capacity	PC	Provocative concentration of methacholine
BHR	Bronchial hyper-responsiveness	GP	General Practitioner	POCT	Point of Care Test
BMI	Body Mass Index	GLI	Global Lung Initiative	QoL	Quality of Life
C-ACT	Childhood Asthma Control Test	H	Hours	RTI	Respiratory Tract Infection
CI	Confidence Interval	IC	Integrated care	SD	Standard Deviation
CNS	Central Nervous System	ICS	Inhaled corticosteroid	SES	Socio-economic status
COPD	Chronic Obstructive Pulmonary Disease	Kg	Kilogram	UK	United Kingdom
CRP	C-Reactive Protein	kg/m ²	Kilogram per metre squared	USA	United States of America
		LABA	Long-acting beta-agonist		

**** EDITOR'S CHOICE ****

Oseltamivir treatment for influenza in adults: a meta-analysis of randomised controlled trials

THE LANCET

Joanna Dobson, Richard J Whitley, Stuart Pocock, Arnold S Monto. *Lancet* 2015; 385: 1729–37; [http://dx.doi.org/10.1016/S0140-6736\(14\)62449-1](http://dx.doi.org/10.1016/S0140-6736(14)62449-1)

Oseltamivir selectively blocks the enzymatic activity of all influenza viruses, making it useful in prophylaxis and treatment for both seasonal and pandemic disease. Concerns exist however about its efficacy and adverse effects and whether these outweigh the benefits.

This group conducted a meta-analysis of all available randomised treatment trials of oseltamivir in adults. They used individual patient data and included both published and unpublished trials thereby overcoming previous concerns regarding potential publication bias. They analysed participants who were identified as influenza-infected (intention-to-treat infected population), all treated participants (intention-to-treat population) and safety populations.

Data were included from nine trials (4328 patients). In the intention-to-treat infected population, they noted a 21% shorter time to alleviation of all symptoms for oseltamivir versus placebo recipients. The median times to alleviation were 97.5 h for oseltamivir and 122.7 h for placebo groups (difference –25.2 h,

95% CI –36.2 to –16.0). For the intention-to-treat population, the estimated treatment effect was attenuated but remained highly significant (median difference –17.8 h).

In the intention-to-treat infected population, they noted fewer lower respiratory tract complications requiring antibiotics more than 48 h after randomisation (4.9% oseltamivir vs 8.7% placebo, risk difference –3.8%, 95% CI –5.0 to –2.2) and also fewer admissions to hospital for any cause (0.6% oseltamivir, 1.7% placebo, risk difference –1.1%, 95% CI –1.4 to –0.3).

Regarding safety, oseltamivir increased the risk of nausea (9.9% oseltamivir vs 6.2% placebo, risk difference 3.7%, 95% CI 1.8–6.1) and vomiting (8.0% oseltamivir vs 3.3% placebo, risk difference 4.7%, 95% CI 2.7–7.3). They recorded no effect on neurological or psychiatric disorders or serious adverse events.

These findings show that oseltamivir in adults with influenza accelerates time to clinical symptom alleviation, reduces risk of lower respiratory tract complications, and admittance to hospital, but increases the occurrence of nausea and vomiting.

Effects of weight loss on airway responsiveness in obese adults with asthma



Smita Pakhale, Justine Baron, Robert Dent, Katherine Vandemheen, Shawn D. Aaron. *CHEST* 2015;147(6): 1582–1590; <http://dx.doi.org/10.1378/chest.14-3105>

Research suggests that the incidence of asthma is 1.47 times greater in obese people than non-obese people and that a three-unit increase in BMI is associated with a 35% increase in the risk of asthma. Few studies have examined whether weight loss in obese people with asthma improves asthma outcomes and of these few rely on appropriate physiological tests for asthma to determine study eligibility (ie, airway hyper-responsiveness (AHR), reversibility of airway obstruction).

This Canadian, prospective, controlled, parallel-group study, followed 22 obese participants with asthma aged 18 to 75 years with a BMI ≥ 32.5 kg/m² and AHR (provocative concentration of methacholine causing a 20% fall in FEV₁ [PC 20] < 16 mg/mL). Sixteen participants followed a behavioural weight reduction program for 3 months, and six served as control subjects.

The primary outcome was change in AHR over 3 months. Changes in lung function, asthma control, and quality of life were secondary outcomes.

At study entry, participant mean (SD) age was 44 (SD 9) years, 95% were women, and mean BMI was 45.7 (SD 9.2) kg/m². After 3 months, mean weight loss was 16.5 (SD 9.9) kg in the intervention group, and the control group had a mean weight gain of 0.6 (SD 2.6) kg.

There were significant improvements in PC 20 (P=0.009), FEV₁ (P=0.009), FVC (P=0.010), asthma control (P < 0.001), and asthma

quality of life (P=0.003) in the intervention group, but these parameters remained unchanged in the control group. Physical activity levels also increased significantly in the intervention group but not in the control group.

Weight loss in obese adults with asthma can improve asthma severity, AHR, asthma control, lung function, and quality of life. These findings support the need to actively pursue healthy weight-loss measures in this population.

Influenza vaccination for NHS staff: attitudes and uptake

BMJ Open Respiratory Research

Dinesh Shrikrishna, Siân Williams, Louise Restrict and Nicholas S Hopkinson. *BMJ Open Resp Res* 2015;2:e000079. <http://dx.doi.org/10.1136/bmjresp-2015-000079>

Annual vaccination against influenza (flu) is recommended for all UK NHS staff to reduce the risk of contracting the virus and transmitting it to patients. However, vaccination uptake in England for the 2013–2014 was only 54.8% for healthcare workers with direct patient contact.

The aim of this study was to investigate staff attitudes to flu vaccination to see how this may influence their decision to be vaccinated. An online survey was sent to staff members across 6 NHS trusts, asking if staff had been vaccinated in the preceding flu season (2013–2014); the survey included questions about beliefs and attitudes to the vaccination, scored on a 5-point Likert scale.

3059 NHS staff members responded to the survey. 68% reported being vaccinated in the preceding year. Doctors were the most likely group to have been vaccinated (doctors 72%, nurses 65%, clerical 63%, other staff members 68%).

Using a stepwise regression model, the survey response retained as a positive predictor of having been vaccinated was 'people working in healthcare should have the flu vaccination every year' ($p < 0.001$), and the responses retained as negative predictors were 'the flu vaccination will make me unwell' ($p < 0.001$) and 'the flu vaccination was too much trouble for me' ($p < 0.001$). Doctors had a greater tendency to disagree with the statement that 'the flu vaccination will make me unwell' compared to other staff members.

Addressing NHS staff beliefs around the need for vaccination, and removing practical barriers to vaccination, may help increase uptake. Alleviating the concerns of particular staff groups regarding adverse effects of the vaccine may also be of benefit in improving uptake.

Interaction effect of psychological distress and asthma control on productivity loss?

Grégory Moullec, J. Mark FitzGerald, Roxanne Rousseau, Wenjia Chen, Mohsen Sadatsafavi, the Economic Burden of Asthma (EBA) study team. *Eur Respir J* 2015;45:1557–1565
<http://dx.doi.org/10.1183/09031936.00141614>

Little is known about the potential synergistic effect of comorbid psychological distress and uncontrolled asthma on productivity loss.

This Canadian population-based study [conducted as part of a larger 1-year longitudinal study the Economic Burden of Asthma (EBA) study] aimed to narrow this evidence gap by quantifying productivity loss, in terms of presenteeism and absenteeism, as a function of level of asthma control and presence of psychological distress, in employees with asthma.

300 adults with asthma were prospectively recruited between Dec 2010 and Aug 2012. Psychological distress and productivity loss due to absenteeism and presenteeism were measured using validated instruments, and asthma control was ascertained using the 2010 Global Initiative for Asthma management strategy. The authors used two-part regression models to study the contribution of uncontrolled asthma and psychological distress to productivity loss.

Compared with reference group (controlled asthma + no psychological distress), people with uncontrolled asthma but no psychological distress had a weekly productivity loss of Canadian \$286 (95%CI \$276–297). People with psychological distress had a weekly productivity loss of Canadian \$465 (\$445–485) even if their asthma was well controlled. Having uncontrolled asthma made no further difference to the productivity loss Canadian \$449 (437–462).

In patients without psychological distress, uncontrolled asthma was associated with an increased productivity loss, but made no further difference in the presence of psychological distress. This finding can be explained by the fact that the contribution of psychological distress to productivity loss is so large that there is no room for synergy with asthma control.

Future studies should assess the impact of interventions that modify psychological distress in patients with asthma.

Monitoring strategies in children with asthma: a randomised controlled trial

Thorax

Sandra Voorend-van Bergen, Anja A Vaessen-Verberne, Hein J Brackel, Anneke M Landstra, Norbert J van den Berg, Wim C Hop, Johan C de Jongste, Peter J Merkus, Mariëlle W Pijnenburg. *Thorax* 2015;70:543–550. <http://dx.doi.org/10.1136/thoraxjnl-2014-206161>

While monitoring of asthma control in children is recommended the best monitoring strategy is not known.

The 'Better Asthma Treatment: Monitoring with ACT and Nitric oxide' study randomised controlled, partly blinded, parallel group multicentre trial studied two monitoring strategies for their ability to improve asthma outcomes in comparison with standard care: web-based monthly monitoring with the (Childhood) Asthma Control Test (ACT or C-ACT) and 4-monthly monitoring of FeNO.

280 children aged 4–18 years (mean age 10.4 years, 66% boys) with a diagnosis of asthma, and treated in one of seven hospitals were randomised to one of the three groups. In the web group, treatment was adapted according to ACT obtained via a website at 1-month intervals; in the FeNO group according to ACT and FeNO, and in the standard care group according to the ACT at 4-monthly visits. The primary endpoint was the change from baseline in the proportion of symptom-free days.

After 1 year, neither of the novel strategies had improved the number of symptom-free days more than standard care. However, monthly web-based ACTs resulted in a clinically relevant decrease of ICS dose, while maintaining asthma control. FeNO monitoring modestly improved asthma control selectively in children aged <12 years without the need for higher ICS doses.

The children were all under hospital care and almost half of all eligible children refused to participate in the study, which may limit the conclusions due to selection bias.

Neither of the two novel monitoring approaches improved asthma control, though seemed to achieve control with reduced doses of ICS. Future studies should test the combined strategy of web-based monitoring in combination with FeNO measurements during clinic visits.

Narrative review of primary care point-of-care testing (POCT) and antibacterial use in respiratory tract infection (RTI)

BMJ Open Respiratory Research

Jonathan Cooke, Christopher Butler, Rogier Hopstaken, Matthew Scott Dryden, Cliodna McNulty, Simon Hurding, Michael Moore and David Martin Livermore. *BMJ Open Resp Res* 2015;2:e000086. <http://dx.doi.org/10.1136/bmjresp-2015-000086>

Antimicrobial resistance is a global healthcare and economic problem. There is a need to avoid unnecessary prescriptions many of which are for respiratory tract infections (RTIs) in the community. Studies suggest little benefit is achieved from the prescription of antibiotics, except in elderly patients at high risk of pneumonia. Potentially, primary care prescribers can use point-of-care testing to inform their management of disease.

This narrative review of the literature was undertaken to ascertain the value of C reactive protein (CRP) and procalcitonin measurements to guide antibacterial prescribing in adult patients presenting

to GP practices with symptoms of respiratory tract infection (RTI). Studies included randomised controlled trials, controlled before and after studies, cohort studies and economic evaluations.

Many studies demonstrated that the use of CRP tests in patients presenting with RTI symptoms reduces antibiotic prescribing by 23.3% to 36.16%. Procalcitonin is not currently available as a POCT but has shown value in patients admitted to hospital with RTI. GPs and patients report good acceptability for a CRP point-of-care testing and economic evaluations show cost-effectiveness of CRP over existing RTI management in primary care. NICE pneumonia guidelines now make recommendations for use of CRP in assisting in diagnosis.

Point-of-care tests increase diagnostic precision for GPs in the better management of patients with RTI. CRP point-of-care testing can improve targeting of antibacterial prescribing by GPs and contribute to national antimicrobial resistance strategies. Health services need to develop ways to ensure funding is transferred in order for point-of-care testing to be implemented.

Airflow limitation by the Global Lungs Initiative equations in a cohort of very old adults

Eralda Turkeshi, Bert Vaes, Elena Andreeva, Catharina Matheï, Wim Adriaenssen, Gijs Van Pottelbergh and Jean-Marie Degryse. *Eur Respir J* 2015;46:123–132 <http://dx.doi.org/10.1183/09031936.00217214>

Correct identification of airflow limitation in very old adults is important as it may reduce inappropriate COPD diagnosis and use of COPD medications, along with their side-effects.

The cut-off for forced expiratory volume in 1 second (FEV₁)/forced vital capacity (FVC) defining airflow limitation for chronic obstructive pulmonary disease (COPD) is still contested. ATS/ERS support the use of the lower limit of normal (LLN) at the 5th percentile of the frequency distribution of values of a reference "healthy" never-smoker population of equivalent age and sex. The fixed cut-off approach leads to over diagnosis of COPD in older adults, while the LLN cut-off approach is dependent on age-specific reference values. These have been lacking for adults aged >80 years or have been extrapolated from younger populations, until recently when the Global Lungs Initiative (GLI) all-age reference equations for different ethnic groups and populations aged 3–95 years were made available.

This Belgian population-based prospective cohort study assessed airflow limitation prevalence by the lower limit of normal (LLN) of Global Lungs Initiative (GLI) 2012 reference values and its predictive ability for all-cause mortality and hospitalisation in 411 very old adults (aged ≥80 years) compared with the fixed cut-off.

9.2% had airflow limitation by GLI-LLN and 27% by fixed cut-off, without good agreement (kappa coefficient ≤0.40) with GP-reported COPD (9%). Only airflow limitation by GLI-LLN was independently associated with mortality (adjusted hazard ratio 2.10, 95%CI 1.30 to 3.38). FEV₁/FVC <0.70 but ≥GLI-LLN (17.8%) had no significantly higher risk for mortality or hospitalisation.

In a cohort of very old adults, airflow limitation by GLI-LLN has lower prevalence than by fixed cut-off, independently predicts all-cause mortality and does not miss individuals with significantly higher all-cause mortality and hospitalisation.

Effects of quitting cannabis on respiratory symptoms

Robert J. Hancox, Hayden H. Shin, Andrew R. Gray, Richie Poulton and Malcolm R. Sears. *Eur Respir J* 2015;46:80–87 <http://dx.doi.org/10.1183/09031936.00228914>

Moderate levels of cannabis use are associated with proximal airway inflammation and symptoms of bronchitis. These associations persist after adjusting for tobacco smoking and also occur in those who only smoke cannabis, indicating that cannabis can cause bronchitis independently of tobacco. Little is known about the persistence of symptoms after stopping cannabis use.

This group from New Zealand studied the effect of quitting cannabis use in the Dunedin Multidisciplinary Health and Development Study, a population-based birth cohort of 1,037 people (52% male). Participants have been followed throughout childhood and into adulthood. Follow-up rates have been high with 95% of the surviving cohort continuing to participate in the most recent assessment at age 38 years. A previous analysis of this study found that many cannabis users had symptoms of bronchitis at the age of 21 years.

Participants were asked about cannabis and tobacco use at ages 18, 21, 26, 32 and 38 years. Symptoms of morning cough, sputum production, wheeze, dyspnoea on exertion and asthma diagnoses were ascertained at the same ages. Frequent cannabis use was defined as ≥52 occasions over the previous year.

Associations between frequent cannabis use and respiratory symptoms were analysed using generalized estimating equations with adjustments for tobacco smoking, asthma, sex and age.

Frequent cannabis use was associated with morning cough (OR 1.97, p<0.001), sputum production (OR 2.31, p<0.001) and wheeze (OR 1.55, p<0.001). Reducing or quitting cannabis use was associated with reductions in the prevalence of cough, sputum and wheeze to levels similar to nonusers.

Frequent cannabis use is associated with symptoms of cough, sputum production and wheeze independently of tobacco smoking. These symptoms tend to improve in people who quit indicating that the airway inflammation caused by cannabis may be largely reversible.

Gait patterns in COPD: the Rotterdam Study

Lies Lahousse, Vincentius J.A. Verlinden, Jos N. van der Geest, Guy F. Joos, Albert Hofman, Bruno H.C. Stricker, Guy G. Brusselle and M. Arfan Ikram. *Eur Respir J* 2015; 46: 88–95 <http://dx.doi.org/10.1183/09031936.00213214>

In patients with COPD, muscle weakness precedes the development of functional limitations, which can result in an increased risk of falling. Even without injuries, a fall can negatively impact on quality of life through fear of falling, which often inhibits performance of activities. Although causes of falls are often multifactorial, the majority occur during walking and gait analysis may be the best tool to predict the risk of falls. Gait is affected by various organ systems, such as the CNS, CVS and musculoskeletal system, and poor gait is a strong risk factor of death. COPD may affect gait through any of these systems.

This large (n=1,094), population-based cohort Rotterdam Study (age ≥ 55 years), investigated associations of COPD with various gait domains (including Rhythm and Pace) and explored a potential link with falling.

Those with COPD (n=196) exhibited worse Rhythm compared with those with normal lung function (n=898), independent of age, sex, height, education, smoking or analgesic use, especially when dyspnoea and severe airflow limitation or frequent exacerbations (Global Initiative for Chronic Obstructive Lung Disease group D: -0.83 SD, 95%CI -1.25 to -0.41 SD) were present.

A lower FEV₁ was associated with worse Rhythm and Pace, including lower cadence and gait velocity, respectively. Importantly, fallers with COPD had significantly worse Rhythm than non-fallers with COPD.

Given the cross-sectional analysis and the retrospective interrogation of falls, causality cannot be inferred: -the worse Rhythm in COPD patients may lead to more falls, or falls in COPD patients may aggravate anxiety and unsecured gait, leading to worse Rhythm.

Further research should investigate the underlying mechanisms of these associations, to enable development of proper intervention strategies to prevent falling in patients with COPD.

Large observer variation of clinical assessment of dyspnoeic wheezing children

Archives of
Disease in Childhood

Jolita Bekhof, Roelien Reimink, Ine-Marije Bartels, Hendriekje Eggink, Paul L P Brand. *Arch Dis Child* 2015;100:649–653. <http://dx.doi.org/10.1136/archdischild-2014-307143>

Evaluating the severity of dyspnoea in children is important in clinical decision-making and evaluation of treatment. The usefulness of the clinical assessment of dyspnoea is strongly determined by its reliability and is often performed by different professionals. The aim of this observational study from the Netherlands was to determine the intraobserver and interobserver variation of common clinical findings in children with acute severe dyspnoea and wheeze.

27 acutely wheezing children (aged 3 months–7 years) were recorded in the emergency department on video before and after treatment with inhaled bronchodilators. These video recordings were independently assessed by nine observers scoring wheeze, prolonged expiratory phase, retractions, nasal flaring and a general assessment of dyspnoea on a Likert scale (0–10). Assessment was repeated after 2 weeks to evaluate intra-observer variation.

Intra-observer variation was modest, and inter-observer variation was large for most clinical findings in children with dyspnoea. The measurement error induced by this variation was too large to distinguish potentially clinically relevant changes in dyspnoea after treatment in two-thirds of observations.

The poor inter-observer reliability of clinical dyspnoea assessment in children limits its usefulness in clinical practice and research, and highlights the need to use more objective measurements in these patients.

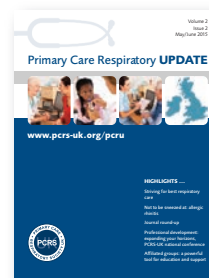
PCRS-UK News Round-Up

TELL US WHAT YOU THINK...

We are keen to hear from you what you think of the *Primary Care Respiratory Update*. Is it useful? Do you like the centre-fold pull out charts that we produce from time to time? Do you like having summaries from other respiratory related journals?

Enter this short (it takes less than five minutes) survey to tell us your views
Visit https://www.surveymonkey.com/r/PCRU_Review to take part

Make sure you submit your feedback before 30th October 2015



NEW PCRS-UK EXECUTIVE MEMBER



PCRS-UK is delighted to announce the election of Oonagh Potts to the PCRS-UK Executive. Oonagh is a nurse practitioner and lead nurse in primary care.

Oonagh is the group leader for the PCRS-UK affiliated group, Fylde Respiratory Forum, and she is a regional trainer for Education for Health.

Commiserations to those who were unsuccessful in the election. However, we look forward to their input through other committees within PCRS-UK.

NEW PCRS-UK REGIONAL LEADS



Jackie Abbey and Basil Penney have recently been appointed regional leads for PCRS-UK for Wales and the North East respectively.

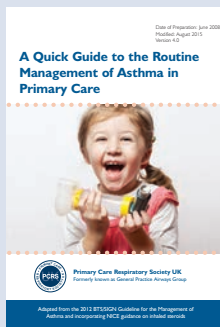
You can see who your regional lead is at <https://www.pcrs-uk.org/pcrs-uk-regional-leads> and contact them directly via our Members Directory. Simply log in to the PCRS-UK website and then go to <http://www.pcrs-uk.org/directory>.

Regional leads are shown with the symbol



LATEST RESOURCES

PCRS-UK Quick Guide to the Diagnosis and Management of Asthma



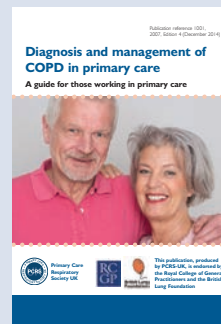
PCRS-UK is delighted to announce the revision of the PCRS-UK Quick Guide to the Diagnosis and Management of Asthma. This 'Quick Guide' to the routine management of asthma in primary care is based on the British Thoracic Society (BTS) and Scottish Intercollegiate Guideline Network (SIGN) British Guideline on the Management of Asthma, May 2008, revised edition published October 2014 <https://www.brit-thoracic.org.uk/guidelines-and-quality-standards/asthma-guideline/> supported by the recommendations of the Royal College of Physicians National Review of Asthma Deaths and the guidance published by NICE on the use of inhaled steroids in the management of asthma TA131 and TA1384 and the NICE Quality Standard for Asthma.

This Quick Guide also takes into consideration the approach proposed by NICE in the draft guideline on the diagnosis and monitoring of asthma that has recently been put out for consultation. At various points we allude to the draft NICE recommendations.

The Quick Guide attempt to demystify asthma diagnosis and provide sound pragmatic and succinct guidance on diagnosing

this condition in children and adults and explores current management supported by evidence-based guidance. To download your copy of the guideline visit <https://www.pcrs-uk.org/resource/Guidelines-and-guidance/AQG>

PCRS-UK Quick Guide to the Diagnosis and Management of COPD



The ever popular PCRS-UK Quick Guide to the Diagnosis and Management of COPD has had a minor revision to take account on new drug therapies available in COPD. The Quick Guide, endorsed by the Royal College of General Practitioners and British Lung Foundation, offers concise and structured information based on national guidance for the diagnosis and management of COPD in an easily digestible format. Download your copy of the guideline at <https://www.pcrs-uk.org/resource/Guidelines-and-guidance/QG1COPD>

Severe Asthma E-Learning course now available

A Severe Asthma programme developed by Respiratory Education UK in conjunction with PCRS-UK and funded by Novartis is now available. This eLearning programme is free of charge and designed for practitioners in primary care who have some understanding of asthma management. It will take you approximately 45 minutes to complete and will allow you to print off your own certifi-

PCRS-UK News Round-Up



cate of completion. To access the on-line course please visit <http://lms.innovesolutions.co.uk/shared/start/key:GSKMFPJU>

GOVERNMENT SIGNALS NEW STRATEGY TO DRIVE DOWN SMOKING RATES

The government has indicated that it is committed to developing a new five year tobacco control strategy as its previous

approach Healthy Lives, Healthy People comes to an end. Health minister Jane Ellison, made the comments at the launch of the ASH (Action on Smoking and Health) Smoking Still Kills report,¹ which has been endorsed by PCRS-UK.

The report outlines ambitious targets to reduce smoking rates in the UK and sets out recommendations for working with the Government and local authorities to put them into practice.

It recommends that healthcare professionals should adhere to NICE smoking cessation guidance² when helping smokers to quit. This advises using brief interventions and referring patients to smoking cessation services where appropriate. The report adds that clinicians should increase the support and information available to smokers, who are unable to quit and should encourage them to switch to less harmful sources of nicotine, in

line with the principles set out in the NICE guidance on tobacco harm reduction.³ Following input from PCRS-UK, ASH included a recommendation that training on providing very brief advice on smoking cessation should be included in the core curricula of all education programmes for healthcare professionals.

PCRS-UK evidence-based online smoking cessation resources provide essential information for clinicians who want to use the most effective ways of helping patients to quit visit <https://www.pcrs-uk.org/smoking-cessation>.

1. Smoking Still Kills ASH 2015. www.ash.org.uk/files/documents/ASH_962.pdf
2. National Institute for Health and Care Excellence (NICE). Smoking cessation guidance. February 2008. <http://www.nice.org.uk/guidance/ph10>
3. NICE Quality Standard. Smoking tobacco harm reduction approaches July 2015 <http://www.nice.org.uk/guidance/qs92>

Other News

ASTHMA UK - SPIKE IN ASTHMA ADMISSIONS PREDICTED AS NEW SCHOOL YEAR BEGINS

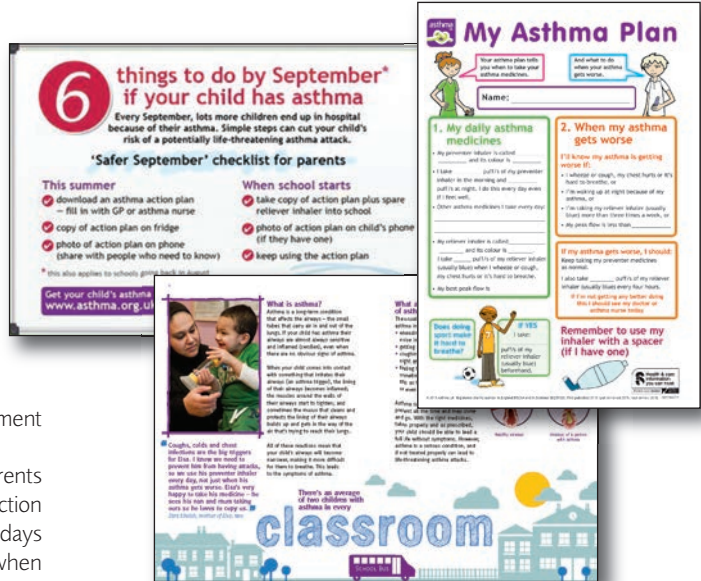
Every September there is a spike in childhood asthma admissions according to Asthma UK as pupils return to school.

There are various theories for this peak in admissions:

- Children not taking their preventer inhalers during the summer holidays
- High levels of environmental allergens in late summer
- Higher stress levels as a result of returning to school
- Exposure to respiratory viral infections in the school environment

Over the summer holidays Asthma UK has been encouraging parents of primary school aged children to download one of its new child action plans so they can establish good routines over the summer holidays to avoid the likelihood of being affected by the above factors when they return to school. Asthma UK action plans for children have recently been updated to reflect the latest BTS/SIGN asthma guidelines and include new pointers for parents about how to get the most from their child's action plan, including:

- Make it easy for you and your family to find it when you need it
- Take a photo and keep it on your mobile (and your child's mobile if they have one)
- Stick a copy on your fridge door
- Share your child's action plan with school, grandparents and babysitter (a printout or a photo).



To download a free copy of the new action plan go to www.asthma.org.uk/downloads

Some of you may have taken part in the PLEASANT study in which the parents of children with asthma in intervention practices were sent a letter in August reminding them to maintain their child's preventer medication to protect them against this surge in asthma attacks just after the school terms starts. The study is complete and the results are awaited... Watch this space!

British Lung Foundation Professionals



Bethany Bateman at BLF highlights the role of BLF professionals

British Lung Foundation (BLF) professionals are health care professionals (HCPs) with a special interest in respiratory care. We want to create a network of BLF Professionals, pooling all of our passion to fight lung disease, helping us to work together and subsequently achieve more.

There are hundreds of HCPs around the country supporting the BLF work in lots of ways. They are talking to those affected by lung disease at our Breathe Easy support groups. They are helping raise awareness by speaking to the media and writing blogs for our website. They're informing our work as we endeavour to influence government and NHS policy for the better, and they are ensuring we are giving people living with lung disease accurate and helpful information through our leaflets and website. For all the projects and campaigns with which HCPs help, we have set up our BLF Professionals network - to recognise the hard work HCPs are doing to fight lung disease, and to ensure that patients and colleagues know about it too.

So many HCPs are doing a wonderful job supporting their own patients, but want to be able to reach out and make a difference in the wider respiratory world. Being a part of BLF Professionals gives them an opportunity to do so.

Of course there are various benefits for HCPs who join the BLF Professionals network, such as bursaries for work shadowing, priority booking for study days, and £30 reimbursement against PCRS-UK membership. There are also opportunities for joint working with our regional managers, and evaluations that enable BLF Professionals to feed into the future of the programme (as well as help us secure more funding for it). However, most HCPs

join the programme for the difference they can make to patients and to the wider fight to reduce the impact of lung disease.

A BLF Professional badge can even inspire greater confidence in patients. Here's what Jo Wright, whose husband Dave is living with idiopathic pulmonary fibrosis (IPF), has to say about being supported by a BLF Professional, Sarah: "I think I feel more confident that my nurse is a BLF Professional as it makes me feel she would have more expertise, more contacts and greater access to resources. The name BLF conjures up an image of a very professional and knowledgeable organisation, giving me more peace of mind – I find it very comforting to know my nurse is part of a well-known organisation. I can't praise Sarah enough for how caring and supportive she is. I want to thank her and all BLF Professionals for the work they are doing and the awareness they are bringing to others."

Of course, we don't want to wear out HCPs by asking them to do everything. By telling us the type of activity they're interested in – media work, patient support, information review, or anything else – we know who we can call on when we need help and when we have opportunities with which they are most likely to want to get involved. However, in order to take advantage of all the opportunities we have, we need the network to grow. We want to do everything we can to support patients and fight lung disease across the UK, and we need your help to do it.

To find out more about becoming a BLF Professional, visit the BLF website <https://www.blf.org.uk/Page/blf-professionals>



Supporting you and your patients

We provide a range of support and information for people living with COPD and other lung conditions.

Our support includes:

- The BLF Helpline: **03000 030 555**
- A national network of **Breathe Easy** support groups
- Comprehensive COPD information online: **www.blf.org.uk/COPD**
- A range of leaflets and booklets for your patients: **www.blf.org.uk/publications**
- COPD patient passport available in print and online: **www.blf.org.uk/passport**

Leading
the **fight**
against
lung disease

Helping you develop your services

We also provide support and advice on service improvements and redesign across the respiratory pathway.

We offer:

- Bespoke training packages
- Awareness campaigns to support early diagnosis
- Organise patient engagement
- And much more:
www.blf.org.uk/hcp

To find out more, please contact:

- **020 7688 5555**
- **enquiries@blf.org.uk**

Delivering Excellence Locally

Featuring initiatives led by PCRS-UK members around the UK, supported by PCRS-UK programmes and tools

Delivering Excellence Locally in West Yorkshire



Francesca Robinson talks to **Dr Anuj Handa**

Strong clinical relationships are being forged and best practice shared in Yorkshire by the PCRS-UK affiliated West Yorkshire Respiratory Network.

The network's quarterly meetings bring together clinical respiratory leads from primary, community and secondary care and public health with key personnel from Clinical Commissioning Group (CCG) and commissioning support units.

The group has been looking at variation in health care outcomes by comparing different CCGs and then drilling down into different practices within the CCGs.

At a recent meeting one of the CCGs with few avoidable respiratory admissions shared information about the systems they have in place.

Action points from the meeting for CCGs included:

- Identify key points in the patient pathway where improvement is needed, look at those issues and work collaboratively to find solutions
- Consider what needs to happen at system level, not just in primary or secondary care.

At the next meeting each CCG brought three problems with which they struggle in their area and then worked with their locality colleagues in small groups on how they were going to tackle them. The issues included quality of spirometry, respiratory education of primary care professionals, disparities in availability of pulmonary rehabilitation and communication across sectors within localities. Feedback on the success of those measures will be shared at the next meeting.

Dr Anuj Handa, a Huddersfield GP and PCRS-UK Lead for West Yorkshire, says: "The Network was able to review the five-year strategic plan for each CCG in West Yorkshire area, looking at the focus for respiratory care. This enabled the network to generate a work plan based on the common themes in West Yorkshire; to be used to focus the aspects of respiratory care that local CCGs are interested in and need to tackle."

"How to improve commissioning for QIPP (Quality Innovation Productivity Prevention) will be the focus of the next session. We will be moving the meeting away from a lecture and workshop format with less talking from the front and more working in locality groups to commit to specific pieces of improvement work."

"We have made a list of best practice in our area, for example introducing an asthma template for SystmOne practices, to share with the group. In addition we will looking at how services can be improved using the PCRS-UK improvement tools including the EQUIP tools."

Sheffield CCG recently joined the group after meetings were promoted through mailings to PCRS-UK members across the region and advertised on the PCRS-UK website. The network is now working to spread the word and extend its representation into the Humber region.

The West Yorkshire Respiratory Network recently affiliated to PCRS-UK and Dr Handa says support from PCRS-UK enabled wider marketing of the meetings as well as the opportunity to talk with experienced leaders including Stephen Gaduzo, and to get input from Bronwen Thompson on making the best use of NHS policy has really helped the group to develop. Local support from Lisa Chandler a former strategic health authority respiratory manager has also been invaluable.

Delivering Excellence Locally in the East of England



Francesca Robinson interviews **Daryl Freeman**

A new venture to bring together healthcare professionals with a special interest in respiratory care has been getting underway in the East of England.

This has been a strong area for networks of nurses for some time, and several local groups affiliated to PCRS-UK provide a forum where ideas can be shared and mutual support given. The only formal Strategic Clinical Network with a focus on respiratory disease is also in the East of England area. So it seemed the perfect time and place to bring together respiratory interested professionals from across the area.

The inaugural meeting in Peterborough was held alongside the East Anglian Thoracic Society (EATS) meeting. The meeting started with a presentation on the diagnosis and management of interstitial lung disease from Professor Sherwood Burge followed by a session on palliative care by Geraldine Burge, a discussion about integrated care and a debate about NHS England's Five Year Forward View. Community nurses shared news about the projects with which they are involved.

Dr Daryl Freeman, a Norfolk GP, East of England PCRS-UK lead and Clinical Director NHS England- Midlands & East (East), said the meeting was oversubscribed and feedback was very positive. One person said they would have driven 150 miles to hear Professor Sherwood's presentation. Many of those present did not know much about the Five Year Forward View and felt the meeting provided them with a useful update.

"There was tremendous enthusiasm from everyone who came to the meeting. We have to travel a long way from East Anglia to national meetings in other parts of the country so there is a real need for a local meeting to enable us to share good practice in respiratory care."

"The plan for the next meeting in November is to have a primary care session probably focussing on sleep disorders and asthma-COPD overlap syndrome before the tea break, then to join the EATS members for combined sessions to include respiratory prescribing across boundaries to highlight the issues we face when prescribing for patients in primary, community and secondary care settings," said Dr Freeman.

Affiliated Groups in East of England

- Peterborough Nurses Respiratory Interest Group
- The West Norfolk Respiratory Nurse Group
- North Norfolk FORD
- FORD (Focus on Respiratory Disease)
- South Norfolk FORD
- Suffolk Respiratory Group
- Barking, Dagenham & Havering Respiratory Interest Group
- Thanet Respiratory Interest Group

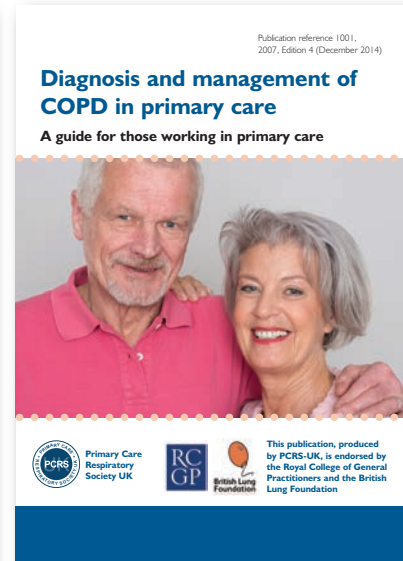
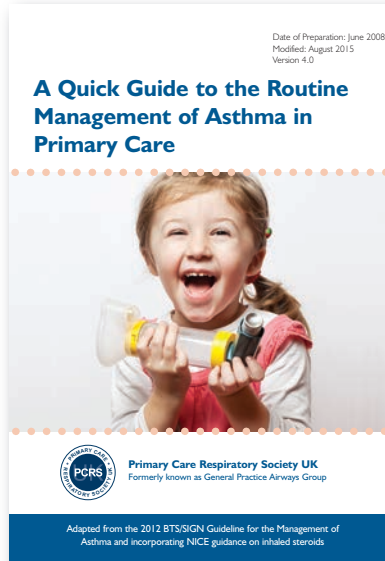
To contact a group leader log-in to the PCRS-UK website and go to the Members Directory at <https://www.pcrs-uk.org/directory>

Affiliating your locality network to PCRS-UK offers an opportunity to share the work you do on a wider platform. PCRS-UK can offer advice and support and promote any events to its members in the local area. If you are interested in affiliating your group please contact us for more information at info@pcrs-uk.org

PCRS-UK Resources and Improvement Tools: Reviewing diagnosis

In addition to the newly updated Quick Guides for the Diagnosis and Management of asthma (<https://www.pcrs-uk.org/resource/Guidelines-and-guidance/AQG>) and COPD (<https://www.pcrs-uk.org/resource/Guidelines-and-guidance/QGCOPD>) both of which offer sound, evidence based guidance on how to diagnose these conditions.

PCRS-UK has also developed two Practice Improvement Worksheets to support health-care professionals:-



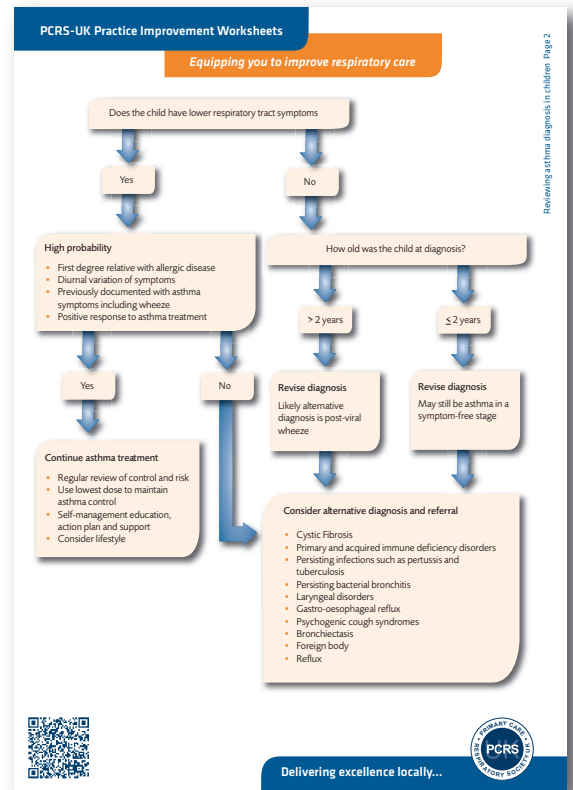
Reviewing Asthma Diagnosis in Childhood

Many children suffer from lower respiratory symptoms in early childhood. For some, this will be asthma, but for many, they have had a complex post-viral illness, often associated with Respiratory Syncytial Virus (RSV), resulting in intermittent symptoms of cough, wheeze and breathlessness.

It is important not to label children as having asthma if they do not have a confirmed diagnosis of asthma. Young adults with an erroneous label of childhood asthma may find that their career options are inappropriately limited as a consequence of an early and inaccurate diagnosis of asthma.

Confirming a diagnosis in very young children can be difficult, and it is therefore important to review children with a diagnosis of asthma or wheezing in infancy as they get older. This simple worksheet with a practical flow chart enables healthcare practitioners to review a prior diagnosis and consider possible alternative diagnoses.

The worksheet, available to members of PCRS-UK can be downloaded at <https://www.pcrs-uk.org/resource/Improvement-tools/reviewing-asthma-diagnosis-children-improvement-worksheet>



Accurate Diagnosis of COPD

Early and accurate diagnosis of COPD is important, as all too often people do not have their diagnosis confirmed until they are well down the trajectory of functional decline. There are a range of different diseases (both respiratory and non-respiratory) that present with symptoms similar to COPD, and these should be considered and excluded as part of a making a COPD diagnosis. This simple worksheet offers a practical flow chart to support busy healthcare practitioners on the steps to consider when diagnosing COPD.

The worksheet, available to members of PCRS-UK, can be downloaded at <https://www.pcrs-uk.org/resource/Improvement-tools/accurate-diagnosis-copd-improvement-worksheet>

What every GP should know about rarer lung conditions

Rarer respiratory conditions are, by definition, rare and each practice will only have a handful of such patients. The most important things that a GP needs to know are:

- Limits of your own knowledge.
- How such patients present.
- Referral pathways to secondary care.
- What the local Community Respiratory Team can provide

This simple, two-page guide includes brief advice on rarer lung conditions including their presentation, investigations and other steps to take when considering these rarer disorders such as interstitial lung disease, sarcoidosis, pulmonary vasculitis, bronchiectasis pulmonary hypertension, tuberculosis and aspergilliosis.

You can download your copy of the opinion sheet at <https://www.pcrs-uk.org/resource/Opinion-sheets/what-every-gp-should-know-about-rarer-lung-conditions-opinion-sheet>

Our excerpt from *npj Primary Care Respiratory Medicine* this issue features an article on the diagnosis and management of interstitial lung disease see page 40.

New respiratory leaders inspired by PCRS-UK leadership event

Creating a case for change and getting it heard was the title of the PCRS-UK respiratory leaders' event in June that gave delegates the opportunity of creating, building and delivering a business case.

The two-day event helped delegates to look at the structure of a business case, review real life examples then develop and deliver an actual business case.

This was the second respiratory leaders' event that Clare Cook, clinical lead for respiratory services at Bristol Community Health had attended along with her colleague and operational lead Laura Turner. "We use these events as an opportunity to have an away day together to plan projects and take a 'temperature check' of other services similar to ours. It is definitely worth the budget and the time for both of us to go," she said.

She added: "I am new in my leadership post, as is Laura, and it helps to learn from other people's experiences and check our priorities against other people's priorities.

"I found one of the speakers, Catherine Blackaby, Senior Improvement Manager with NHS Improving Quality, very inspiring. She delivered a presentation on how to put together a business case. I have mirrored that process about providing evidence for the things I want to achieve from the business plan and was recently successful in getting an uplift of £100,000 for our budget."

Dr Helen Ward, community lead and consultant respiratory physician at the Royal Wolverhampton Hospitals NHS Trust and a member of

the British Thoracic Society integrated working party, says she found the meeting very informative - the presentations on developing a business case and on recent policy updates were particularly useful.

"There were some very good leaders and experts in the room and there was a lot of knowledge shared around the tables where we sat.

"I'm a relatively new consultant, so a leadership course was really useful not just from the business aspect but also to meet people within primary care who have passion for respiratory - that is really valuable for me from a networking point of view.

"The meeting was very welcoming and the people running it came across as being very nurturing of the people coming through - they are there to support you as leaders because that support is not always available elsewhere," said Dr Ward.

Dr Noel Baxter, GP and member of the PCRS-UK executive, who helps to run the respiratory leaders programme, said: "It was great to chair a meeting with such a diversity of respiratory interested health professionals learning from each other and working together. I am confident that this course is contributing to more coordinated care for patients with complex health problems."

The next respiratory leaders meeting will be held on 6th-7th November 2015 at the Lea Marston Hotel, Sutton Coldfield. The theme is "Delivering value for patients and demonstrating outcomes". For more information visit our website at <https://www.pcrs-uk.org/respiratory-clinical-leadership-workshop>

PCRS-UK Affiliated Group Leaders Meeting 15th October 2015 Scanning the Horizon – How to get ahead of the game

Whether you're an existing leader of a PCRS-UK Affiliated Group, or you're thinking about setting one up make sure you don't miss out on our annual meeting for group leaders.

This year's event is a really practical workshop-style meeting to help inspire potential new group leaders and support existing group leaders to get the most of your meetings, help you with some of the challenges of running meetings and give you some useful tips and ideas for topics to share with your local networks all relevant to primary care and respiratory health.

The meeting also offers you an opportunity to network with other experienced affiliated group leaders and share ideas and tips.

Chaired by Carol Stonham and facilitated by Ren Lawlor and Allwin Mercer, this half-day interactive event also features guest speaker Beverley Bostock Cox. The meeting runs from 12.00 – 17.30 is

FREE of charge to PCRS-members and will include sessions on:-

- Revalidation and appraisal – supporting your group nurse members to prepare for forthcoming revalidation and appraisal
- Getting the best out of your meetings:-
 - Chairing your meetings
 - Dealing with difficult/disruptive/disengaged delegates
 - Planning ahead
 - Making your speakers feel valued
 - Evaluating your meetings
- Making the case for training and development and getting your voice heard

To find out more about the meeting, download a programme and register visit our website at <https://www.pcrs-uk.org/leaders-events>

Spirometry goes vertical

The future of spirometry testing in primary care



Introducing the new **SpiroConnect** PC based spirometer, designed by the inventor of the original turbine spirometer, Chris Lawson, Micro Medical's Technical Director for 27 years.

With its patented vertical turbine, this new and improved spirometer is more sensitive to the low flow rates seen in COPD and Asthma patients.

What's more, it is the most comprehensively integrated with the EMIS, SystemOne and InPS clinical systems, filing over 40 user selectable Read codes and a PDF spirometry report with a single click at the end of the test.

SpiroConnect • Safer • Faster • Easier • More accurate

To find out more about our special introductory and trade in offers

Visit: www.numed.co.uk **Call:** 0114 2433896 **Email:** sales@numed.co.uk

Equip yourself to take the lead in respiratory primary care



- Interactive skills & knowledge based workshops, free to attend for PCRS-UK members
- Supportive and safe environment to develop and practice a range of skills and network with like minded colleagues
- Access to a faculty of experienced leaders - all practising primary care clinicians who understand the realities and challenges of driving improvement

Next event: Delivering value for patients and demonstrating outcomes
6th-7th November 2015,
Lea Marston Hotel, Sutton Coldfield

- Want to make a real difference for respiratory patients in or beyond your practice, ward or clinic or consider a wider population?
- Would you like to take help and push forward work with your colleagues seeing patients in their community to influence and improve respiratory services in your area?
- All it takes is just one person to take the initiative. And with our help, that person could be you.



Open for Submissions

The only fully-indexed scientific journal devoted to the management of respiratory diseases in primary care.



npj Primary Care Respiratory Medicine is an online-only, open access journal, publishing papers representing important advances of significance to specialists within the fields of primary care and respiratory medicine.

npj Primary Care Respiratory Medicine publishes internationally-relevant open research that is essential to the future of primary care management of respiratory and respiratory-related allergic diseases.

Submit your next manuscript and benefit from:

- Strong editorial values
- Competitive turnaround times
- Wide dissemination and high visibility
- Editorial Summaries
- Compliance with international open access funding mandates

In partnership with



EDITORS-IN-CHIEF

Professor Aziz Sheikh

The University of Edinburgh, Edinburgh, UK

Dr. Paul Stephenson

Honorary Clinical Research Fellow, Allergy and Respiratory Research Group, Centre for Population Health Sciences, The University of Edinburgh, Edinburgh, UK

FREQUENCY OF PUBLICATION

Continuous, new content published weekly

2014 IMPACT FACTOR

2.504*

All content is indexed within Web of Science, and PubMed/Medline

*2014 Journal Citation Report (Thomson Reuters, 2015), formerly published under *Primary Care Respiratory Journal*

Part of the Nature Partner Journals series

npj nature partner journals

PERSPECTIVE **OPEN****Interstitial lung disease: raising the index of suspicion in primary care**Joseph D Zibrak¹ and David Price²

Interstitial lung disease (ILD) describes a group of diseases that cause progressive scarring of the lung tissue through inflammation and fibrosis. The most common form of ILD is idiopathic pulmonary fibrosis, which has a poor prognosis. ILD is rare and mainly a disease of the middle-aged and elderly. The symptoms of ILD—chronic dyspnoea and cough—are easily confused with the symptoms of more common diseases, particularly chronic obstructive pulmonary disease and heart failure. ILD is infrequently seen in primary care and a precise diagnosis of these disorders can be challenging for physicians who rarely encounter them. Confirming a diagnosis of ILD requires specialist expertise and review of a high-resolution computed tomography scan (HRCT). Primary care physicians (PCPs) play a key role in facilitating the diagnosis of ILD by referring patients with concerning symptoms to a pulmonologist and, in some cases, by ordering HRCTs. In our article, we highlight the importance of prompt diagnosis of ILD and describe the circumstances in which a PCP's suspicion for ILD should be raised in a patient presenting with chronic dyspnoea on exertion, once more common causes of dyspnoea have been investigated and excluded.

npj Primary Care Respiratory Medicine (2014) **24**, 14054; doi:10.1038/npjpcrm.2014.54; published online 11 September 2014

INTRODUCTION

Interstitial lung disease (ILD) is an umbrella term, synonymous with diffuse parenchymal lung disease, for a large group of lung diseases affecting the tissue and space around the air sacs of the lungs, which cause progressive scarring of lung tissue through inflammation and fibrosis.¹ ILD is uncommon compared to other pulmonary problems seen in general practice. The prevalence of ILD has been estimated as 81/100,000 in males and 67/100,000 in females.²

Primary care physicians (PCPs) often misdiagnose ILD as chronic obstructive pulmonary disease (COPD), bronchitis, emphysema, asthma, heart disease or other common diseases with similar symptoms.³ In a US survey of 1,583 patients, 54.6% reported a delay of ≥ 1 year between the onset of breathing problems and receiving a diagnosis of pulmonary fibrosis.³

A diagnosis of ILD should be confirmed by a pulmonologist working with a radiologist experienced in the differential diagnosis of ILD, but PCPs have a crucial role to play in recognising the need for further evaluation. This paper describes the circumstances in which a PCP's suspicion for ILD should be raised in a patient with chronic dyspnoea and the importance of referring patients who may have ILD promptly to a pulmonologist to confirm the diagnosis.

WHAT IS ILD?

The American Thoracic Society and European Respiratory Society define ILD as a heterogeneous group of non-neoplastic disorders resulting from damage to the lung parenchyma by inflammation and fibrosis that diminish the lung's capacity for alveolar gas diffusion.¹ The latest classification of ILDs is shown in Figure 1.^{1,4} Most ILDs are 'restrictive' pulmonary disorders, i.e., the lungs have a reduced ability to expand on inhalation. This is in contrast to 'obstructive' pulmonary disorders such as asthma, COPD and

emphysema, in which the airways of the lungs become narrowed or blocked so the patient cannot exhale completely. Some forms of ILD are associated with environmental or occupational exposures (e.g., asbestosis or silicosis), the use of certain drugs⁵ or with connective tissue disorders.^{2,6,7} Others, including the idiopathic interstitial pneumonias, have no known cause.¹ Some cases of ILD do not meet the specific definitions for any form of ILD and are considered 'unclassifiable'. This may be the case when there are non-specific or conflicting clinical, radiological or histopathological findings, or when patients are unable or unwilling to undergo diagnostic procedures.⁸

Idiopathic pulmonary fibrosis (IPF) is the most common idiopathic interstitial pneumonia.¹ IPF also has the worst prognosis; the 5-year survival rate (20–40%)⁹ is lower than many common cancers.¹⁰ While most cases of IPF are sporadic, familial forms of the disease may account for ~5–20% of cases.^{11,12} The prevalence of IPF in the US has been estimated as 14.0–27.9 cases/100,000 using narrow case definitions and as 42.7–63 cases/100,000 using broad case definitions.¹³ In Europe, the estimated prevalence of IPF was 1.25–23.4 cases/100,000.¹³ The incidence of IPF increased with age and was higher in men than in women (10.7 vs. 7.4 per 100,000 patient-years in the US).¹³

RECOGNISING ILD

As highlighted in the international guidelines for the diagnosis of ILD, a diagnosis of ILD should be made by review of clinical, pulmonary, radiological and possibly histopathological features.⁴ However, PCPs have a key role to play in identifying patients who may have ILD and referring them to a pulmonologist or an ILD clinic with a multidisciplinary team. PCPs can also help detect non-idiopathic forms of ILD by reviewing relevant occupational, environmental and drug exposures, as well as by assessing the presence of extrapulmonary symptoms suggestive of connective tissue disease.

¹Beth Israel Deaconess Medical Centre, Harvard Medical School, Boston, MA, USA and ²Division of Applied Health Sciences, University of Aberdeen, Aberdeen, UK.
Correspondence: JD Zibrak (jzibrak@caregroup.harvard.edu)

Received 1 October 2013; revised 2 July 2014; accepted 8 July 2014

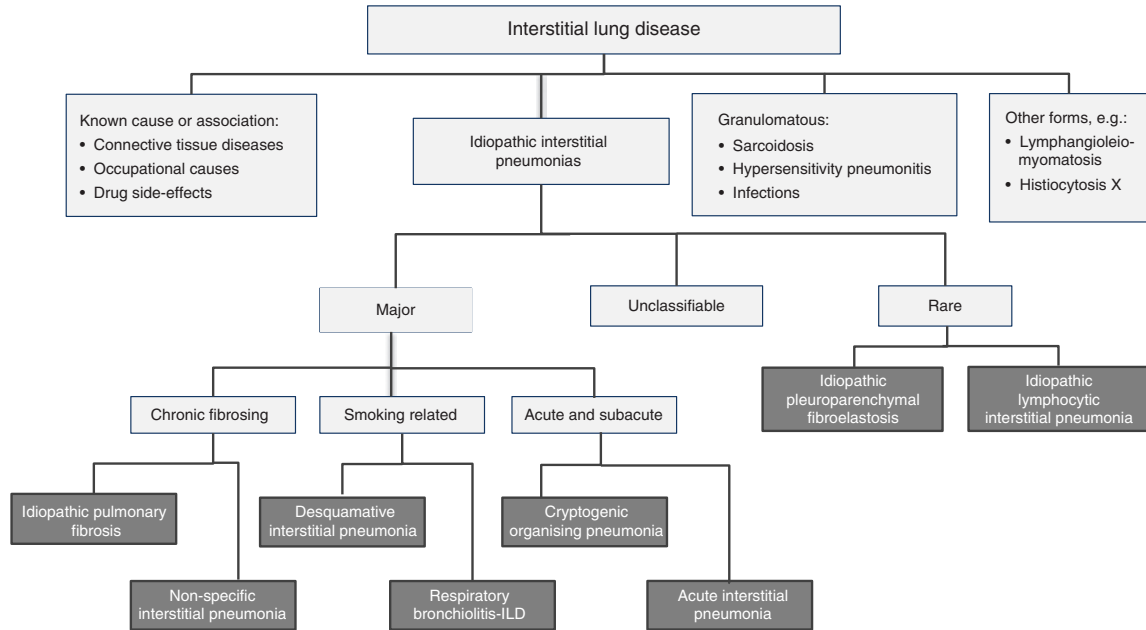


Figure 1. Classification of ILDs.⁴

One of the challenges for PCPs in determining whether a patient may have ILD is the lack of specificity of its symptoms. Breathlessness and cough are seen in many diseases that commonly occur in middle-aged and elderly patients, most notably COPD and heart failure. Once non-respiratory causes of breathlessness have been excluded, PCPs should consider the possibility of ILD in all middle-aged or elderly patients who present with unexplained chronic dyspnoea on exertion, particularly those who are more breathless than would be expected based on their lung function and other contributory factors such as obesity, or who have a cough of more than several months' duration. The suspicion of ILD should be raised in patients with a presumptive obstructive lung disease or congestive heart failure who fail to benefit from regular therapy, and in patients with a non-specific chest X-ray.

A diagnosis of asthma or COPD should be confirmed using the appropriate diagnostic criteria. If diagnostic criteria are not met, further assessments should be performed to ensure that the correct diagnosis is made and patients are not given inappropriate therapies. A diagnosis of COPD should be confirmed by spirometry. Obstructive lung disease is characterised by a disproportionate reduction of maximal airflow from the lung in relation to the maximal volume that can be displaced from the lung (forced vital capacity (FVC)), defined by a significantly reduced forced expiratory volume in 1s (FEV₁)/FVC ratio.¹⁴ Spirometry may be normal in the early phases of ILD, and patients with both pulmonary fibrosis and emphysema often have lung volumes within normal limits.¹⁵ However, restrictive lung disease tends to be characterised by a reduction in total lung capacity and a normal or elevated FEV₁/vital capacity ratio.¹⁶ Patients with ILD tend to show a reduced FVC, which declines as the disease progresses,¹⁷ and many experience oxygen desaturation with only minimal exertion. PCPs often order chest radiographs in patients with chronic respiratory symptoms. As patients with COPD have largely normal chest radiographs, the presence of even subtle abnormalities such as increased interstitial markings or nodular opacities warrants further investigation. Patients presenting with a history of ≥6 months' breathlessness on

exertion with no evidence of obstruction should be assessed for conditions that may cause breathlessness, such as chronic heart failure, asthma, COPD, bronchiectasis, pleural effusion, abdominal splinting and anaemia.¹⁸ If these conditions are not present, patients should not only have spirometry and diffusing capacity for carbon monoxide measured, but also be referred for a high-resolution computed tomography scan. High-resolution computed tomography scans are vital for the diagnosis of ILD and allow for differentiation among the types of ILD.¹⁹

PCPs should take a comprehensive medical history in all patients presenting with chronic dyspnoea, including questions about their employment history, as some forms of ILD such as asbestosis and silicosis are linked to environmental factors,¹ and individuals who have worked with livestock or birds, or been subjected to high exposure to dust or fumes, may have a higher risk of IPF.²⁰ A medical history may reveal the use of medications that may cause drug-induced ILD (DILD), such as chemotherapeutic agents (e.g., cyclophosphamide, carmustine, busulphan), antibacterial agents (e.g., sulphonamide, nitrofurantoin) or cardiovascular agents (e.g., amiodarone).⁵ The likelihood of developing DILD is idiosyncratic and largely unpredictable and the underlying mechanisms are not fully understood.⁵ PCPs should also assess for extrapulmonary symptoms, such as fatigue, joint pain and stiffness, which may suggest connective tissue disease.²⁰

An early clinical test that can aid diagnosis of ILD is lung auscultation. Patients with many types of ILD commonly present with inspiratory 'crackles'. These crackles, which are discontinuous, short sounds similar to those heard when slowly separating Velcro, are heard even in the early stages of ILD when the patient takes slow, deep breaths.²¹ Unlike the crackles occasionally heard in healthy elderly subjects, which disappear after several deep breaths, these fine crackles are heard consistently at end inspiration and originate from the basal areas of the lung, progressing to the upper zones as disease becomes more severe.²¹ Although higher in frequency,²¹ crackles associated with ILD may be confused with the crackles heard in patients with heart failure. Finger clubbing, where reduced oxygenation of the blood

causes the fingertips and fingernails to spread out and become rounder, is seen in 25–50% of patients with IPF.²²

The differential features of ILD and COPD are shown in Table 1.

IMPORTANCE OF PROMPT DIAGNOSIS OF ILD

Patients with ILD who are misdiagnosed may receive inappropriate and ineffective treatment. Such treatment may be harmful physically (e.g., patients may experience adverse effects of medication without gaining any benefit) or psychologically (e.g., it is disheartening for patients to be given several treatments that do not work). There are few recommended treatments for ILD, although corticosteroids appear to be effective in the treatment of cryptogenic organising pneumonia and desquamative interstitial pneumonia.²³ The latest international guidelines for the management of IPF, issued in 2011, did not recommend any specific pharmacological treatments for the chronic treatment of IPF, but long-term oxygen therapy was strongly recommended for patients with clinically significant resting hypoxaemia and a weak recommendation was made for pulmonary rehabilitation in the majority of patients.¹²

The anti-fibrotic agent pirfenidone has been approved for the treatment of IPF in Japan, India, China and Canada and for the treatment of mild to moderate IPF in Europe, but has not been approved in the US. An additional Phase III trial of pirfenidone has recently reported results and showed that pirfenidone reduced the decline in FVC percent predicted.²⁴ There are several other agents in clinical development for the treatment of IPF, the most advanced of which is the tyrosine kinase inhibitor nintedanib, for which the results of two Phase III INPULSIS trials have recently been announced.²⁵ In both the INPULSIS trials, nintedanib significantly reduced the decline in FVC over the 52-week treatment period, consistent with a slowing of disease progression. In contrast, warfarin and triple therapy with prednisone, azathioprine and N-acetylcysteine have been shown to be harmful for patients with IPF,^{26,27} highlighting the importance of potential therapies being evaluated in controlled trials before they are used in clinical practice. Acetylcysteine alone appears not to be harmful, but to have no efficacy as a treatment for IPF.²⁸

PCPs treating patients with ILD should address comorbidities such as gastroesophageal reflux disease, pulmonary hypertension (PH) and emphysema. The prevalence of gastroesophageal reflux disease in patients with ILD²⁹ and IPF³⁰ is very high and there is some evidence that the use of gastroesophageal reflux disease treatments in patients with IPF is a predictor of longer survival time.³¹ PH is also common in patients with ILD.³² In patients with IPF, PH has a reported incidence of 32–84% and is associated with decreased exercise capacity and worse prognosis.³³ Patients with IPF, emphysema and PH have a particularly dismal prognosis.¹⁵ Patients with IPF may also have an increased risk of acute coronary syndrome and deep vein thrombosis.³⁴

The clinical course and prognosis of the different types of ILD are highly variable.¹ Even if pharmacological treatment is not appropriate, it is important that a patient receives the correct diagnosis so that their physician can provide appropriate information and support. Data from a US-based survey of 1,448 patients with IPF or caregivers of patients with IPF highlight that there is a need for improved patient education regarding diagnosis and management.³ Less than half of the respondents felt well informed about treatment options, and the role of pulmonary rehabilitation, supplemental oxygen and lung transplantation.³ Receiving a diagnosis with such a poor prognosis may be difficult for the patient, and they may need time to start to come to terms with the diagnosis before they can fully comprehend the information they receive. Patients may also benefit from contacting patient organisations for advice and support.

Table 1. Differential features of ILD and COPD

	ILD	COPD
<i>Physical examination</i>		
Inspiratory crackles ²¹	✓	×
Wheezing ¹⁴	×	✓
Finger clubbing ²²	✓	×
<i>Spirometry</i> ¹⁶		
Reduction in TLC	✓	×
Reduced FEV ₁ /FVC ratio	×	✓
Response to bronchodilators	×	✓
Chest radiographs ¹	Show specific patterns of abnormality	Largely normal

Abbreviations: COPD, chronic obstructive pulmonary disease; FEV₁, forced expiratory volume in 1s; FVC, forced vital capacity; ILD, interstitial lung disease.

The majority of patients with ILD should have the opportunity for referral to a pulmonary rehabilitation programme. Comprehensive programmes involve not only exercise training with aerobic conditioning, strength and endurance training and respiratory therapy, but also behaviour modification techniques.³⁵ In a large cohort of 402 patients with IPF, pulmonary rehabilitation had a positive effect on patients' pulmonary function and quality of life.³⁶

At present, lung transplant is the only therapy that has been shown to improve survival in patients with IPF, but it is available only to selected patients.²² The eligibility criteria for lung transplantation vary between and even within countries, but international guidelines stress that the early referral of patients with IPF for transplantation should be promoted.³⁷ Prompt diagnosis of IPF also makes it more likely that a patient will meet the criteria for participation in a clinical trial; most clinical trials in IPF are restricted to patients with mild or moderate impairment of lung function.

CONCLUSIONS

ILD presents with symptoms similar to other chronic respiratory conditions and is frequently misdiagnosed in primary care, most commonly as COPD and heart failure. Prompt diagnosis of ILD is important to ensure that patients receive appropriate care and support, and to enable them to be considered for lung transplantation or clinical trials. PCPs have a key role to play in expediting the diagnosis of ILD by referring middle-aged/elderly patients with unexplained chronic dyspnoea and cough who do not meet the diagnostic criteria for other diseases to an ILD centre or pulmonologist with expertise in this group of disorders.

ACKNOWLEDGEMENTS

The authors acknowledge the medical writing assistance provided by Clare Ryles and Wendy Morris of Fleishman-Hillard Group, 40 Long Acre, London, during the preparation of this review.

CONTRIBUTIONS

JDZ and DP drafted this article, with medical writing assistance from Clare Ryles and Wendy Morris of Fleishman-Hillard Group. The authors are fully responsible for all content and editorial decisions, were involved at all stages of manuscript development, and have approved the final version of the review, which reflects the authors' interpretation and conclusions.

COMPETING INTERESTS

JDZ is a member of the InterMune Pharmaceuticals Clinical Advisory Board. DP has been a consultant to and/or member of the advisory boards for Almirall, AstraZeneca, Boehringer Ingelheim, Chiesi, GlaxoSmithKline, Medapharma, Merck, Mundipharma, Napp, Novartis, Nycomed, Pfizer, Sandoz and Teva. He or his research team have received grants and support for research in respiratory disease from the following organisations in the last 5 years: UK National Health Service, Aerocrine, AstraZeneca, Boehringer Ingelheim, Chiesi, GlaxoSmithKline, Merck, Mundipharma, Novartis, Nycomed, Orion, Pfizer, Takeda and Teva. He has received payment for lectures from Activaero, Almirall, AstraZeneca, Boehringer Ingelheim, Chiesi, Cipla, GlaxoSmithKline, Kyorin, Merck, Mundipharma, Napp, Novartis, Pfizer and Teva, and has been paid for development of educational materials by Boehringer Ingelheim and GlaxoSmithKline. He has been reimbursed for travel expenses by Boehringer Ingelheim, Merck, Mundipharma Napp, Novartis and Teva. He holds shares in AKL, which produces phytopharmaceuticals and owns 80% of Research in Real Life and its subsidiary social enterprise Optimum Patient Care.

FUNDING

The authors have not received payment for this article. Medical writing assistance was funded by Boehringer Ingelheim Pharmaceuticals.

REFERENCES

- American Thoracic Society. American Thoracic Society/European Respiratory Society international multidisciplinary consensus classification of the idiopathic interstitial pneumonias. *Am J Respir Crit Care Med* 2002; **165**: 277–304.
- Coults DB, Zumwalt RE, Black WC, Sobonya RE. The epidemiology of interstitial lung diseases. *Am J Respir Crit Care Med* 1994; **150**: 967–972.
- Collard HR, Tino G, Noble PW, Shreve MA, Michaels M, Carlson B *et al*. Patient experiences with pulmonary fibrosis. *Respir Med* 2007; **101**: 1350–1354.
- Travis WD, Costabel U, Hansell DM, King TE Jr, Lynch DA, Nicholson AG *et al*. An official American Thoracic Society/European Respiratory Society statement: update of the international multidisciplinary classification of the idiopathic interstitial pneumonias. *Am J Respir Crit Care Med* 2013; **188**: 733–748.
- Schwaiblmair M, Behr W, Haeckel T, Markl B, Foerg W, Berghaus T. Drug induced interstitial lung disease. *Open Respir Med J* 2012; **6**: 63–74.
- de Boer S, Wilsher M. Review series: aspects of interstitial lung disease. *Sarcoidosis. Chron Respir Dis* 2010; **7**: 247–258.
- Olson AL, Swigris JJ, Sprunger DB, Fischer A, Fernandez-Perez ER, Solomon J *et al*. Rheumatoid arthritis-interstitial lung disease-associated mortality. *Am J Respir Crit Care Med* 2011; **183**: 372–378.
- Ryerson CJ, Urbania TH, Richeldi L, Mooney JJ, Lee JS, Jones KD *et al*. Prevalence and prognosis of unclassifiable interstitial lung disease. *Eur Respir J* 2013; **42**: 750–757.
- Kim DS, Collard HR, King TE Jr. Classification and natural history of the idiopathic interstitial pneumonias. *Proc Am Thorac Soc* 2006; **3**: 285–292.
- Vancheri C, Failla M, Crimi N, Raghu G. Idiopathic pulmonary fibrosis: a disease with similarities and links to cancer biology. *Eur Respir J* 2010; **35**: 496–504.
- Lawson WE, Loyd JE. The genetic approach in pulmonary fibrosis: can it provide clues to this complex disease? *Proc Am Thorac Soc* 2006; **3**: 345–349.
- Raghu G, Collard HR, Egan JJ, Martinez FJ, Behr J, Brown KK *et al*. An official ATS/ERS/JRS/ALAT statement: idiopathic pulmonary fibrosis: evidence-based guidelines for diagnosis and management. *Am J Respir Crit Care Med* 2011; **183**: 788–824.
- Nalysnyk L, Cid-Ruzafa J, Rotella P, Esser D. Incidence and prevalence of idiopathic pulmonary fibrosis: review of the literature. *Eur Respir Rev* 2012; **21**: 355–361.
- Global Strategy for the Diagnosis, Management and Prevention of COPD. Global Initiative for Chronic Obstructive Lung Disease (GOLD) 2014. (2014). <http://www.goldcopd.org> (accessed 11 May 2014).
- Cottin V, Le PJ, Prevot G, Mal H, Humbert M, Simonneau G *et al*. Pulmonary hypertension in patients with combined pulmonary fibrosis and emphysema syndrome. *Eur Respir J* 2010; **35**: 105–111.
- Pellegrino R, Viegi G, Brusasco V, Crapo RO, Burgos F, Casaburi R *et al*. Interpretative strategies for lung function tests. *Eur Respir J* 2005; **26**: 948–968.
- Ley B, Collard HR, King TE Jr. Clinical course and prediction of survival in idiopathic pulmonary fibrosis. *Am J Respir Crit Care Med* 2011; **183**: 431–440.
- National Institute for Health and Care Excellence (NICE). Clinical Knowledge Summary on breathlessness (2014). <http://cks.nice.org.uk/breathlessness#scenario-recommendation:7> (accessed 11 May 2014).
- Lynch DA, Travis WD, Muller NL, Galvin JR, Hansell DM, Grenier PA *et al*. Idiopathic interstitial pneumonias: CT features. *Radiology* 2005; **236**: 10–21.
- Baumgartner KB, Samet JM, Coultas DB, Stidley CA, Hunt WC, Colby TV *et al*. Occupational and environmental risk factors for idiopathic pulmonary fibrosis: a multicenter case-control study. Collaborating Centers. *Am J Epidemiol* 2000; **152**: 307–315.
- Cottin V, Cordier JF. Velcro crackles: the key for early diagnosis of idiopathic pulmonary fibrosis? *Eur Respir J* 2012; **40**: 519–521.
- Borchers AT, Chang C, Keen CL, Gershwin ME. Idiopathic pulmonary fibrosis—an epidemiological and pathological review. *Clin Rev Allergy Immunol* 2011; **40**: 117–134.
- Bradley B, Branley HM, Egan JJ, Greaves MS, Hansell DM, Harrison NK *et al*. Interstitial lung disease guideline: the British Thoracic Society in collaboration with the Thoracic Society of Australia and New Zealand and the Irish Thoracic Society. *Thorax* 2008; **63**: v1–58.
- King TE Jr, Bradford WZ, Castro-Bernardini S, Fagan EA, Glasspole I, Glassberg MK *et al*. A phase 3 trial of pirfenidone in patients with idiopathic pulmonary fibrosis. *N Engl J Med* 2014; **370**: 2083–2092.
- Richeldi L, du Bois RM, Raghu G, Azuma A, Brown KK, Costabel U *et al*. Efficacy and safety of nintedanib in idiopathic pulmonary fibrosis. *N Engl J Med* 2014; **370**: 2071–2082.
- Raghu G, Anstrom KJ, King TE Jr, Lasky JA, Martinez FJ. Prednisone, azathioprine, and N-acetylcysteine for pulmonary fibrosis. *N Engl J Med* 2012; **366**: 1968–1977.
- Noth I, Anstrom KJ, Calvert SB, de Andrade J, Flaherty KR, Glazer C *et al*. A placebo-controlled randomized trial of warfarin in idiopathic pulmonary fibrosis. *Am J Respir Crit Care Med* 2012; **186**: 88–95.
- Idiopathic Pulmonary Fibrosis Clinical Research Network, Martinez FJ, de Andrade JA, Anstrom KJ, King TE Jr, Raghu G. Randomized trial of acetylcysteine in idiopathic pulmonary fibrosis. *N Engl J Med* 2014; **370**: 2093–2101.
- Soares RV, Forsythe A, Hogarth K, Sweiss NJ, Noth I, Patti MG. Interstitial lung disease and gastroesophageal reflux disease: key role of esophageal function tests in the diagnosis and treatment. *Arq Gastroenterol* 2011; **48**: 91–97.
- Raghu G, Freudenberger TD, Yang S, Curtis JR, Spada C, Hayes J *et al*. High prevalence of abnormal acid gastro-oesophageal reflux in idiopathic pulmonary fibrosis. *Eur Respir J* 2006; **27**: 136–142.
- Lee JS, Ryu JH, Elicker BM, Lydell CP, Jones KD, Wolters PJ *et al*. Gastroesophageal reflux therapy is associated with longer survival in patients with idiopathic pulmonary fibrosis. *Am J Respir Crit Care Med* 2011; **184**: 1390–1394.
- Hallowell RW, Reed RM, Fraig M, Horton MR, Girgis RE. Severe pulmonary hypertension in idiopathic nonspecific interstitial pneumonia. *Pulm Circ* 2012; **2**: 101–106.
- Pitsiou G, Papakosta D, Bouros D. Pulmonary hypertension in idiopathic pulmonary fibrosis: a review. *Respiration* 2011; **82**: 294–304.
- Hubbard RB, Smith C, Le JI, Gribbin J, Fogarty AW. The association between idiopathic pulmonary fibrosis and vascular disease: a population-based study. *Am J Respir Crit Care Med* 2008; **178**: 1257–1261.
- Holland A, Hill C. Physical training for interstitial lung disease. *Cochrane Database Syst Rev* 2008; (4): CD006322.
- Huppmann P, Szczepanski B, Boensch M, Winterkamp S, Schönheit-Kenn U, Neurohr C *et al*. Effects of inpatient pulmonary rehabilitation in patients with interstitial lung disease. *Eur Respir J* 2013; **42**: 444–453.
- Orens JB, Estenne M, Arcasoy S, Conte JV, Corris P, Egan JJ *et al*. International guidelines for the selection of lung transplant candidates: 2006 update—a consensus report from the Pulmonary Scientific Council of the International Society for Heart and Lung Transplantation. *J Heart Lung Transplant* 2006; **25**: 745–755.



This work is licensed under a Creative Commons Attribution-NonCommercial-ShareAlike 4.0 International License. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in the credit line; if the material is not included under the Creative Commons license, users will need to obtain permission from the license holder to reproduce the material. To view a copy of this license, visit <http://creativecommons.org/licenses/by-nc-sa/4.0/>



Take Action and Improve Your Clinical Skills!

Many patients present co-morbidities and this requires a flexible and confident healthcare service delivered by professionals equipped with the right clinical skills and knowledge.

Our education and training is designed to support you in delivering high quality care in the following areas:

Allergy	Cardiovascular
Diabetes	Musculoskeletal
Professional Development	Respiratory

Whether it is an asthma update, performing and interpreting quality assured spirometry, or building your diabetes prevention and screening knowledge, our education and training helps you take action and improve your clinical skills.

Visit our website to see forthcoming courses in your area of interest, including dates across the UK, or talk to our team about bringing a module or workshop to your region.

Education for Health provides a specialist portfolio of workshops and resources aimed at transforming all levels of patient centred care.

**Book
in your
area**

Can't see a date that suits?

Education for Health runs training throughout the UK to respond to local demand. Talk to our NHS Liaison Managers to discuss setting up a course in your area: www.educationforhealth.org/contact



Join the conversation: #TakeAction



www.educationforhealth.org
www.respiratoryeduk.com

intermedical
CARDIO RESPIRATORY



Spirometry made simple with **spirolab**.

The next generation diagnostic spirometer.
Designed & developed specifically
for UK respiratory professionals.

MIR
spirolab
from *intermedical*

See more at www.spirolab.co.uk

or call 01732 522444 for more information

"The moment
I picked it up
I knew how
to use it."*^{1,2}

Intuitive design^{1,2}

- Intuitive to use^{1,2}
- Ready in one flip of the cover
- For adult asthma and COPD^{**3}



**DuoResp[®]
Spiromax[®]**
budesonide/formoterol



**MEDICAL
DESIGN
EXCELLENCE
AWARDS[®]**
2015 SILVER WINNER
Drug-Delivery Devices
and Combination Products

Please refer to the Summary of Product Characteristics (SmPC) for full details of the Prescribing Information. DuoResp[®] Spiromax[®] (budesonide/formoterol) 160mcg/4.5mcg inhalation powder and DuoResp[®] Spiromax[®] (budesonide/formoterol) 320mcg/9mcg inhalation powder. **Abbreviated Prescribing Information. Presentation:** DuoResp[®] Spiromax[®] 160/4.5: Each delivered dose contains 160mcg of budesonide and 4.5mcg of formoterol fumarate dihydrate. This is equivalent to a metered dose of 200mcg budesonide and 6mcg of formoterol fumarate dihydrate. DuoResp[®] Spiromax[®] 320/9: Each delivered dose contains 320mcg of budesonide and 9mcg of formoterol fumarate dihydrate. This is equivalent to a metered dose of 400mcg budesonide and 12mcg of formoterol fumarate dihydrate. Inhalation powder. **Indications: Asthma:** Treatment of asthma, where use of a combination (inhaled corticosteroid and long-acting β_2 -adrenoceptor agonist) is appropriate. COPD: Symptomatic treatment of patients with severe COPD (FEV₁ <50% predicted normal) and a history of repeated exacerbations, who have significant symptoms despite regular therapy with long-acting bronchodilators. **Dosage and administration:** For use in adults ≥ 18 years. Not for use in children <18 years of age. **Asthma:** Not intended for the initial management. If an individual patient should require a combination of doses other than those available in the combination inhaler, appropriate doses of β_2 -adrenoceptor agonists and/or corticosteroids by individual inhalers should be prescribed. The dose should be titrated to the lowest dose at which effective control of symptoms is maintained. When control of symptoms is achieved titrate to the lowest effective dose, which could include once daily dosing. DuoResp[®] Spiromax[®] 160/4.5: maintenance therapy – regular maintenance treatment with a separate reliever inhaler. Adults: 1-2 inhalations twice daily (maximum of 4 inhalations twice daily). DuoResp[®] Spiromax[®] maintenance and reliever therapy – regular maintenance treatment and as needed in response to symptoms: should be considered for patients with: (i) inadequate asthma control and in frequent need of reliever medication (ii) previous asthma exacerbations requiring medical intervention. Adults: The

recommended maintenance dose is 2 inhalations per day, given either as one inhalation morning and evening or as 2 inhalations in either the morning or evening. For some patients a maintenance dose of 2 inhalations twice daily may be appropriate. Patients should take 1 additional inhalation as needed in response to symptoms. If symptoms persist after a few minutes, an additional inhalation should be taken. Not more than 6 inhalations should be taken on any single occasion. A total daily dose of up to 12 inhalations could be used for a limited period. Patients using more than 8 inhalations daily should be strongly recommended to seek medical advice. DuoResp[®] Spiromax[®] 320/9: Only to be used as maintenance therapy. Adults: 1 inhalation twice daily (maximum of 2 inhalations twice daily). COPD: Adults: 1 inhalation twice daily. Elderly patients (≥ 65 years old): No special requirements. Patients with renal or hepatic impairment: No data available. **Contraindications:** Hypersensitivity to the active substance or to any of the excipients. **Precautions and warnings:** If treatment is ineffective, or exceeds the highest recommended dose, medical attention must be sought. Patients with sudden and progressive deterioration in control of asthma or COPD should undergo urgent medical assessment. Patients should have their rescue inhaler available at all times. The reliever inhalations should be taken in response to symptoms and are not intended for regular prophylactic use e.g. before exercise. For such, a separate rapid-acting bronchodilator should be considered. Patients should not be initiated during an exacerbation. Serious asthma-related adverse events and exacerbations may occur. If asthma symptoms remain uncontrolled or worsen, patients should continue treatment and seek medical advice. If paradoxical bronchospasm occurs, treatment should be discontinued immediately. Paradoxical bronchospasm responds to a rapid-acting inhaled bronchodilator and should be treated straightaway. Systemic effects may occur, particularly at high doses prescribed for long periods. Potential effects on bone density should be considered, particularly in patients on high doses for prolonged periods that have co-existing risk factors for osteoporosis. Prolonged treatment with high doses of inhaled corticosteroids may result in clinically

significant adrenal suppression. Additional systemic corticosteroid cover should be considered during periods of stress. Treatment should not be stopped abruptly. Transfer from oral steroid therapy to a budesonide/formoterol fumarate fixed-dose combination may result in the appearance of allergic or arthritic symptoms which will require treatment. In rare cases, tiredness, headache, nausea and vomiting can occur due to insufficient glucocorticosteroid effect and temporary increase in the dose of oral glucocorticosteroids may be necessary. To minimise risk of oropharyngeal Candida infection patients should rinse mouth with water. Administer with caution in patients with thyrotoxicosis, phaeochromocytoma, diabetes mellitus, untreated hypokalaemia, or severe cardiovascular disorders. The need for, and dose of inhaled corticosteroids should be re-evaluated in patients with active or quiescent pulmonary tuberculosis, fungal and viral infections in the airways. Additional blood glucose controls should be considered in diabetic patients. Hypokalaemia may occur at high doses. Particular caution is recommended in unstable or acute severe asthma. Serum potassium levels should be monitored in these patients. As with other lactose containing products the small amounts of milk proteins present may cause allergic reactions. **Interactions:** Concomitant treatment with potent CYP3A4 inhibitors should be avoided. If this is not possible the time interval between administration should be as long as possible. Not recommended with β -adrenoceptor blockers (including eye drops) unless compelling reasons. Concomitant treatment with quinidine, disopyramide, procainamide, phenothiazines, antihistamines (terfenadine), Monoamine Oxidase Inhibitors (MAOIs) and Tricyclic Antidepressants (TCAs) can prolong the QTc-interval and increase the risk of ventricular arrhythmias. L-Dopa, L-thyroxine, oxycotin and alcohol can impair cardiac tolerance. Concomitant treatment with MAOIs, including agents with similar properties, may precipitate hypertensive reactions. Patients receiving anaesthesia with halogenated hydrocarbons have an elevated risk of arrhythmias. Hypokalaemia may increase the disposition towards arrhythmias in patients taking digitalis glycosides. **Pregnancy and lactation:** Use only when benefits outweigh potential risks. Budesonide is excreted in breast milk;

at therapeutic doses no effects on infants are anticipated. **Effects on ability to drive and use machines:** No or negligible influence. **Adverse reactions:** Since DuoResp[®] Spiromax[®] contains both budesonide and formoterol, the same pattern of adverse reactions as reported for these substances may occur. No increased incidence of adverse reactions has been seen following concurrent administration of the two compounds. **Serious:** Immediate and delayed hypersensitivity reactions, e.g. exanthema, urticaria, pruritus, dermatitis, angioedema and anaphylactic reaction, Cushing's syndrome, adrenal suppression, growth retardation, decrease in bone mineral density, hypokalaemia, hyperglycaemia, aggression, psychomotor hyperactivity, anxiety, sleep disorders, depression, behavioural changes, cataract and glaucoma, tachycardia, cardiac arrhythmias, e.g. atrial fibrillation, supraventricular tachycardia and extrasystoles, angina pectoris, prolongation of QTc-interval, variations in blood pressure, bronchospasm and paradoxical bronchospasm. **Common:** Candida infections in the oropharynx, headache, tremor, palpitations, mild irritation in the throat, coughing and hoarseness. Consult the Summary of Product Characteristics in relation to other side effects. **Overdose:** An overdose of formoterol may lead to: tremor, headache, palpitations. Symptoms reported from isolated cases are tachycardia, hyperglycaemia, hypokalaemia, prolonged QTc-interval, arrhythmia, nausea and vomiting. Supportive and symptomatic treatment may be indicated. **Price per pack:** DuoResp[®] Spiromax[®] 160/4.5 and DuoResp[®] Spiromax[®] 320/9: £29.97. **Legal Category:** POM. **Marketing Authorisation Numbers:** DuoResp[®] Spiromax[®] 160/4.5: EU/1/14/920/001. DuoResp[®] Spiromax[®] 320/9: EU/1/14/920/004. **Marketing Authorisation Holder:** Teva Pharma BV, Computerweg 10, 3542 DR Utrecht, The Netherlands. **Date of Preparation:** May 2014. **Job Code:** UK/MED/14/0019. **References:** 1. Kychik R, Kreimendahl F. Presented at the 7th IPCRG World Conference, 2014. 2. Plusa T, Bijou P. *Int Rev Allergol Clin Immunol Family Med*, 2015; 21(1): 21-24. 3. DuoResp Spiromax[®] Summary of Product Characteristics.

Adverse events should be reported. Reporting forms and information can be found at www.mhra.gov.uk/yellowcard. Adverse events should also be reported to Teva UK Limited on 0207 540 7117 or medinfo@teva.uk

*Instructions for use should be followed as per the patient information leaflet.

**DuoResp Spiromax[®] is licensed for use in adults 18 years of age and older only.

Approval code: UK/DUO/15/0007(1)0

Date of preparation: August 2015