

BMJ Open
Respiratory
Research

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

Jonathan Cooke,^{1,2} Christopher Butler,^{3,4} Rogier Hopstaken,⁵ Matthew Scott Dryden,⁶ Clodna McNulty,⁷ Simon Hurding,^{8,9} Michael Moore,¹⁰ David Martin Livermore^{11,12}

To cite: Cooke J, Butler C, Hopstaken R, *et al*. Narrative review of primary care point-of-care testing (POCT) and antibacterial use in respiratory tract infection (RTI). *BMJ Open Res* 2015;**2**: e000086. doi:10.1136/bmjresp-2015-000086

Received 3 March 2015
Revised 14 April 2015
Accepted 15 April 2015

ABSTRACT

Antimicrobial resistance is a global problem and is being addressed through national strategies to improve diagnostics, develop new antimicrobials and promote antimicrobial stewardship. A 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 that were included were 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 point-of-care testing (POCT), but has shown value for patients with RTI admitted to hospital. GPs and patients report a good acceptability for a CRP POCT and economic evaluations show cost-effectiveness of CRP POCT over existing RTI management in primary care. POCTs increase diagnostic precision for GPs in the better management of patients with RTI. CRP POCT can better target 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 POCT to be implemented.

BACKGROUND

Antimicrobial resistance (AMR) is a global healthcare and economic problem, and should be a priority for all the World's governments.¹ The Director General of the WHO, Margaret Chan, has stated with respect to AMR that—'No action today means no cure tomorrow'.² The UK Chief Medical Officer has, likewise, repeatedly emphasised the threat of AMR³ and the recently published UK Strategy for AMR has three aims: ¹ improving knowledge and understanding of AMR; ² conserving and stewarding the effectiveness of existing treatments; and ³ stimulating the development of new antibiotics, diagnostics and novel therapies.⁴ Substantial new or redeployed funding

will be needed in order to deliver this strategy, as the amount spent on research in this area is relatively small.⁵ The UK public has recognised the importance of AMR by voting for the challenge of the Longitude Prize 2014 to be to create "a cheap, accurate, rapid and easy-to-use point-of-care diagnostic test for bacterial infections".⁶ The aim is to find the test that will have the greatest impact globally on antibiotic resistance and the prize will be open until 2019. <http://www.longitudeprize.org/challenge/antibiotics>.

Community prescribing of antibacterial medication varies considerably across European countries and in general practice there are individual community and country-level associations in the prescribing of antibacterials—the prevalence of resistance underscoring the need to avoid unnecessary prescription many of which are for respiratory tract infections (RTIs) in the community.⁷ RTIs are the most frequent infections encountered in primary care and most people presenting to a GP with an acute uncomplicated RTI still receive an antibiotic prescription, with many doctors and patients believing that this is the 'right thing' to do.⁸ This is despite the facts (1) that most RTIs are viral, (2) that there is only limited evidence to support the use of antibiotics in acute bronchitis, sore throat, sinusitis and otitis media^{9–12} and (3) the evidence from systematic reviews and other studies suggests little, if any, benefit is achieved from the prescription of antibiotics, except in elderly patients at high risk of pneumonia.^{9 13–15} A recent study of GP prescribing in England found that the likelihood of GPs prescribing antibiotics for coughs and colds increased by 40% between 1999 and 2011, despite Government recommendations to reduce prescribing for illnesses largely caused by viruses.¹⁶ Clarity of diagnosis was questioned in a cross-sectional study, which showed there were considerable differences in GPs'



CrossMark

For numbered affiliations see end of article.

Correspondence to

Jonathan Cooke;
j.cooke@imperial.ac.uk

diagnosis of pneumonia between Denmark and Spain;¹⁷ moreover, there is evidence from diagnostic studies to show the poor accuracy of clinical diagnosis of pneumonia in general practice (with radiographic proof as reference standard).^{18–20}

So, how can the prescribing of antibiotics for RTIs in primary care be made more appropriate? How might improvements in diagnostic testing promote this?

POINT-OF-CARE TESTS

Potentially, primary care prescribers can use point-of-care testing (POCTs) to inform their management of disease, especially if these tests can be performed within the duration of a patient visit, with results obtained within 5 min of taking the sample. If sufficiently sensitive and specific, these tests can offer objective precision to clinical assessment of the patient's signs and symptoms.

A recent survey of Dutch GPs reported the most common point-of-care tests currently used by family physicians were: blood glucose (96%), urine leucocytes or nitrite (96%), urine pregnancy (94%), haemoglobin (58%), and CRP (48%). The most commonly desired point-of-care tests were: D-dimer (70%), troponin (65%), brain natriuretic peptide (BNP; 62%), chlamydia (60%) and International Normalised Ratio (INR; 54%). Family practitioners expected point-of-care tests to have a positive effect on patient satisfaction (93%), diagnostic certainty (89%), antibiotics use (84%) and substitution to primary care (78%). They considered the proven effect on clinical management (46%), and the tests' reliability (35%) to be important aspects of point-of-care tests. Respondents wanted point-of-care tests to help them diagnose acute conditions, such as acute thromboembolic disorders (D-dimers), cardiac disorders (troponin, BNP) and infections (CRP, chlamydia).²¹

However, are there any useful POCTs to help predict RTI in general practice and to reduce unnecessary antibiotic prescribing, as advocated in the UK AMR Strategy?

NARRATIVE REVIEW

Aim

To consider whether POCTs improve antibacterial prescribing, whether they are acceptable for patients and GPs, and their cost-effectiveness for the National Health Service (NHS).

Design and setting

Narrative review of the literature on POCT and antibacterial use in RTI in a primary care setting.

Method

A literature search was undertaken to review the literature on biomarker Point-of-Care Tests (POCT) for adults presenting to GPs with symptoms of RTI in order to:

- Determine whether POCTs can reduce antibacterial prescribing;

- Ascertain the safety and acceptability of POCTs for patients and GPs;
- Determine the cost-effectiveness of POCTs in an NHS setting;

Papers reviewed were in English, and published between 1995 and 2015.

Search terms were: *CRP, C-reactive protein, biomarkers, procalcitonin, infections, antibiotics, antimicrobials, primary care, point of care testing, infection respiratory tract.*

Databases searched were: EMBASE, Excerpta Medica (Ovid), Journals@Ovid Full Text (Ovid), PubMed, MEDLINE (Ovid).

Highwire Press, nature.com, ScienceDirect—All Content—V.4 (Elsevier), SpringerLink, Wiley Interscience Journals, NHS Evidence, The Cochrane Collaboration.

RESULTS

The first search yielded 1022 references. This was further refined down to 76 references by focusing on systematic reviews, clinical trials and economic analyses in human adults, and in RTIs presenting in primary care. Excluded studies included laboratory studies, studies in children, duplications and non-systematic reviews.

The pivotal studies of randomised controlled trials, cluster randomised controlled trials, economic evaluations and systematic reviews are shown in [table 1](#).

Measurement of C reactive protein (CRP) and procalcitonin biomarkers have been advocated as additional tests to assist proper clinical examination, especially (1) where there is a high degree of diagnostic uncertainty, (2) for patients who are very worried and/or demanding an antibiotic, and (3) to differentiate the seriously ill from the non-seriously ill. Procalcitonin has greater sensitivity and specificity for distinguishing bacterial and viral infection in hospital settings, and multiple analyses point to its value in guiding hospital antibiotic use for patients with pneumonia, leading to reductions in antibiotic use without harm to patients.^{22–23} However, studies show that procalcitonin does not yet add sufficient value for decision-making in primary care,^{24–26} whereas CRP POCT appears better in correctly predicting the absence of radiographic pneumonia.²⁴ Moreover, procalcitonin has not yet proven to be suitable for deployment as a POCT in general practice, having a turnaround time (depending on the system) of 18–30 min; CRP POCTs, by contrast, have a turnaround of <5 min, thus giving a result within the ambit of a standard NHS GP consultation.

CRP Background

CRP is a major acute-phase plasma protein which is produced in response to infection or tissue injury. It has Ca²⁺-dependent binding specificity for phosphocholine (PCh), a constituent of many bacterial and fungal polysaccharides, and of most biological cell membranes. The main biological function of CRP is determined by this ability to recognise pathogens and damaged host cells,

Table 1 Studies that examine antibiotic prescribing for patients presenting with symptoms of RTI to GPs who use POCT biomarkers compared with normal care

Author and date	Patient group	Study type	Outcomes	Key results
Diederichsen <i>et al</i> 2000 ³⁴	35 general practices, County of Funen, Denmark. PATIENTS: 812 patients with respiratory infection	Randomised controlled trial	Frequency of antibiotic prescriptions and morbidity 1 week after the consultation, as stated by the patients	The frequency of antibiotic prescriptions was 43% (179/414) in a CRP group compared with 46% (184/398) in the control group (OR=0.9, NS)
Cals <i>et al</i> 2009 ⁴⁰	40 general practitioners from 20 practices in the Netherlands recruited 431 patients with lower respiratory tract infection	Cluster randomised controlled trial	Main outcome measures The primary outcome was antibiotic prescribing at the index consultation	General practitioners in a CRP test group prescribed antibiotics to 31% of patients compared with 53% in the no test group (42% lower $p=0.02$). Patients' recovery and satisfaction were similar in both study groups
Cals <i>et al</i> 2010 ³³	258 patients were enrolled (107 LRTI and 151 rhinosinusitis) by 32 family physicians in the Netherlands	Randomised controlled trial	Antibiotic prescribing rate, recovery and patient satisfaction	Patients in a CRP-assisted group were prescribed fewer antibiotics (43.4%) than control patients (56.6%; 23.5% lower) after the index consultation (relative risk (RR) =0.77; 95% CI 0.56 to 0.98). Recovery was similar across groups. Satisfaction with care was higher in patients managed with CRP assistance ($p=0.03$)
Little <i>et al</i> 2013 ¹⁴	Patients presenting with upper or lower RTI in primary-care practices in six European countries	Cluster randomised trial	Antibiotic prescribing rate	GP antibiotic prescribing was lower with CRP training than without (33% vs 48%—31% lower, adjusted risk ratio 0.54, 95% CI 0.42 to 0.69) and with enhanced-communication training than without (36% vs 45%, 0.69, 0.54–0.87). The combined intervention was associated with the greatest reduction in prescribing rate (CRP risk ratio 0.53, 95% CI 0.36 to 0.74, $p<0.0001$; enhanced communication 0.68, 0.50–0.89, $p=0.003$; combined 0.38, 0.25–0.55, $p<0.0001$)
Andreeva and Melbye 2014 ³⁰	179 patients with acute cough/LRTI (including acute bronchitis, pneumonia and infectious exacerbations of COPD or asthma) from 18 Russian GP practices	Open cluster randomised clinical trial	Antibiotic prescribing rates, referral for chest X-ray and recovery rate	The antibiotic prescribing rate was 37.6% in the CRP group, which was significantly lower than that in the control group (58.9%; $p=0.006$). Referral for chest X-ray was also significantly lower in the intervention group (55.4%) than in the control group (75.6%) ($p=0.004$). The recovery rate, as recorded by the GPs, was 92.9% and 93.6% in the intervention and control groups, respectively
Systematic reviews Huang <i>et al</i> 2013 ⁵⁵	13 studies in 10 005 patients presenting to GPs with RTI	Systematic review and meta-analysis	Association between point-of-care CRP testing and antibiotic prescribing in respiratory tract infections	POC CRP testing was associated with a significant reduction in antibiotic prescribing at the index consultation (RR 0.75, 95% CI 0.67 to 0.83)

Continued

Table 1 Continued

Author and date	Patient group	Study type	Outcomes	Key results
Aabenhus <i>et al</i> 2014 ³¹	6 trials (3284 participants; 139 children) Patients presenting with symptoms of acute respiratory infections in primary care	Systematic review (Cochrane Review)	To assess the benefits and harms of point-of-care biomarker tests of infection to guide antibiotic treatment in patients in primary care presenting with symptoms of RTI	Reduction in the use of antibiotic treatments was found in studies comparing C reactive protein (631/1685) with standard of care (785/1599)
Economic evaluations				
Cals <i>et al</i> 2011 ⁴⁶	Cost-effectiveness analysis with a time horizon of 28 days in 431 patients with LRTIs recruited by 40 GPs	Economic analysis alongside a factorial, cluster randomised trial	Healthcare costs. Cost-effectiveness, using the primary outcome measure antibiotic prescribing at index consultation, was assessed by incremental cost-effectiveness ratios (ICER)	The total mean cost per patient in the usual care group was €35.96 with antibiotic prescribing of 68%, €37.58 per patient managed by GPs using CRP tests (antibiotic prescribing 39%, ICER €5.79), €25.61 per patient managed by GPs trained in enhanced communication skills (antibiotic prescribing 33%, dominant) and €37.78 per patient managed by GPs using both interventions (antibiotic prescribing 23%, ICER €4.15)
Oppong 2013 ⁴⁷	Patients with acute cough and LRTI in primary care settings in Norway and Sweden	Economic analysis alongside an observational study	Antibiotic use, cost, and patient outcomes (symptom severity after 7 and 14 days, time to recovery, and EQ-5D)	POCCRP testing was associated with a cost per quality-adjusted life year (QALY) gain of €9391. At a willingness-to-pay threshold of €30 000 per QALY gained, there is a 70% probability of CRP being cost-effective
Hunter 2015 ⁴⁸	Three different strategies of CRP testing (GP plus CRP; practice nurse plus CRP; and GP plus CRP and communication training) for patients with RTI symptoms compared with current standard GP practice without CRP testing	Economic analysis using decision tree and Markov model	Quality-adjusted life years (QALYs) and cost per 100 patients, together with the number of antibiotic prescriptions and RTIs for each group	Compared with current standard practice, the GP plus CRP and practice nurse plus CRP test strategies result in increased QALYs and reduced costs, while the GP plus CRP testing and communication training strategy is associated with increased costs and reduced QALYs. Additionally, all three CRP arms led to fewer antibiotic prescriptions and infections over 3 years

CRP, C reactive protein; COPD, chronic obstructive pulmonary disease; LRTI, lower respiratory tract infection; RTI, respiratory tract infection.

and to mediate their elimination by recruiting the complement system along with phagocytic cells. PCh, the principal CRP ligand, is widely distributed in the teichoic acids, capsular carbohydrates, and lipopolysaccharides of bacteria and other microorganisms.²⁷ CRP was discovered and named because of its reactivity with the PCh residues of Cpolysaccharide (PnC), the teichoic acid of *Streptococcus pneumoniae*.²⁸ CRP is synthesised by the liver in response to factors released by macrophages and fat cells (adipocytes). Like many acute phase proteins, CRP is normally present at trace levels in serum, but increases rapidly and dramatically in response to a variety of infectious or inflammatory stimuli.²⁹ Its levels typically are highest in patients with a bacterial infection and are lower in those with viral infection. Rapid tests for CRP were introduced into general practice about 20 years ago and are widely used as a diagnostic in the Netherlands and Nordic countries, mostly for RTI.³⁰

CRP evidence

A recent Cochrane review concluded that a point-of-care biomarker (eg, CRP) to guide antibiotic treatment of acute respiratory infections (ARIs) in primary care can significantly reduce antibiotic use, [figure 1](#).³¹ Studies included in the final analysis included randomised and cluster randomised controlled trials in England, Wales, the Netherlands, Poland, Spain, Belgium, Denmark and Russia to demonstrate that using CRP POCT in primary care can significantly reduce the initial prescribing rate of antibiotics. Reported reductions were 36.16% (37.6% vs 58.9%) in patients presenting with acute cough/lower respiratory tract infection, LRTI (including acute bronchitis, pneumonia and infectious exacerbations of chronic obstructive pulmonary disease (COPD) or asthma),³⁰ 31.25% (33% vs 48%) in upper and lower

RTI,¹⁴ 24.4% (59% vs 78%) in sinusitis,³² 23.3% (43.4% vs 56.6%) in LRTI or rhinosinusitis³³ and 6.5% (43% vs 46%) in respiratory infections.³⁴ In an uncontrolled observational study, Swiss GPs who used CRP as part of a clinical assessment of patients presenting with acute cough were observed to prescribe antibiotics in 22% of consultations which is considerably lower than studies from other countries.³⁵ These studies, together with systematic reviews, offer guidance on the value of measured levels of CRP as both positive and negative prognostic indicators of whether to administer antibiotics or not.^{36 37} Interpretive and prescribing criteria, have been developed from a consensus of collaborators in two consortia: the 'Improving Management of Patients with Acute cough by C-Reactive Protein Point of Care Testing and Communication Training (IMPACT) Programme' and the Genomics to Combat Resistance against Antibiotics in Community-acquired LRTI in Europe (GRACE) consortium.^{14 38} These groups propose:

- ▶ CRP levels <20 mg/L—Self-limiting LRTI. Withhold antibiotics
- ▶ CRP 21–50 mg/L—Most patients have self-limiting LRTI. Assessment of signs, symptoms, risk factors and CRP is important. Withhold antibiotics in most cases.
- ▶ CRP 51–99 mg/L—Assessment of signs, symptoms, risk factors and CRP is crucial. Withhold antibiotics in the majority of cases and consider delayed antibiotics in the minority of cases.
- ▶ CRP >100 mg/L—Severe infection. Prescribe antibiotics¹⁴

Similar criteria have been proposed by other groups, including the UK National Institute for Health and Care Excellence (NICE). These are given below and compared in [table 2](#).

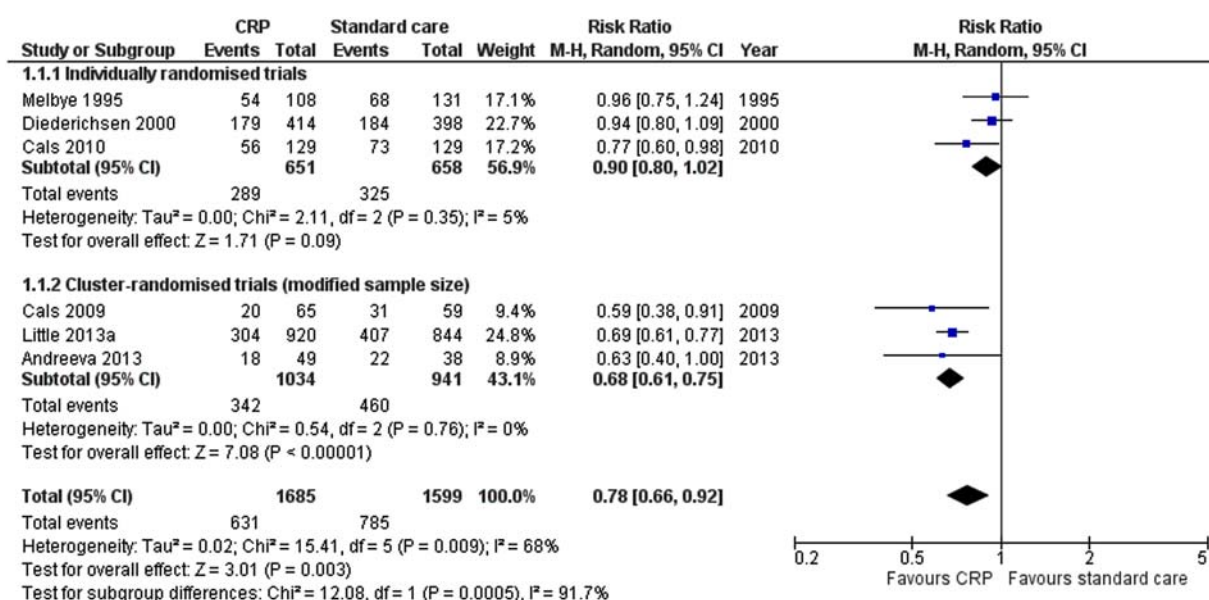


Figure 1 Forest plots for randomised and cluster randomised controlled trials from Cochrane review Biomarkers as point-of-care tests to guide prescription of antibiotics in patients with acute respiratory infections in primary care.³¹

Table 2 Criteria for management of RTIs in general practice with CRP POCT after proper clinical examination of the patient

Management	Draft NICE guidance for Pneumonia ⁴⁹	Dutch GP practice guideline ⁵⁴	GRACE study ¹⁴	ERS ⁵¹
Self-limiting RTI. Do not routinely offer antibiotic therapy. Pneumonia unlikely. Give education Majority of patients have self-limiting LRTI. Assessment of signs, symptoms, risk factors and CRP is important. Withhold antibiotics, in most cases Assessment of signs, symptoms, risk factors and CRP is crucial. Withhold antibiotics in the majority of cases and consider delayed antibiotics in the minority of cases	CRP less than 20 mg/L	CRP less than 20 mg/L	CRP less than 20 mg/L	CRP less than 20 mg/L
Consider a delayed antibiotic prescription. Clinical presentation decisive. Prescribe antibiotics only in patients with a high risk of complications*	CRP between 20 and 100 mg/L	CRP between 20 and 100 mg/L	CRP 21–50 mg/L	
Severe infection. High risk of pneumonia. Offer antibiotic therapy	CRP greater than 100 mg/L	CRP greater than 100 mg/L	CRP 51–99 mg/L	CRP greater than 100 mg/L

*Complicated respiratory tract infection.

A complicated respiratory tract infection is an infection with an increased risk of a complicated course (mortality or hospital admission). Two groups are distinguished.

1. Patients with a probable diagnosis of pneumonia based on:

► Acute cough AND:

- Being severely ill, with for example, tachypnoea, tachycardia, hypotension (systolic blood pressure <90, diastolic blood pressure <60 mm Hg) or confusion;
- Being moderately ill and unilateral auscultatory abnormalities (however, the absence of auscultatory abnormalities does not rule out pneumonia);
- An infiltrate on chest X-ray;
- A course >7 days with fever and cough (without abnormalities on physical examination).
- For values between 20 and 100 mg/L, the clinical presentation determines policy, along with risk factors such as heart failure, diabetes mellitus, COPD, asthma, age <3 months or >75 years;

2. Patients with another risk factor for a complicated course.

► Other risk factors—particularly age and comorbidities—should also be included in the evaluation of patients with acute cough. The following factors increase risk of hospital admission and mortality:

- Age <3 months or >75 years;
- In children, cardiovascular and pulmonary conditions (except asthma);
- In adults: heart failure, severe COPD, diabetes mellitus (esp. with use of insulin), neurological conditions, severe renal insufficiency;
- A disrupted immune system (owing to oncological conditions, renal insufficiency).⁵⁴

CRP, C reactive protein; COPD, chronic obstructive pulmonary disease; POCT, point-of care-test; RTI, respiratory tract infection; NICE, National Institute for Health and Care Excellence.

Both general practitioners (GPs) and patients appear to find using the test acceptable, with one Dutch study showing CRP POCT to have little effect on GP workload in 50% of practices.³⁹ Patients were satisfied to be provided with the results of a reassuringly low CRP POCT rather than receiving an antibiotic prescription.^{39–41} CRP testing has a role as an adjunct in effective communication with patients; clinicians and patients all recommend seeing the CRP in conjunction with the overall assessment and caution against over reliance on CRP results in isolation of clinical assessment.^{42–43}

In elderly patients with bacterial infections, CRP POCT reportedly had a sensitivity of diagnosing bacterial infection of 80.7% and a specificity of 96%, with a positive predictive value 91.9% and negative predictive value 89.8%.⁴⁴

While most of these results are promising, a few studies have sounded a note of caution, particularly on the ability of CRP to distinguish between bacterial and viral infection. One Dutch study of LRTIs found that a CRP >20 mg/L (OR 2.1–4.6), along with an erythrocyte sedimentation rate >50 (OR 2.3–3.3), were independent

predictors for viral or bacterial LRTI, but could not delineate between these two aetiologies.⁴⁵

Is withholding antibiotics based on CRP POCT safe?

In a Russian study of patients with acute cough/LRTI (including acute bronchitis, pneumonia and infectious exacerbations of COPD or asthma), participants were randomised by GPs to normal care or to care informed by a CRP POCT. As well as a significant reduction in antibiotic prescribing, the referral rate for chest radiography was significantly lower in the intervention group (55.4%) as compared to the control group (76%; $p=0.004$).³⁰ Another Dutch study showed that GPs' use of CRP POCT significantly reduced antibiotic prescribing for LRTIs, without compromising patients' recovery and satisfaction with care.⁴⁰ This study also enhanced general consultation skills and showed that proper discussion of the test results with the patient offered an important contribution to better antibiotic stewardship.

While near patient tests can add diagnostic precision to a standard clinical examination, these are not 100% reliable.²⁴ Any strategy reducing antibiotic use may have rare adverse consequences due to untreated infection. Use of the test is not a substitute for a proper clinical examination and 'safety net' advice for those not recovering as expected.

Economic evaluations of CRP POCTs

In an observational study of the presentation, management, and outcomes of patients with acute cough and LRTI in primary care settings in the Netherlands, the total mean cost per patient in the 'usual care' group was €35.96, with antibiotics prescribed to 68% of patients; €37.58 per patient managed by GPs using CRP POCT (antibiotic prescribing, 39%); €25.61 per patient managed by GPs trained in enhanced communication skills (antibiotic prescribing 33%); and €37.78 per inpatient managed by GPs using both interventions (antibiotic prescribing, 23%). It is less clear how such outcomes translate between national healthcare systems varying in consultation format, and in the choice and cost of the particular antibiotics prescribed.⁴⁶ In a primary care study undertaken in Norway and Sweden, CRP POCT testing was associated with non-significant reductions in antibiotic prescribing ($p=0.078$) and increased cost ($p=0.092$). Despite this, CRP POCT was also associated with a cost per quality-adjusted life year (QALY) gain of £9391. At a Willingness-to-Pay threshold of £30 000 per QALY gained, there is a 70% probability of CRP POCT being cost-effective.⁴⁷

Finally, in a modelling study undertaken in England, 3 models of CRP POCT use for patients presenting with RTI symptoms were subjected to Markov analysis: GP plus CRP, Practice nurse plus CRP POCT and GP plus CRP POCT and communication training. All these models were found to be more cost-effective than not using any CRP POCT.⁴⁸

Limitations of study

These include, the heterogeneity of presenting conditions of patients for both upper and lower RTIs in the studies evaluated and the different healthcare systems that provide primary care services in different countries. However, these limitations are somewhat ameliorated by the designs of the studies selected—randomised controlled and cluster randomised controlled trials.

Clinical guidelines for use of CRP in RTI investigation for GPs

National Institute of Health and Care Excellence

Current *draft* clinical guidelines for pneumonia from the NICE propose a number of actions to help GPs to manage patients who present with symptoms of RTI.⁴⁹ These include:

Consider a point-of-care C-reactive protein test for patients presenting with lower respiratory tract infection in primary care if it is not clear after clinical assessment whether antibiotics should be prescribed. Use the results of the C-reactive protein test to guide antibiotic prescribing as follows:

Do not routinely offer antibiotic therapy if the C-reactive protein concentration is less than 20 mg/L.

Consider a delayed antibiotic prescription (a prescription for use at a later date if symptoms worsen) if the C-reactive protein concentration is between 20 mg/L and 100 mg/L.

Offer antibiotic therapy if the C-reactive protein concentration is greater than 100 mg/L.

These NICE guidelines were developed following a systematic review and meta-analysis of the clinical and cost-effective published evidence from randomised controlled trials. The stratification differs from the advice of the IMPAC3T and GRACE consortia (above) only in not subdividing the patients with CRP concentrations between 20 and 100 mg/L.

Public Health England

Current guidance from Public Health England offers strategies for the management of acute upper and lower RTIs in general practice, including community-acquired pneumonia.⁵⁰ This guidance is expected to be updated in line with the NICE guidance, given above, including the use of CRP as a biomarker.

European guidance

The European Respiratory Society has produced Guidelines for the management of adult lower RTI which include the recommendation:

In patients with a suspected pneumonia, a test for serum-level of C-reactive protein (CRP) can be done. A CRP level of <20 mg/L at presentation, with symptoms for

>24 h, makes the presence of pneumonia highly unlikely, a level of >100 mg/L makes pneumonia likely.⁵¹

Table 2 compares the CRP-guided management of RTI based on the various slightly differing sets of guideline criteria outlined above.

Which countries use CRP POCT as a prognostic tool in general practice?

Countries that use CRP POCT widely include: Denmark, Norway, Sweden, Germany, the Netherlands, Switzerland and Finland; countries that use CRP POCT to some extent include: Estonia, Slovenia, Latvia, Czech Republic, Hungary and Austria.⁵² Interestingly, most of these countries have lower systemic antibacterial consumption in the community than does the UK.⁵³ The Netherlands has particularly addressed antimicrobial

stewardship in primary care, and has one of the lowest ambulatory antibiotic prescribing rates for humans in Europe⁵³ (figure 2). Its national guidelines (NHG Guidelines)⁵⁴ resemble those of GRACE and IMPACT, and the draft NICE guidance, summarised in table 2.

Making comparisons between countries is not complete without considering how different countries pay for POCTs. In the UK, GPs would have to meet the additional costs of CRP POCT in order to encourage uptake of this technology; the alternative funding mechanisms would need to be identified, for instance, by using 'enhanced service' contracting arrangements.

CONCLUSION

There is considerable and accumulating evidence that measurements of CRP and procalcitonin are both

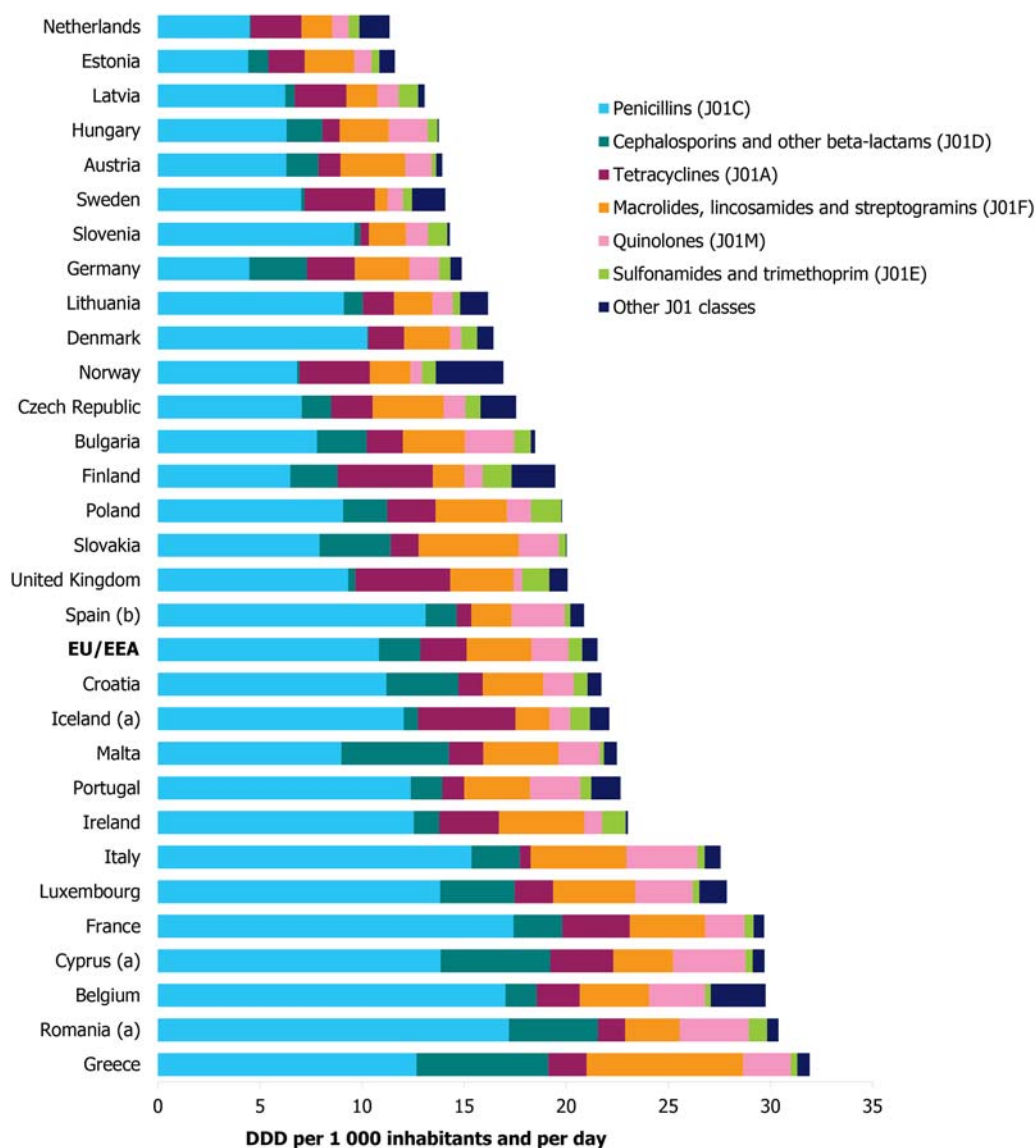


Figure 2 Consumption of antibacterials for systemic use (ATC group J01) at anatomical therapeutic chemical (ATC) group level 3 in the community, European Union/European Economic Area countries, 2012, expressed as defined daily dose (DDD) per 1000 inhabitants and per day. From European Centre for Disease Prevention and Control.⁵³

clinically valuable and cost-effective diagnostics. Procalcitonin is not currently available as a POCT but has shown value in patients admitted to hospital with RTI. CRP POCT offers GPs a simple test that can be performed within 5 min and this helps to distinguish whether community patients with RTI need antibiotics or not. The test is acceptable to both patients and GPs, and the consequential reduction in the prescribing of antibiotics will contribute to global strategies in promoting the better stewardship of antimicrobials.

Author affiliations

¹Division of Infectious Diseases, Department of Medicine, The Centre for Infection Prevention and Management, Imperial College London, London, UK

²Faculty of Medical and Human Sciences, University of Manchester, Manchester, UK

³Nuffield Department of Primary Care Health Sciences, University of Oxford, Oxford, UK

⁴Cochrane Institute of Primary Care & Public Health, School of Medicine, Cardiff University, Cardiff, UK

⁵Saltro Diagnostic Centre for Primary Care, Utrecht, The Netherlands

⁶Department of Microbiology, Hampshire Hospitals NHS Foundation Trust, Winchester, UK

⁷Public Health England, Primary Care Unit, Microbiology Department, Gloucestershire Royal Hospital, Gloucester, UK

⁸Scottish Government, Clinical Lead Therapeutics Branch, Edinburgh, UK

⁹Medicines Management Adviser NHS Lothian, Edinburgh, UK

¹⁰Primary Care and Population Sciences, University of Southampton Aldermore Health Centre, Aldermore Close, UK

¹¹Norwich Medical School, University of East Anglia, Norwich, UK

¹²Antimicrobial Resistance & Healthcare Associated Infections Reference Unit, Public Health England, London, UK

Contributors All authors have contributed to this paper.

Funding This study was supported through an educational grant from Alere International.

Competing interests JC chaired the Antimicrobial Stewardship Subgroup of the Department of Health Advisory Committee on Antimicrobial Resistance and Healthcare Associated Infection (travel expenses only), and has chaired, presented and received honoraria at meetings supported by Astellas, Cubist, Alere and HHI. DML is partly self-employed and consults for numerous pharmaceutical and diagnostic companies, including Achaogen, Adenium, Allegra, Astellas, AstraZeneca, Bayer, Basilea, bioMerieux, bioVersys, Cubist, Curetis, GSK, Longitude, Merck, Meiji Seika, Pfizer, Roche, Tetrphase, VenatoRx and Wockhardt; he holds grants from AstraZeneca, Basilea, Cubist, Meiji Seika, Merck and VenatoRx; has received lecture honoraria or travel reimbursement from AstraZeneca, Curetis, GSK, J&J, Leo, Meiji, Merck, Novartis, Pfizer and Tetrphase, and holds shares in Dechra, GSK, Merck and Pfizer, collectively amounting to <10% of portfolio value.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement No additional data are available.

Open Access This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>

REFERENCES

1. WHO Advisory Group on Integrated Surveillance of Antimicrobial Resistance (AGISAR). Antimicrobial resistance: global report on surveillance 2014 April 2014. 257p.
2. Chan M. *Antimicrobial resistance: no action today, no cure tomorrow. World Health Day 2011 Combat antimicrobial resistance*. Geneva: Director-General of the World Health Organization, 2011.
3. Davies SC. *Annual Report of the Chief Medical Officer. Volume Two, 2011. Infections and the rise of antimicrobial resistance*. London: Department of Health, 2013.
4. Department of Health. *UK 5 year antimicrobial resistance strategy 2013 to 2018*. London: Department of Health 2013.
5. Bragginton EC, Piddock LJV. UK and European Union public and charitable funding from 2008 to 2013 for bacteriology and antibiotic research in the UK: an observational study. *Lancet Infect Dis* 2014;14:857–68.
6. Longitude Prize 2014—Antibiotics 2014. <http://www.longitudeprize.org/>
7. Goossens H, Ferech M, Vander Stichele R, *et al*. Outpatient antibiotic use in Europe and association with resistance: a cross-national database study. *Lancet* 2005;365:579–87.
8. National Institute for Health and Clinical Excellence. Respiratory tract infections—antibiotic prescribing for self-limiting respiratory tract infections in adults and children in primary care. NICE clinical guideline 69, 2008.
9. Smith SM, Fahey T, Smucny J, *et al*. Antibiotics for acute bronchitis. *Cochrane Database Syst Rev* 2014;3:CD000245.
10. Spinks A, Glasziou PP, Del Mar CB. Antibiotics for sore throat. *Cochrane Database Syst Rev* 2013;11:CD000023.
11. Ahovuo-Saloranta A, Rautakorpi UM, Borisenko OV, *et al*. Antibiotics for acute maxillary sinusitis in adults. *Cochrane Database Syst Rev* 2014;2:CD000243.
12. Venekamp RP, Sanders S, Glasziou PP, *et al*. Antibiotics for acute otitis media in children. *Cochrane Database Syst Rev* 2013;1:CD000219.
13. Butler CC, Hood K, Verheij T, *et al*. Variation in antibiotic prescribing and its impact on recovery in patients with acute cough in primary care: prospective study in 13 countries. *BMJ* 2009;338:2242.
14. Little P, Stuart B, Francis N, *et al*. Effects of internet-based training on antibiotic prescribing rates for acute respiratory-tract infections: a multinational, cluster, randomised, factorial, controlled trial. *Lancet* 2013;382:1175–82.
15. Petersen I, Johnson AM, Islam A, *et al*. Protective effect of antibiotics against serious complications of common respiratory tract infections: retrospective cohort study with the UK General Practice Research Database. *BMJ* 2007;335:982.
16. Hawker JI, Smith S, Smith GE, *et al*. Trends in antibiotic prescribing in primary care for clinical syndromes subject to national recommendations to reduce antibiotic resistance, UK 1995–2011: analysis of a large database of primary care consultations. *J Antimicrob Chemother* 2014;69:3423–30.
17. Christensen SF. Marked differences in GPs' diagnosis of pneumonia between Denmark and Spain: a cross-sectional study. *Prim Care Respir J* 2013;22:454–8.
18. Melbye H, Straume B, Aasebo U, *et al*. The diagnosis of adult pneumonia in general practice. The diagnostic value of history, physical examination and some blood tests. *Scand J Prim Health Care* 1988;6:111–17.
19. Hopstaken RM, Muris JW, Knotterus JA, *et al*. Contributions of symptoms, signs, erythrocyte sedimentation rate, and C-reactive protein to a diagnosis of pneumonia in acute lower respiratory tract infection. *Br J Gen Pract* 2003;53:358–64.
20. van Vugt SF, Verheij TJ, de Jong PA, *et al*. Diagnosing pneumonia in patients with acute cough: clinical judgment compared to chest radiography. *Eur Respir J* 2013;42:1076–82.
21. Cals JW, Schols AM, van Weert HC, *et al*. [Point-of-care testing in family practices: present use and need for tests in the future]. *Ned Tijdschr Geneesk* 2014;158:A8210.
22. Simon L, Gauvin F, Amre DK, *et al*. Serum procalcitonin and C-reactive protein levels as markers of bacterial infection: a systematic review and meta-analysis. *Clin Infect Dis* 2004;39:206–17.
23. Saeed K, Dryden M, Bourne S, *et al*. Reduction in antibiotic use through procalcitonin testing in patients in the medical admission unit or intensive care unit with suspicion of infection. *J Hosp Infect* 2011;78:289–92.
24. van Vugt SF, Broekhuizen BD, Lammens C, *et al*. Use of serum C reactive protein and procalcitonin concentrations in addition to symptoms and signs to predict pneumonia in patients presenting to primary care with acute cough: diagnostic study. *BMJ* 2013;346:f2450.
25. Schuetz P, Muller B, Christ-Crain M, *et al*. Procalcitonin to initiate or discontinue antibiotics in acute respiratory tract infections. *Cochrane Database Syst Rev* 2012;9:CD007498.
26. Aabenhus R, Jensen JU. Procalcitonin-guided antibiotic treatment of respiratory tract infections in a primary care setting: are we there yet? *Prim Care Respir J* 2011;20:360–7.

27. Volanakis JE. Human C-reactive protein: expression, structure, and function. *Mol Immunol* 2001;38:189–97.
28. Tillett WS, Francis T. Serological reactions in pneumonia with a non-protein somatic fraction of pneumococcus. *J Exp Med* 1930;52:561–71.
29. Clyne B, Olshaker JS. The C-reactive protein1. *J Emerg Med* 1999;17:1019–25.
30. Andreeva E, Melbye H. Usefulness of C-reactive protein testing in acute cough/respiratory tract infection: an open cluster-randomized clinical trial with C-reactive protein testing in the intervention group. *BMC Fam Pract* 2014;15:80.
31. Aabenhus R, Jensen Jens-Ulrik S, Jørgensen Karsten J, *et al.* Biomarkers as point-of-care tests to guide prescription of antibiotics in patients with acute respiratory infections in primary care. *Cochrane Database Syst Rev* 2014;11:CD010130. <http://onlinelibrary.wiley.com/doi/10.1002/14651858.CD010130.pub2/abstract>
32. Bjerrum L, Gahrn-Hansen B, Munck AP. General practitioners who use CRP have a lower antibiotic prescribing rate to patients with sinusitis—secondary publication. *Ugeskr Laeger* 2005;167:2775–7.
33. Cals JW, Marjolein JC, Schot Sanne AM, *et al.* Point-of-care C-reactive protein testing and antibiotic prescribing for respiratory tract infections: a randomized controlled trial. *Ann Fam Med* 2010;8:124–33.
34. Diederichsen HZ, Skamling M, Diederichsen A, *et al.* Randomised controlled trial of CRP rapid test as a guide to treatment of respiratory infections in general practice. *Scand J Prim Health Care* 2000;18:39–43.
35. Streit S, Frey P, Singer S, *et al.* Clinical and haematological predictors of antibiotic prescribing for acute cough in adults in Swiss practices inverted question mark an observational study. *BMC Fam Pract* 2015;16:15.
36. van der Meer V, Neven AK, van den Broek PJ, *et al.* Diagnostic value of C reactive protein in infections of the lower respiratory tract: systematic review. *BMJ* 2005;331:26.
37. Falk G, Fahey T. C-reactive protein and community-acquired pneumonia in ambulatory care: systematic review of diagnostic accuracy studies. *Fam Pract* 2009;26:10–21.
38. Cals JW, Hopstaken RM, Butler CC, *et al.* Improving management of patients with acute cough by C-reactive protein point of care testing and communication training (IMPAC3T): study protocol of a cluster randomised controlled trial. *BMC Fam Pract* 2007;8:15.
39. Cals JW, Chappin FH, Hopstaken RM, *et al.* C-reactive protein point-of-care testing for lower respiratory tract infections: a qualitative evaluation of experiences by GPs. *Fam Pract* 2010;27:212–18.
40. Cals JW, Butler CC, Hopstaken RM, *et al.* Effect of point of care testing for C reactive protein and training in communication skills on antibiotic use in lower respiratory tract infections: cluster randomised trial. *BMJ* 2009;338:b1374.
41. Kavanagh KE, O'Shea E, Halloran R, *et al.* A pilot study of the use of near-patient C-reactive protein testing in the treatment of adult respiratory tract infections in one Irish general practice. *BMC Fam Pract* 2011;12:93.
42. Butler CC, Simpson S, Wood F. General practitioners' perceptions of introducing near-patient testing for common infections into routine primary care: a qualitative study. *Scand J Prim Health Care* 2008;26:17–21.
43. Wood F, Brookes-Howell L, Hood K, *et al.* A multi-country qualitative study of clinicians' and patients' views on point of care tests for lower respiratory tract infection. *Fam Pract* 2011;28:661–9.
44. Liu A. Serum C-reactive protein as a biomarker for early detection of bacterial infection in the older patient. *Age Ageing* 2010;39:559–65.
45. Hopstaken RM, Stobberingh EE, Knottnerus JA, *et al.* Clinical items not helpful in differentiating viral from bacterial lower respiratory tract infections in general practice. *J Clin Epidemiol* 2005;58:175–83.
46. Cals JW, Ament AJ, Hood K, *et al.* C-reactive protein point of care testing and physician communication skills training for lower respiratory tract infections in general practice: economic evaluation of a cluster randomized trial. *J Eval Clin Pract* 2011;17:1059–69.
47. Oppong R. Cost-effectiveness of point-of-care C-reactive protein testing to inform antibiotic prescribing decisions. *Br J Gen Pract* 2013;63:e465–71.
48. Hunter R. Cost-effectiveness of point-of-care C-reactive protein tests for respiratory tract infection in primary care in England. *Adv Ther* 2015;32:69–85.
49. National Institute for Health and Clinical Excellence (NICE). *Pneumonia: diagnosis and management of community-and hospital-acquired pneumonia in adults (draft guidance)*. Guideline development Group, 2014.
50. Public Health England and the British Infection Association. *Managing common infections: overview of guidance for primary care*. Public Health England and the British Infection Association, 2014.
51. Woodhead M, Blasi F, Ewig S, *et al.* Guidelines for the management of adult lower respiratory tract infections—full version. *Clin Microbiol Infect* 2011;17(Suppl 6):E1–59.
52. Alere. Use of CRP POCT in European countries. 2014.
53. European Centre for Disease Prevention and Control (ECDC). *ECDC Surveillance report. Surveillance of antimicrobial consumption in Europe*. Stockholm:2014.
54. Verlee L, Verheij TJ, Hopstaken RM, *et al.* Summary of NHG practice guideline 'Acute cough'. *Ned Tijdschr Geneeskde* 2012;156:A4188. Complete guideline now available in English.
55. Huang Y, Chen R, Wu T, *et al.* Association between point-of-care CRP testing and antibiotic prescribing in respiratory tract infections: a systematic review and meta-analysis of primary care studies. *Br J Gen Pract* 2013;63:787–94.

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

Jonathan Cooke, Christopher Butler, Rogier Hopstaken, Matthew Scott Dryden, Cliona McNulty, Simon Hurding, Michael Moore and David Martin Livermore

BMJ Open Resp Res 2015 2:
doi: 10.1136/bmjresp-2015-000086

Updated information and services can be found at:
<http://bmjopenrespres.bmj.com/content/2/1/e000086>

These include:

References

This article cites 45 articles, 16 of which you can access for free at:
<http://bmjopenrespres.bmj.com/content/2/1/e000086#BIBL>

Open Access

This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>

Email alerting service

Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
<http://group.bmj.com/group/rights-licensing/permissions>

To order reprints go to:
<http://journals.bmj.com/cgi/reprintform>

To subscribe to BMJ go to:
<http://group.bmj.com/subscribe/>