

Protecting and improving the nation's health

Start Smart - Then Focus Antimicrobial Stewardship Toolkit for English Hospitals

Updated March 2015

About Public Health England

Public Health England exists to protect and improve the nation's health and wellbeing, and reduce health inequalities. It does this through world-class science, knowledge and intelligence, advocacy, partnerships and the delivery of specialist public health services. PHE is an operationally autonomous executive agency of the Department of Health.

Public Health England Wellington House 133-155 Waterloo Road London SE1 8UG Tel: 020 7654 8001 www.gov.uk/phe Twitter: @PHE_uk Facebook: www.facebook.com/PublicHealthEngland

Prepared by: ESPAUR SSTF Implementation subgroup (see Appendix 3 for membership) For queries relating to this document, please contact: ESPAUR@phe.gov.uk

© Crown copyright 2015

You may re-use this information (excluding logos) free of charge in any format or medium, under the terms of the Open Government Licence v2.0. To view this licence, visit OGL or email psi@nationalarchives.gsi.gov.uk. Where we have identified any third party copyright information you will need to obtain permission from the copyright holders concerned. Any enquiries regarding this publication should be sent to ESPAUR@phe.gov.uk

Published March 2015 PHE publications gateway number: 2014828



Contents

About Public Health England	2
Executive summary	4
Introduction	9
Aim Context Antimicrobial Stewardship Programme An assessment of the Trust's Antimicrobial Stewardship Activities An Antimicrobial Stewardship Management Team/Committee A ward-focused antimicrobial team Evidence-based antimicrobial prescribing guidelines Quality Assurance Measures/Audits and Feedback Education and Training Why use the toolkit? Start Smart	9 9 11 12 13 13 15 17 17 18
Then Focus	19
Components of Best Practice for Antimicrobial Prescribing (Treatment)	20
Components of Best Practice for Antimicrobial Prescribing (peri-operative prophylaxis)	23
Appendix 1	25
Appendix 2 - Other relevant toolkits and resources	25
Appendix 3: List of ESPAUR SSTF implementation subgroup members	26

Executive summary

This document is an update of the guidance published in 2011. It takes into account recommendations from the Annual Report of the Chief Medical Officer (CMO) 2011 (published in March 2013),¹ the UK Five Year Antimicrobial Resistance Strategy 2013 to 2018,² the Cochrane Review - interventions to improve antibiotic prescribing practices for hospital inpatients 2013³ and the English Surveillance Programme for Utilisation and Resistance (ESPAUR) report 2014.⁴ It also acknowledges the forthcoming Cochrane update and draft antimicrobial stewardship guideline produced by the National Institute for Health and Care Excellence (NICE).^{5,6}

The prevalence of antimicrobial resistance (AMR) has risen alarmingly over the last 40 years, and few truly novel antimicrobials have been developed. This has led to increased pressure on existing antibiotics and greater challenges in treating patients. Inappropriate use of antimicrobials increases the risk to patients of colonisation and infection with resistant organisms and subsequent transmission to other patients.

Antimicrobial stewardship is an important element of the both the UK Five Year Antimicrobial Resistance Strategy² and the 2011 CMO report.¹ The aims of such stewardship initiatives are to improve the safety and quality of patient care and to contribute significantly to reductions in the emergence and spread of AMR. These aims are ultimately achieved by improving antimicrobial prescribing through an organised antimicrobial management program. A Start Smart - then Focus approach is recommended for all antibiotic prescriptions.

Criterion 9 of the Health and Social Care Act 2008: Code of Practice on the prevention and control of infections and related guidance outlines the standards for compliance for registered organisations to provide evidence of prudent prescribing and antimicrobial stewardship. 'The updated Code of Practice places greater emphasis on antimicrobial resistance and stewardship. The Code states: "Procedures should be in place to ensure prudent prescribing and antimicrobial stewardship. There should be an ongoing programme of audit, revision and update. In healthcare this is usually monitored by the antimicrobial management team or local prescribing advisors".

The Health and Social Care Act 2008. Code of practice for the NHS on the

¹ Davies S. Annual Report of the Chief Medical Officer 2011: Volume Two. Infections and the Rise of Antimicrobial Resistance. http:// www.dh.gov.uk/health/2013/03/cmo-vol2/ (10 Aug 2014, date last accessed).

UK Five Year Antimicrobial Resistance Strategy 2013 to 2018;

https://www.gov.uk/government/uploads/system/uploads/attachment data/file/244058/20130902 UK 5 year AMR strategy.p df (10 Aug 2014, date last accessed).

³ Davey P, Brown E, Charani E, et al. Interventions to improve antibiotic prescribing practices for hospital inpatients. Cochrane Database Syst Rev 2013;(4):CD003543. http://dx.doi.org/10.1002/14651858.CD003543.pub3. PHE. English Surveillance Programme for Antimicrobial Utilisation and Resistance (ESPAUR) report 2014.

https://www.gov.uk/government/publications/english-surveillance-programme-antimicrobial-utilisation-and-resistance-espaur-

report ⁵ Davey P, Peden C, Brown E et al. interventions to improve antibiotic prescribing practices for hospital inpatients (updated protocol) http://onlinelibrary.wiley.com/doi/10.1002/14651858.CD011236/full (27 Aug 2014 date last accessed)

NICE guidance: Antimicrobial stewardship https://www.nice.org.uk/guidance/indevelopment/gid-antimicrobialstewardship (19 Feb 2015 date last accessed)

prevention and control of health care associated infections and related guidance. Department of Health 2010. http://www.dh.gov.uk/en/Publicationsandstatistics/Publications/PublicationsPolicyAndGuidance/DH 122604

Outcome 8 of the CQC's Essential Standards for Quality and Safety on cleanliness and infection control also points to the Code of Practice⁸.

Improving antimicrobial prescribing and stewardship is dependent on strong clinical leadership. Within local Trusts, medical teams, in particular consultants should take a leadership role for antimicrobial quality improvement in their specialist areas. This should be done in collaboration with a consultant microbiologist/infectious diseases specialist and the antimicrobial pharmacist. Such initiatives should also seek to engage with junior doctors in order to develop a wider understanding of antimicrobial stewardship throughout the organisation.

This toolkit provides an outline of evidence-based antimicrobial stewardship in the secondary healthcare setting. It is recommended that the AMS management team/committee or equivalent use this toolkit along with the Code of Practice and *'Clostridium difficile:* how to deal with the problem'. This needs to be accompanied with a robust programme of auditing activities that promote safe and appropriate use of antimicrobials.^{7,9}

These activities will form part of the quality improvement strategy for patient safety and help to reduce inappropriate prescribing and optimise antibiotic use. Implementation of this toolkit and the audit programme can be used as evidence of meeting criterion 9 of the Code of Practice on the prevention and control of infections when seeking registration with the Care Quality Commission.

Figures 1 and 2 show the Start Smart, Then Focus treatment algorithm and the antimicrobial stewardship surgical prophylaxis algorithm.

Start Smart - this means:

- do not start antimicrobial therapy unless there is clear evidence of infection
- take a thorough drug allergy history
- initiate prompt effective antibiotic treatment within one hour of diagnosis (or as soon as possible) in patients with severe sepsis or life-threatening infections. Avoid inappropriate use of broad-spectrum antibiotics
- comply with local antimicrobial prescribing guidance
- document clinical indication (and disease severity if appropriate), drug name, dose and route on drug chart and in clinical notes*
- include review/stop date or duration
- obtain cultures prior to commencing therapy where possible (but do not delay therapy)
- prescribe single dose antibiotics for surgical prophylaxis where antibiotics have been shown to be effective (figure 2)
- document the exact indication on the drug chart (rather than stating long term prophylaxis) for clinical prophylaxis

* Inclusion of these in both the drug chart and in clinical notes may clarify the patient treatment pathway thus aiding in the improvement of patient outcomes and in medico-legal outcomes such as for *C. difficile* apportionment.

⁸ http://www.cqc.org.uk/sites/default/files/documents/gac_-_dec_2011_update.pdf

⁹ Department of Health and the Health Protection Agency. Clostridium difficile infection: How to deal with the problem. London: Department of Health. 2008.

http://www.dh.gov.uk/prod_consum_dh/groups/dh_digitalassets/documents/digitalasset/dh_093218.pdf

Then Focus – this means:

- reviewing the clinical diagnosis and the continuing need for antibiotics at 48*-72 hours and documenting a clear plan of action the 'antimicrobial prescribing decision'
- the five 'antimicrobial prescribing decision' options are:
 - 1. **Stop** antibiotics if there is no evidence of infection
 - 2. Switch antibiotics from intravenous to oral
 - 3. Change antibiotics ideally to a narrower spectrum or broader if required
 - 4. **Continue** and document next review date or stop date
 - 5. Outpatient Parenteral Antibiotic Therapy (**OPAT**)¹⁰
 - it is essential that the review and subsequent decision is clearly documented in the clinical notes and on the drug chart where possible eg stop antibiotic

* Due to advances in rapid diagnostics it may be possible to review prior to 48 hours after first dose.

It is recommended that as a minimum, providers should develop an action plan and monitor adherence to Start Smart Then Focus principles regularly in all clinical areas (at least annually). In particular monitoring:

- evidence of documenting indication and duration (or review date) on the drug chart
- evidence of antimicrobial stewardship review of antibiotics at 48-72 hours after initiation and documentation of the antimicrobial prescribing decision (one of five options) on the drug chart (or in the clinical notes – see Figure 1)
- the time between the onset of sepsis related hypotension and administration of appropriate antibiotics (this may be part of 'Surviving Sepsis' related audits within the Trust)
- adherence with local guidance on the choice of antibiotic therapy (or documented reason for non-compliance)
- antimicrobial resistance and consumption trends

¹⁰ British Society for Antimicrobial Chemotherapy. OPAT Good Practice Recommendations Available at: http://eopat.com/opat-standards

Figure 1: Antimicrobial Stewardship (AMS) – Treatment algorithm



Advocating patient safety and auditing of antimicrobial stewardship in hospitals should be based around the principles stated in this AMS algorithm. Examples of audit tools are shared in Appendix 1

Figure 2: Antimicrobial Stewardship (AMS) – Surgical prophylaxis algorithm



Advocating patient safety and auditing of antimicrobial stewardship in hospitals should be based around the principles stated in this AMS algorithm. Examples of audit tools are shared in Appendix 1. Deviations from the NICE guideline should be evidence based, with prolonged prophylaxis needing evidence of benefit.

Introduction

Aim

The aim of this toolkit is to provide an evidence-based outline for antimicrobial stewardship (AMS) in the secondary healthcare setting.

Context

The Code of Practice on the prevention and control of infections and related guidance applies to all providers of healthcare and adult social care under the Health and Social Care Act 2008.⁷ The Code sets out the 10 criteria against which a registered provider will be judged on how it complies with the registration requirement for cleanliness and infection control. Antimicrobial resistance and stewardship has been strengthened within the updated document and it also provides a range of information to support providers in complying with the regulations.

The law states that the Code must be taken into account by the Care Quality Commission (CQC) when it makes decisions about registration against the cleanliness and infection control requirements. The regulations also state that providers must have regard to the Code when deciding how they will comply with the registration requirements. So by following the Code, registered providers will be able to show that they meet the requirement set out in the regulations.

This toolkit will help healthcare providers assess whether they meet Criterion 9 of the Health and Social Care Act 2008: Code of Practice on the prevention and control of infections and related guidance.⁷

Guidance on meeting this criterion states: 'Procedures should be in place to ensure prudent prescribing and antimicrobial stewardship. There should be an ongoing programme of audit, revision and update. In healthcare this is usually monitored by the antimicrobial management team or local prescribing advisors'.⁷

Current evidence clearly demonstrates that the inappropriate use of broad-spectrum antimicrobials is associated with the selection of AMR bacteria. This includes extended-spectrum beta-lactamase (ESBL)-producing Gram-negative bacteria,^{11,12} Methicillin resistant *Staphylococcus aureus* (MRSA)^{13,14,15,16} and the induction of *Clostridium difficile* infection

¹² Hawkey, P., Jones, A. The Changing Epidemiology of Resistance. J Antimicrob Chemother. 2009; 64:Suppl 1:i3–10 ¹³ Lucet JC, Chevret S, Durand-Zaleski I, Chastang C, Regnier B. Prevalence and risk factors for carriage of methicillinresistant *Staphylococcus aureus* at admission to the intensive care unit: results of a multicenter study. Arch Intern Med 2003:163:181-8

¹¹ Livermore DM. Has the era of untreatable infections arrived? J Antimicrob Chemother. 2009;64(Suppl 1):29-36

¹⁴ Tacconelli E, De Angelis, Cataldo MA, et al. Does antibiotic exposure increase the risk of methicillin-resistant *Staphylococcus aureus* (MRSA) isolation? A systematic review and meta-analysis. J Antimicrob Chemother. 2008; 61(1): 26-38

¹⁵ Dancer SJ. The effect of antibiotics on meticillin-resistant *Staphylococcus aureus*. J Antimicrob Chemother. 2008; 61:246-253

¹⁶ Liebowitz LD and Blunt MC. Modification in prescribing practices for third generation cephalosporins and ciprofloxacin is associated with a reduction in meticillin- resistant *Staphylococcus aureus* bacteraemia rate. J.Hosp Inf 2008; 69:328-336

(CDI)^{3,7,17,18,19} and can cause long-lasting harmful changes to the body's protective microbial flora.^{20,21}

Broad-spectrum antibiotics (including cephalosporins, fluoroquinolones, co-amoxiclav, piperacillin-tazobactam, carbapenems and clindamycin) have been most associated with CDI, but all antibiotics should be avoided unless there are clear clinical indications for their use. Antibiotics should be used for the shortest duration possible that gives an appropriate clinical outcome. They should also be managed within a multifactorial programme (including infection prevention and control precautions) aimed at reducing healthcare-associated infections (HCAI) and improving antimicrobial use.6,7

A stong and robust antimicrobial stewardship programme is seen as a key component in the reduction of some HCAIs in support of patient safety. The draft NICE guideline on antimicrobial stewardship recommends that organisations should establish an antimicrobial stewardship programme taking account of the resources needed to support good antimicrobial stewardship across all care settings.⁶

There are a number of reports which highlight the failure to embed stewardship programmes into local practice. These include a National Audit Office report, published in 2009, which suggested that one-third of Trusts in England did not have a robust strategy to review antimicrobial prescriptions automatically within a defined period.²² A separate national survey of antimicrobial stewardship activites by the English Surveillance Programme for Antimicrobial Utilization and Resistance (ESPAUR) revealed that although a large majority (87.9%) of Trusts reported reviewing the Start Smart Then Focus (SSTF) document formally or informally; only 48% of Trusts report implementing a SSTF action plan after a review. In addition, whilst 79% of Acute Trusts collate data on at least one of the recommended audits in SSTF, there is a low uptake of audits that can be correlated to patient outcomes (eg time to first dose in severe sepsis, post prescription review and documentation at 48 hours).⁴

Like any change and improvement activity, especially those linked to patient safety, the success of an antimicrobial stewardship program is dependent on the support of hospital management and senior clinical staff. It is no longer the responsibility of specialists alone to champion the stewardship efforts within an organisation. Ultimately the Trust Board, managers and staff are all responsible for establishing, maintaining and supporting a coordinated approach to antimicrobial stewardship.

¹⁷ Wistrom, J., S. R. Norrby, E. B. Myhre, S. Eriksson, G. Granstrom, L. Lagergren, G. Englund, C. E. Nord, and B. Svenungsson. 2001. Frequency of antibiotic-associated diarrhoea in 2462 antibiotic-treated hospitalized patients: a prospective study. J. Antimicrob. Chemother. 47:43-50

Freeman J, Bauer MP, Baines SD, Corver J, Fawley WN, Goorhuis B, Kuijper EJ, Wilcox MH The changing epidemiology of Clostridium difficile infections. Clin. Microbiol. Rev. 2010. 23:529-549.

Nelson RL, Kelsey P, Leeman H, Meardon N, Patel H, Paul K, Rees R, Taylor B, Wood E, Malakun R. Antibiotic treatment for Clostridium difficile-associated diarrhea in adults. Cochrane Database of Systematic Reviews 2011, Issue 9. Art. No.: CD004610. DOI: 10.1002/14651858.CD004610.pub4

Blaser M. Antibiotic overuse: Stop the killing of beneficial bacteria. Nature. 2011 476: 393-394

²¹ Hviid A, Svanström H, Frisch M Antibiotic use and inflammatory bowel diseases in childhood Gut. 2011 Jan;60(1):49-54

Antimicrobial Stewardship Programme

Current guidance and published evidence recommends an antimicrobial stewardship programme should include the following: 1,2,3,6,7,23,24,25,26,27,28,29,30,31

- an assessment of the Trust's antimicrobial stewardship activities
- an antimicrobial stewardship management team/committee
- a ward-focused antimicrobial team
- evidence-based antimicrobial prescribing guidelines
- quality assurance measures/audits and feedback
- education and training

An assessment of the Trust's Antimicrobial Stewardship Activities

Trusts should demonstrate that there has been an assessment of the organisation's antimicrobial stewardship activities against the Start Smart Then Focus AMS toolkit as well as developing an action plan in order to provide an assurance to the Trust Board of safe, effective and appropriate antimicrobial prescribing (Table 1).

Table 1: It is recommended that as a minimum, providers should develop an action plan and monitor adherence to Start Smart Then Focus principles regularly in all clinical areas (at least annually). In particular monitoring:

- evidence of documenting indication and duration (or review date) on the drug chart
- evidence of antimicrobial stewardship review of antibiotics at 48-72 hours after initiation and documentation of the antimicrobial prescribing decision (one of five options) on the drug chart (or in the clinical notes – see Figure 1)
- the time between the onset of sepsis related hypotension and administration of appropriate antibiotics – this may be part of 'Surviving Sepsis' related audits

²³ Dellit TH, Owens RC, McGowan JE et al. Infectious Diseases Society of America and the Society for Healthcare Epidemiology of America Guidelines for Developing an Institutional Program to Enhance Antimicrobial Stewardship. Clin Infect Dis. 2007;44:159-77.

²⁴ Cooke J. Alexander K. Charani E et al. Antimicrobial stewardship: an evidence-based, antimicrobial self-assessment toolkit (ASAT) for acute hospitals. J Antimicrob Chemother. 2010;65(12):2669-73

Available at: http://www.researchdirectorate.org.uk/uhsm/asat/asat.asp

Thern J, Strauss R, Steib-Bauert M et al. Selection of hospital antimicrobial prescribing quality indicators: a consensus among German antibiotic stewardship (ABS) networkers. Infection, 2014;42(2): 351-362

Center for Disease Control and Prevention (CDC); Checklist for Core Elements of Hospital Antibiotic Stewardship Programs. Available at: http://www.cdc.gov/getsmart/healthcare/pdfs/checklist.pdf , (10 Aug 2014, date last accessed).

Buyle FM, Metz-Gercek S, Mechtler R et al. Development and validation of potential structure indicators for evaluating antimicrobial stewardship programmes in European hospitals. European journal of clinical microbiology & infectious diseases, 2013; 32(9): 1161-1170.

²⁸ Dumartin C, Rogues AM, Amade o B et al. Antibiotic usage in south-western French hospitals: trends and association with antibiotic stewardship measures. J Antimicrob Chemother 2011; 66: 1631-7.

²⁹ Bruce J, MacKenzie FM, Cookson B et al. Antibiotic stewardship and consumption: findings from a pan-European hospital study. J Antimicrob Chemother 2009; 64: 853-60

Australian Commission on Safety and Quality in Health Care. 2011. Antimicrobial stewardship in Australian hospitals. Available at: http://www.safetyandguality.gov.au/publications/antimicrobial-stewardship/ (10 Aug 2014, date last accessed).

Society for Healthcare Epidemiology of America, and Infectious Diseases Society of America. "Policy Statement on Antimicrobial Stewardship by the Society for Healthcare Epidemiology of America (SHEA), the Infectious Diseases Society of America (IDSA), and the Pediatric Infectious Diseases Society (PIDS)." Policy Statement 33.4 (2012): 322-327.

within the Trust

- adherence with local guidance on the choice of antibiotic therapy (or documented reason for non-compliance)
- antimicrobial resistance and consumption trends

The use of a self-assessment tool such as the antimicrobial self-assessment toolkit (ASAT) may be a useful additional resource to enable such an appraisal.²⁴ There are also a number of published quality indicators and check-lists that may be useful.²⁵⁻²⁹ While the individual elements in these tools and checklists may be helpful in improving antibiotic use, not all elements may be feasible in all hospitals.

An Antimicrobial Stewardship Management Team/Committee

It is recommended that a multidisciplinary management team/committee be set up to develop and implement the organisation's antimicrobial stewardship programme for all adults and children admitted to hospital.^{6,7}

This multidisciplinary group may have different names (for example the antimicrobial/antibiotic stewardship committee/group or management team). The name of the management committee is less important than the core functions of the group, which are described below. The suggested minimum core membership should include: a consultant microbiologist/infectious diseases specialist, an antimicrobial pharmacist, an acute care physician, a surgeon, a senior member of the pharmacy management team, an anaesthetist, a paediatrician, a senior nurse and primary care representation (to ensure a whole healthcare economy approach). The aim is to ensure a multidisciplinary approach and improve engagement across the organisation.³²

The management team/committee should report antimicrobial stewardship activities to the Trust Board via the organisation's Director of Infection Prevention and Control (DIPC)/Infection Control Committee and/or the Drugs and Therapeutic Committee (or equivalent).

The key roles of the AMS management team/committee are to:

- ensure that evidence-based local antimicrobial guidelines are in place and reviewed regularly or when new evidence is published
- ensure regular auditing of the guidelines, antimicrobial stewardship practice and quality assurance measures
- report a regular formal review of the organisation's retrospective antibiotic consumption data (especially highlighting the use of broad-spectrum antibiotics such as cephalosporins, co-amoxiclav, piperacillin-tazobactam, fluoroquinolones and carbapenems). PHE and ESPAUR measure antimicrobial consumption as Defined Daily Doses (DDD) per 100 admissions. The presentation of DDD per admissions rather than bed-days reflects hospital activity for admissions rather than those who are in hospital only. This

³² Cortoos PJ, De Witte K, Peetermans WE et al. Opposing expectations and suboptimal use of a local antibiotic hospital guideline: a qualitative study. J Antimicrob Chemother 2008; 62(1): 189-195.

measurement would allow comparison with national data and consistent benchmarking between Trusts

• identify actions to address non-compliance with local guidelines, general antimicrobial stewardship issues and other prescribing issues

A ward-focused antimicrobial team

A ward-focused antimicrobial team is recommended in addition to the AMS management team/committee. This should include the antimicrobial pharmacist and consultant microbiologist/infectious diseases specialist that report to the AMS management team/committee. The ward-focused team would be expected to review prescriptions at ward level as part of multi-disciplinary antimicrobial stewardship ward rounds.

Evidence-based antimicrobial prescribing guidelines

It is recommended that each organisation draw up a local antimicrobial stewardship policy and develop local antimicrobial guidelines based on national guidance (for example from the British National Formulary, NICE or Public Health England).

These local guidelines should be evidence-based, relevant to the local healthcare setting and take into account local antibiotic resistance patterns. They should cover diagnosis and treatment of common infections and prophylaxis of infection. Prescribers should adhere to these guidelines and compliance should be monitored and supported by senior clinicians and pharmacists. Responsibility for guideline implementation should reside with the AMS management team/committee; Drugs and Therapeutics Committee or equivalent and the Director for Infection Prevention and Control.

The local antimicrobial stewardship policy should contain:

- a policy statement that outlines the need for clear clinical case definitions and associated evidence of infection to minimise unnecessary prescribing of antimicrobials ^{33,34}
- 2. an emphasis of the urgent need to start treatment with effective antibiotic agents for severe sepsis or life-threatening infections ^{35,36,37,38}
- 3. a reminder for prescribers to use antibiotic agent(s) with an adequate spectrum to cover only the expected pathogens for less severe infections. To highlight that

³³ Costello C, Metcalfe C, Lovering A, et al. Effect of antibiotic prescribing in primary care on antimicrobial resistance in individual patients: systemic review and meta-analysis. BMJ 2010; 340:c2096.

³⁴ Saeed K, Dryden M, Bourne S, Paget C, Proud A; 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 Aug;78(4):289-92.

 ³⁵ Kollef MH, Sherman G, Ward S, et al: Inadequate antimicrobial treatment of infections. Chest 1999; 115:462-474
 ³⁶ Kumar A, Roberts D, Wood KE, Light B, Parrillo JE, Sharma S, Suppes R, Feinstein D, Zanotti S, Taiberg L, Gurka D, Kumar

A, Cheang M. Duration of hypotension before initiation of effective antimicrobial therapy is the critical determinant of survival in human septic shock. Crit Care Med. 2006 Jun;34(6):1589-96.

³⁷ Tang CM, Macfarlane JT. Early management of younger adults dying of community acquired pneumonia. Resp Med 1993;87: 289-94

³⁸ Dellinger RP, Levy MM, Rhodes A et al. surviving sepsis campaign: international guidelines for management of severe sepsis and septic shock: 2012. Crit Care Med 2013; 41: 580–637. Available at:

http://www.survivingsepsis.org/SiteCollectionDocuments/Final%2008%20SSC%20Guidelines.pdf

broad-spectrum antibiotics are sometimes not as potent in vitro as their narrowerspectrum counterparts against certain pathogens ^{39,40}

- 4. a reminder for prescribers to consider the risk of resistant pathogens such as MRSA or ESBL-producing organisms and offer alternative treatment regimens accordingly or encourage prescribers to seek expert advice ^{41,42,43}
- 5. a description of the importance of confirming the allergy status of recommended antibiotic agents in patients as there may be a need to offer alternative treatment choices for those who are allergic.⁴⁴ In line with NICE guidance on drug allergy,⁴⁵ patients with a history of such allergies should be assessed, and the allergy label removed where it is not correct, in order to improve patient outcomes
- an outline for prescribers to take appropriate specimens for culture and sensitivity testing prior to commencing antibiotic treatment. However they should not delay starting treatment in patients with severe sepsis or life-threatening infections ³⁸
- 7. a recommendation for intravenous (IV) administration only to patients who are severely ill, unable to tolerate oral treatment, or where oral therapy would not provide adequate coverage or tissue penetration
- 8. an outline for prescribers to review microbiology results daily and to de-escalate to pathogen-directed narrow-spectrum treatment promptly where appropriate ^{46,47}
- 9. a recommendation for prescribers to document the next review date or stop date and switch to the oral route of administration promptly in accordance with local IV-to-oral switch guidance ⁴⁸

Antimicrobial prescribing guidelines should be guided by evidence and local susceptibility data (eq by area team where available).⁴ Guidelines should include the following:

- 1. clinical diagnosis to include: case definition, evidence of infection, severity assessment and relevant microbiology investigations
- recommendations for non-antimicrobial treatment (eg fluid resuscitation or surgery)

³⁹ EUCAST. MIC distributions. http://www.eucast.org/mic_distributions/.

⁴⁰ HPA Antimicrobial Resistance and Prescribing in England, Wales and Northern Ireland, 2008 http://www.hpa.org.uk/web/HPAwebFile/HPAweb_C/1216798080469

Safdar N, Maki DG. The commonality of risk factors for nosocomial colonization and infection with antimicrobial-resistant Staphylococcus aureus, enterococcus, gram-negative bacilli, Clostridium difficile, and Candida. Ann Intern Med. 2002 Jun 4;136(11):834-44.

Owens, R.C., Donskey, C.J., Gaynes, R.P. et al. (2008) Antimicrobial-associated risk factors for Clostridium difficile infection. Clin Infect Dis. 46(Suppl 1), S19-S31 ⁴³ Vernaz N, Sax H, Pittet D, Bonnabry P, Schrenzel J, Harbarth S. Temporal effects of antibiotic use and hand rub

consumption on the incidence of MRSA and Clostridium difficile. J Antimicrob Chemother. 2008 vol. 62(3) pp. 601-607

⁴ Charneski L, Deshpande G, Smith SW. Impact of an antimicrobial allergy label in the medical record on clinical outcomes in hospitalized patients. Pharmacother. 2011;31(8):742-7

NICE Clinical Guideline183. Drug allergy: diagnosis and management of drug allergy in adults, children and young people. http://www.nice.org.uk/guidance/cg183

⁴⁶ Alvarez-Lerma, F Modification of empiric antibiotic treatment in patients with pneumonia acquired in the intensive care unit: ICU-Acquired Pneumonia Study Group. Intensive Care Med 1996;22,387-394

Alvarez-Lerma F, Alvarez B, Luque P, Ruiz F, Dominguez-Roldan JM, Quintana E, Sanz-Rodriguez C; ADANN Study Group. Empiric broad-spectrum antibiotic therapy of nosocomial pneumonia in the intensive care unit: a prospective observational study. Crit Care 2006; 10:R78.

Mertz D, Koller M, Haller P, Lampert ML, Plagge H, Hug B, et al Outcomes of early switching from intravenous to oral antibiotics on medical wards. J Antimicrob Chemother 2009;64:188-199

- empirical antimicrobial treatment recommendations: Initial antimicrobial therapy prior to availability of microbiology results or if a microbiological diagnosis is not going to be possible*
- 4. directed antimicrobial treatment when microbiology results are known and advice to contact clinical microbiologists/infectious diseases specialists if required*
- 5. oral switch guidance to highlight which oral agents to switch to and when ⁴⁸
- 6. duration of therapy for IV and oral agents ⁴⁹
- 7. specific guidance for exceptions and special cases if appropriate
- provide advice regarding monitoring and follow-up and contingency advice for treatment failure 6,48,50
- 9. guidance for prophylaxis for surgery or procedures. These should also include: the aim of prophylaxis eg reduce surgical site infection, where prophylaxis is required and where it is not, distinction between risk groups eg patients colonised with multi-drug resistant organisms such as MRSA, ESBL and CRE, alternatives where penicillin or other allergy exists and recommendation of single dose surgical prophylaxis regimens as appropriate and redosing frequency when more than one dose is required 51,52,53

*Empirical and directed treatment recommendations should specify the choice of drug(s), route of administration and dose. In addition a reminder for prescribers to adjust dosing for specific patient factors eg renal or hepatic impairment.54

Quality Assurance Measures/Audits and Feedback

Procedures should be in place to ensure prudent antibiotic prescribing and antimicrobial stewardship. This will necessitate an ongoing programme of audit, revision and update and should be monitored by the AMS management team/committee.

It is recommended that a multi-disciplinary quality improvement/audit programme for antimicrobial stewardship should be developed and sustained in every Acute Trust. Regular (at least annual) feedback of adherence to audits recommended within the Start Smart Then Focus toolkit should be provided to the Trust Board (as part of the annual infection control committee report), prescribers, lead clinicians, microbiologists/infectious diseases specialists, nurses, pharmacists and the DIPC (see Table 1).

⁴⁹ Chastre J, Wolff M, Fagon JY, Chevret S, Thomas F, Wermert D, Clementi E, Gonzalez J, Jusserand D, Asfar P, Perrin D, Fieux F, Aubas S; PneumA Trial Group. Comparison of 8 vs 15 days of antibiotic therapy for ventilator-associated pneumonia in adults: a randomized trial. JAMA. 2003 Nov 19;290(19):2588-98.

Singh N, Rogers P, Atwood CW, Wagener MM, Yu VL.Short-course empiric antibiotic therapy for patients with pulmonary infiltrates in the intensive care unit. A proposed solution for indiscriminate antibiotic prescription. Am J Respir Crit Care Med. 2000 Aug;162(2 Pt 1):505-11. ⁵¹ World Alliance for Patient Safety. WHO surgical safety checklist. June 2008. Available at:

http://www.who.int/patientsafety/safesurgery/tools_resources/SSSL_Checklist_finalJun08.pdf

NICE Clinical Guideline 74. Surgical Site Infection - Prevention and treatment of surgical site infection Available at: http://www.nice.org.uk/nicemedia/pdf/CG74NICEGuideline.pdf

Sign 104: Antibiotic prophylaxis in surgery. www.sign.ac.uk/guidelines/fulltext/104/index.html. 2008

⁵⁴ Thomas JK, Forrest A, Bharnani SM, et al. Pharmacodynamic evaluation of factors associated with the development of bacterial resistance in acutely ill patients during therapy. Antimicrob Agents Chemother. 1998;42:521-7

The AMS management team/committee and the DIPC should review antibiotic consumption trends regularly (at least annually).

Action should be taken to investigate non-adherence to local protocols for antibiotic prescribing (based on best practice) or unexpected trends in prescribing. It is recommended that these should be documented and reported, for example in minutes of the Antimicrobial Stewardship Committee/Team meetings

Trust-wide six-monthly or annual point prevalence studies (PPS) may be used to collect data to monitor compliance with the organisation's stewardship programme, and provide assurance for the organisation around compliance with the code of practice on prevention and control or infections.

Organisations should consider the formal investigation, via an existing clinical governance framework, of cases of repeated non-compliance (without clinical justification) or inappropriate prescribing, particularly when these result in an adverse patient outcome (eg development of an HCAI, prolonged length of stay, etc.). The Medical Director or Director of Infection Prevention and Control should challenge individuals whose prescribing practice is found to be repeatedly inappropriate.

It is important for providers to monitor patient outcomes to ensure that qualitative or quantitative alterations (changing, reducing, restricting) to antimicrobial prescribing do not have unintended detrimental effects for example increased time to clinical cure, increased mortality or increased readmission rate.^{55,56}

Improving antimicrobial prescribing and stewardship is dependent on strong clinical leadership. Within local Trusts, medical teams, in particular consultants should take a leadership role for antimicrobial quality improvement in their specialist areas. This would be done in collaboration with a consultant microbiologist/infectious diseases specialist and the antimicrobial pharmacist.

These initiatives should also seek to engage with junior doctors in order to develop a wider understanding of antimicrobial stewardship throughout the organisation. This is in agreement with the draft NICE antimicrobial stewardship guideline which recommends that organisations should encourage senior health professionals to promote antimicrobial stewardship within their teams, recognising the influence that senior prescribers can have on prescribing practices of colleagues.

Trusts are encouraged to benchmark antimicrobial quality assurance measures to provide an additional context to their individual data.

⁵⁵ Macgowan A. P. Urch J. Reynolds R. Jacobson S. K. Darley E. S. R. K-1715 - Impact of Changes in Antibiotic Use on the incidence of Clostridium difficile-Associated Diarrhoea (CDAD), in Hospital Mortality and Length of Hospital Stay (LOS). Presentation Abstract ICAAC Conference 2011 Chicago. Available at:

http://www.abstractsonline.com/plan/ViewAbstract.aspx?mID=2789&sKey=335516c4-7a3b-4f00-a2d7-

c884c70d10bb&cKey=5a788c37-0f1d-4ac8-8868-b8815dee61f1&mKey=%7B0C918954-D607-46A7-8073-44F4B537A439%7D ⁵⁶ Wachter RM, Flanders SA, Fee C, Pronovost PJ Public reporting of antibiotic timing in patients with pneumonia: lessons from

³⁰ Wachter RM, Flanders SA, Fee C, Pronovost PJ Public reporting of antibiotic timing in patients with pneumonia: lessons from a flawed performance measure. Ann Intern Med. 2008 Jul 1;149(1):29-32

Education and Training

'There should be mandatory core training in prudent antibiotic use for doctors, pharmacists and nurses <u>in addition</u> to an introductory session on each induction programme. Post-registration, this training should be repeated by all such staff every three years and should specifically cover those antibiotics that are linked to CDI'.⁹

Independent prescribers should use the antimicrobial prescribing and stewardship competencies (developed by the Department of Health advisory committee on antimicrobial resistance and healthcare associated infections (ARHAI) and PHE)^{57,58} to help develop their practice in relation to prescribing antimicrobials.

Nurses have a significant role to play in limiting the threat posed by AMR.^{56,59,60} They should be educated on the importance of avoiding missed and/or omitted doses (to maintain therapeutic levels) and ensuring that all diagnostic tests are carried out promptly. A targeted education strategy may facilitate the role that nurses also play in questioning and highlighting the duration of therapies and prescription of medications where these do not meet with established organisational guidelines. ^{56,57,58}

Why use the toolkit?

The NHS Standard Contract is a key enabler for commissioners to secure improvements in the quality of services for patients and to hold providers of NHS funded care to account. Each provider is required to have an HCAI reduction plan for each contract year (and to comply with its obligations under that plan) that must reflect local and national priorities relating to HCAI including AMR, set out obligations for the management and reduction of HCAI and be agreed between the provider and the commissioner.

It is recommended that organisations should use this toolkit as part of their quality improvement strategy for patient safety, enhancing stewardship in antibiotic usage, and ensuring optimal patient care and safety by reducing inappropriate prescribing. Compliance with this toolkit and auditing can be used as evidence of compliance with criterion 9 of the Code of Practice on the prevention and control of infections and incorporated into the HCAI reduction plan.

⁵⁷ Department of Health and Public Health England. Antimicrobial prescribing and stewardship competencies. London: Department of Health. 2013.

https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/253094/ARHAlprescrcompetencies__2_.pdf ⁵⁸ The Royal College of Nursing: Antimicrobial resistance: RCN position on the nursing contribution http://www.rcn.org.uk/ data/assets/pdf file/0003/590484/004681.pdf

⁵⁹ Edwards R Loveday H, Drumright L N, and Holmes A. Should nurses be more involved in antimicrobial management? J Infect Prev 2011 12: 4-5

⁶⁰ Edwards R, Drumright L N, Kiernan M, Holmes A Covering more territory to fight resistance: considering nurses role in antimicrobial stewardship. J Infect Prev 2011 12: 6-10

Start Smart

- **Do not start antibiotics in the absence of clinical evidence of bacterial infection** If there is evidence/suspicion of bacterial infection, use local guidelines to initiate prompt effective antibiotic treatment within one hour of diagnosis (or as soon as possible) in patients with life-threatening infections such as severe sepsis.³⁸ Avoid inappropriate use of broad-spectrum antibiotics.¹¹⁻²¹
- For antibiotic(s) prescribed, document each of the following on the drug chart and in the clinical notes: clinical indication (including disease severity if appropriate), dose, route and duration or review date.

Antibiotics in hospitals are often continued unnecessarily because clinicians caring for the patient do not have information indicating why the antibiotics were initially commenced and how long they were planned to be continued. This problem is compounded where primary responsibility for patient care is frequently transferred from one clinician to another. Ensuring that all antibiotic prescriptions are always accompanied by an indication and a clear duration or review date will help clinicians change or stop therapy when appropriate. In children the dose of antimicrobials should be prescribed according to the individuals weight/age - refer to local formulary or BNFc

Obtain cultures first where possible

Knowing the antibiotic susceptibility of an infecting organism can help clinicians to prescribe the most appropriate antibiotic. This is useful for narrowing of broad-spectrum therapy, changing therapy to effectively treat resistant pathogens and stopping antibiotics when cultures suggest an infection is unlikely. Cultures are also important for epidemiological surveillance.

Do not delay treatment for patients with life-threatening infections eg severe sepsis.

 Prescribe single dose antibiotics for surgical prophylaxis; where antibiotics have been shown to be effective.

Critical to this advice is that the single dose is administered within the 60 minutes prior to surgical incision or tourniquet inflation to enable peak blood levels to be present at the start of the surgical procedure.⁵¹ Intraoperative redosing is needed to ensure adequate serum and tissue concentrations of the antimicrobial if the duration of the procedure exceeds two half-lives of the antimicrobial or there is excessive blood loss (eg >1500 mL in adults, >25 ml/kg in children).⁶¹ A treatment course of antibiotics may also need to be given (in addition to appropriate prophylaxis) in cases of dirty surgery or infected wounds. The appropriate use and choice of antibiotics should be discussed with infection specialists for each case (see Figure 2 - Surgical Prophylaxis Algorithm).^{51,52}

⁶¹ Bratzler DW, Dellinger EP, Olsen KM et al. (2013). Clinical practice guidelines for antimicrobial prophylaxis in surgery. Am J Health Syst Pharm 2013; 70(3): 195-283

Then Focus

- Review the clinical diagnosis and the continuing need for antibiotics by 48-72 hours and make a clear plan of action the 'antimicrobial prescribing decision'
 Antibiotics are generally started before a patient's full clinical picture is known. By 48-72 hours, when additional information is available, including microbiology, radiographic and clinical information, it is important for clinicians to re-evaluate why the therapy was initiated in the first place and to gather evidence on whether there should be changes to the therapy.
- The five 'antimicrobial prescribing decision' options are Stop, Switch, Change, Continue and OPAT:
 - 1. Stop antibiotics if there is no evidence of infection
 - 2. Switch antibiotics from IV to oral
 - 3. Change antibiotics ideally to a narrower spectrum or broader if required. Prescribers should seek expert advice when necessary
 - 4. Continue and document next review date or stop date for IV and oral antibiotics
 - 5. Outpatient Parenteral Antibiotic Therapy (OPAT).¹⁰

For paediatric patients in particular, the choice of oral antibiotic should account for factors potentially affecting adherence such as dosing frequency and palatability/taste of formulation. Palatable oral drugs in a sensible regimen (up to 3 times per day) should be used where possible and middle of the night dosing of oral antibiotics should be avoided whenever possible, especially following discharge.

 It is essential that the review and subsequent decision be clearly documented in the clinical notes. The decision should also be documented clearly on the drug chart.

Components of Best Practice for Antimicrobial Prescribing (Treatment)

Organisations should develop their own audit strategy/programme and action plan.⁷ Each of the criteria below should be audited at least annually however there may be a need to monitor some aspects more frequently.

It is recommended that as a minimum, providers should develop an action plan and monitor adherence to Start Smart Then Focus principles regularly in all clinical areas (at least annually). In particular monitoring:

- evidence of documenting indication and duration (or review date) on drug chart
- evidence of antimicrobial stewardship review of antibiotics at 48-72 hours after initiation and documentation of the antimicrobial prescribing decision (one of five options) on the drug chart (or in the clinical notes) (Figure 1)
- the time between the onset of sepsis related hypotension and administration of appropriate antibiotics – this may be part of 'Surviving Sepsis' related audits within the Trust
- adherence with local guidance on the choice of antimicrobial therapy (or documented reason for non-compliance)
- antimicrobial resistance and consumption trends

Several components of best practice can be audited as part of the Trust-wide sixmonthly or annual point prevalence studies (PPS) or integrated into existing audit programmes established locally.*

* The tables below along with sample audit tools are available in Annex 1: Resource Materials - examples of audit tools, review stickers and drug charts

Criteria	Description of audit	Rationale for audit
1. Treatment of infection emergencies	Audit the treatment of severe sepsis and septic shock against clinical standards. This should include an audit of the time from the onset of severe sepsis to the administration of the first dose of antibiotic therapy ⁶²	A delay in starting adequate antibiotic therapy in severe infection is associated with increased morbidity and mortality

 Table 2: Components of Best Practice for Antimicrobial Prescribing (Treatment)

⁶² The UK Sepsis Trust. Clinical toolkits for professionals. http://sepsistrust.org/info-for-professionals/clinical-toolkits/

Criteria	Description of audit	Rationale for audit
2.Communication of the decision to prescribe antimicrobials	Audit the documentation of the decision to start antimicrobial therapy along with the indication or provisional diagnosis in the clinical notes and on the drug chart. This should include the clear identification of the prescriber and their contact details	Communication between healthcare teams is vital to ensure safe and effective patient care. This is mandated by the Royal Colleges. The requirement to document prescribing decisions will discourage antimicrobial prescribing where evidence of infection is lacking
3. Microbiology culture and sensitivities (MC&S)	Audit the appropriateness of specimens (for specific infections) obtained for MC&S. This should conform to local guidelines	The availability of appropriate cultures and sensitivities will facilitate the prompt de-escalation of broad- spectrum agents or the tailoring of therapy in cases of treatment failure
4. Antimicrobial consumption	Audit the consumption of antimicrobial agents (or de- escalation audit)	The unnecessary continuation of antimicrobials is associated with HCAIs and contributes to the development of AMR
5. Choice of antimicrobial agent(s)	Audit the choice of antimicrobial therapy. This should be according to local guidelines where available. This audit may also opt to include the dose and route of administration of the antimicrobials prescribed	Inappropriate antimicrobial therapy is associated with HCAIs, the development of AMR and the associated risks of unnecessary drug exposure
6. Review date for prescribed antimicrobials	Audit the review of antimicrobials at 48-72 hours after initiation. This should capture the documentation of the decision to continue current therapy and subsequent specified review or stop date	An expected duration or review date should be documented on antimicrobial prescriptions. This practice will discourage open-ended prescriptions
7. Duration of IV antimicrobial therapy	Where IV antimicrobials are continued at 48-72 hours after initiation, audit the documentation for continuing treatment. Audit the relative consumption of IV and oral antimicrobials	Treatment with IV antimicrobials should not continue beyond 48-72 hours unless recommended by local guideline or consultant microbiologist/infectious diseases specialist Unnecessary continuation of IV treatment increases the risk of line infection

Criteria	Description of audit	Rationale for audit
8. IV-to-oral antimicrobial switch	Audit compliance with local IV to oral switch OR Audit the relative consumption of IV and oral antimicrobials	Treatment with IV antimicrobials should be switched to oral therapy within 24 hours of meeting local switch criteria Unnecessary continuation of IV treatment increases the risk of line infection
9. Total duration of antimicrobial therapy	Audit antimicrobial consumption	Treatment with antimicrobials should not continue beyond 7 days (IV plus oral) unless recommended by a local guideline or consultant microbiologist/infectious diseases specialist Prolonged antibiotic therapy is associated with HCAIs, the development of AMR and other consequences of prolonged drug exposure

Components of Best Practice for Antimicrobial Prescribing (peri-operative prophylaxis)

Annual audit of local surgical prophylaxis practice to include:*

* The tables below along with sample audit tools are available in Annex 1: Resource Materials - examples of audit tools, review stickers and drug charts

Table 3: Components of Best Practice for Antimicrobial Prescribing (peri-operative prophylaxis)

Criteria	Description of audit	Rationale for audit
1. Need for antimicrobial prophylaxis	Audit the indication for antimicrobial prophylaxis. Practice should conform to local quidelines	For certain clean procedures, evidence suggests a lack of benefit of antimicrobials
		The clinical indication should comply with <i>NICE 74: Surgical site infection:</i> <i>Prevention and treatment of surgical</i> <i>site infection</i> ⁵²
2. Choice of antimicrobial agent(s) for perio- operative prophylaxis	Audit the choice of antimicrobial therapy. This should be according to local guidelines where available	Antimicrobial prophylaxis should ensure adequate coverage of expected pathogens according to surgical site. Whenever possible the prophylaxis should avoid cephalosporins, clindamycin and fluoroquinolones
		Where necessary, appropriate alternatives should be prescribed for patients with penicillin/ beta-lactam allergy, or those colonised with resistant organisms eg MRSA
		The choice of antimicrobial agent(s) should be prescribed according to local guidelines.
3. Timing of antimicrobial prophylaxis	Audit the time between the administration of antimicrobial prophylaxis and skin incision	Antimicrobial prophylaxis should be administered within 60-minutes prior to incision (or tourniquet) or according to local guidelines ⁵² . The lowest surgical site infection rates associated with optimal timing of pre-incision administration of antimicrobials

Criteria	Description of audit	Rationale for audit
4.Repeat doses of antimicrobial prophylaxis	Audit cases of multiple or post- operative antimicrobial prophylaxis	Single dose is indicated for majority of procedures and should be implemented unless there is clear evidence that multiple or post-operative dosing improves outcomes. Reason for antimicrobial administration beyond one dose should be documented and/or comply with criteria below agreed criteria
5. MRSA positive patients	Audit MRSA decolonization practice (normally collected by IPC teams)	Decolonisation therapy is recommended prior to surgery and antibiotic prophylaxis should include cover for MRSA

Appendix 1

See separate document: Resource Materials - Examples of audit tools, review stickers and drug charts

Appendix 2 - Other relevant toolkits and resources

PHE does not necessarily endorse the examples of antimicrobial stewardship in this section.

European Antibiotic Awareness Day: 2014 resources:

https://www.gov.uk/government/collections/european-antibiotic-awareness-day-resources

Prescribing competencies:

https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/253094/ARHAIpr escrcompetencies_2_pdf

Start Smart Then Focus Prescribers checklist:

https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/366944/Seconda ry_care_prescribers_checklist.pdf

Antimicrobial Self-Assessment Toolkit:

http://imperial-asat.herokuapp.com/

Sepsis clinical toolkit and audit tools:

http://www.collemergencymed.ac.uk/Shop-Floor/Clinical%20Standards/Sepsis http://sepsistrust.org/info-for-professionals/clinical-toolkits/

TARGET Antimicrobial prescribing toolkit for Primary Care:

http://www.rcgp.org.uk/clinical-and-research/target-antibiotics-toolkit.aspx

Appendix 3: List of ESPAUR SSTF implementation subgroup members

Dr Diane Ashiru-Oredope	Chair of ESPAUR SSTF implementation subgroup; PHE Pharmacist lead for AMRS and HCAI
Dr Gavin Barlow Mr Brian Brown	Consultant in Infection, Hull & East Yorkshire Hospitals NHS Trust National Pharmacy Manager, Care Quality Commission.
Dr Emma Budd	ESPAUR secretariat. PHE AMRS & HCAI programme
Dr Druin Burch	Consultant, Oxford University Hospitals. Representing the Royal College
	of Physicians
Dr Teh-Li Chin	Microbiology Consultant, North Bristol NHS Trust
Dr Oliver Dyar	Junior Doctor representative
Ms Rose Gallagher	Professional Lead - Infection Prevention & Control, Royal College of Nursing
Mr Mark Gilchrist	Consultant pharmacist for infection, Imperial College Healthcare NHS Trust, Representing UK Clinical Pharmacy Association
Mr Kieran Hand	Consultant Pharmacist – anti-infectives, University Hospital Southampton.
	Healthcare Associated infections (ARHAI)
Dr James Hatcher	Infectious Diseases/Medical Microbiology registrar. Imperial College
	London Healthcare NHS Trust
Mr Philip Howard	Consultant Pharmacist – Antimicrobials, Leeds Teaching Hospitals NHS
	Trust. Representing the Royal Pharmaceutical Society
Mr David Ladenheim	Lead SSTF editorial group. Antimicrobial Pharmacist, East & North Herts NHS Trust
Prof Heather Loveday	President, Infection Prevention Society
Prof Alasdair Macgowan	Professor of Antimicrobial Therapeutics, University of Bristol
Ms Kate Morrow	National Patient Safety Lead, NHS England
Dr Bharat Patel	PHE, AMRS & HCAI medical microbiologist
Dr Sanjay Patel	Consultant in Paediatric infectious diseases and immunology,
	Southampton Children's Hospital. Representing the Royal College of
Ms Laura Whitney	Consultant Dharmacist Antimicrobials St. George's Healthcare NHS
	Trust.
Prof Tony Young	Consultant Urological Surgeon, Southend University Hospital. Representing the Royal College of Surgeons



Protecting and improving the nation's health

Start Smart Then Focus Appendix 1 Resource Materials: Examples of audit tools, review stickers and drug charts

Examples provided by English Hospitals

About Public Health England

Public Health England exists to protect and improve the nation's health and wellbeing, and reduce health inequalities. It does this through world-class science, knowledge and intelligence, advocacy, partnerships and the delivery of specialist public health services. PHE is an operationally autonomous executive agency of the Department of Health.

Public Health England Wellington House 133-155 Waterloo Road London SE1 8UG Tel: 020 7654 8001 www.gov.uk/phe Twitter: @PHE_uk Facebook: www.facebook.com/PublicHealthEngland

For queries relating to this document, please contact: ESPAUR@PHE.gov.uk

© Crown copyright 2015

You may re-use this information (excluding logos) free of charge in any format or medium, under the terms of the Open Government Licence v2.0. To view this licence, visit OGL or email psi@nationalarchives.gsi.gov.uk. Where we have identified any third party copyright information you will need to obtain permission from the copyright holders concerned.

Published March 2015 PHE publications gateway number: 2014828



Contents

About Public Health England	2
Executive summary	4
Start Smart Then Focus tools	5
Antimicrobial Stewardship – Antibiotic Treatment Algorithm Antimicrobial Stewardship – Surgical Prophylaxis Algorithm Start Smart Then Focus Prescriber's checklist Examples of Antimicrobial Stewardship resources provided by English hospitals	5 6 7 8
Best practice audit tools Antimicrobial Review Stickers Dedicated Antimicrobial Section on drug chart; separate sheets Dedicated Antimicrobial Section on drug chart – added within regular prescriptions section Sepsis Audit Tool	8 18 20 22 27

Executive summary

This appendix section of the Start Smart Then Focus Antimicrobial Stewardship Toolkit for secondary care contains Antimicrobial Stewardship resource materials

- 1. Start Smart Then Focus tools:
 - Algorithms
 - Prescriber's checklist which can be printed and provided as aide-memoire for prescribers
- 2. Examples of Antimicrobial Stewardship resources provided by English hospitals:
 - audit tools,
 - review stickers,
 - Drug charts with specific antibiotic prescribing sections

PHE present the resources kindly provided by English hospitals as examples only.

Start Smart Then Focus tools

Antimicrobial Stewardship – Antibiotic Treatment Algorithm



Advocating patient safety and auditing of antimicrobial stewardship in hospitals should be based around the principles stated in this algorithm. Examples of audit tools are shared in the following pages

Antimicrobial Stewardship – Surgical Prophylaxis Algorithm



Redose for long surgical procedures

Intraoperative redosing is needed to ensure adequate serum and tissue concentrations of the antimicrobial if the duration of the procedure exceeds two half-lives of the antimicrobial or there is excessive blood loss (i.e., >1500 mL in adults³ or >25ml/kg in children). A treatment course of antibiotics may also need to be given (in addition to appropriate prophylaxis) in cases of dirty surgery or infected wounds³. The appropriate use and choice of antibiotics should be discussed with infection specialists for each case.

DOCUMENT ALL DECISIONS

References:

- 1. NICE clinical guideline 74: Surgical site infection Prevention and treatment of surgical site infection http://www.nice.org.uk/Guidance/CG74
- World alliance for Patient Safety. WHO surgical safety checklist June 2008 <u>http://www.who.int/patientsafety/safesurgery/tools_resources/SSSL_Checklist_finalJun08.pdf?ua=1</u>
- Bratzler DW, Dellinger EP, Olsen KM et al. (2013). Clinical practice guidelines for antimicrobial prophylaxis in surgery. Am J Health Syst Pharm 2013; 70(3): 195-283

Advocating patient safety and auditing of antimicrobial stewardship in hospitals should be based around the principles stated in this algorithm. Examples of audit tools are shared in the following pages

Start Smart Then Focus Prescriber's checklist

This can be printed and provided as an aide-memoire for prescribers

EUROPEAN MARKENESS DA Secondary Care Prescriber's Checklist Antibiotics - Overuse and incorrect use drives resistance
START SMART: do not start antimicrobial therapy unless there is clear evidence of infection take a thorough drug allergy history initiate prompt effective antibiotic treatment within one hour of diagnosis (or as soon as possible) in patients with severe sepsis or life-threatening infections. Avoid inappropriate use of broad-spectrum antibiotics
 comply with local antimicrobial prescribing guidance document clinical indication (and disease severity if appropriate), drug name, dose and route on drug chart and in clinical notes include review/stop date or duration obtain cultures prior to commencing therapy where possible (but do not delay therapy) prescribe single dose antibiotics for surgical prophylaxis where antibiotics have been shown to be effective document the exact indication on the drug chart (rather than stating long term prophylaxis) for clinical prophylaxis
THEN FOCUS: At 48 – 72 hours; review the patient and make a clinical decision "the Antimicrobial Prescribing Decision" on the need for on-going antibiotic therapy. Does patient's condition and/or culture result(s) necessitate: Stop of antibiotic therapy (if no evidence of infection) Switch from intravenous to oral therapy Change: de-escalation/substitution/addition of agents Continuation of current therapy Outpatient Parenteral Antibiotic Therapy (OPAT)
Reference: Antimicrobial Stewardship Toolkit for Secondary Care: Start Smart – then Focus Available at: https://www.gov.uk/government/publications/antimicrobial-stewardship-start-smart-then-focus SSTF was developed by Public Health England and the Department of Health expert advisory committee on Antimicrobial Resistance and Healthcare Associated Infection (ARHAI)

Available at:

https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/366944/Secondary_care_prescriber s_checklist.pdf

Examples of Antimicrobial Stewardship resources provided by English hospitals

Best practice audit tools

Form on APX_HAPP Patient III Formane Summe Summe Summe Summe Summe Summe Summe Summe Summe Summe Summe Summe Summe Standard 2: Antibiotic choice according to guideline *Care Group Cereral Surgery Secary OF *Standard 2: Antibiotic choice according to guideline *Standard 2: Appropriate dose prescribed *Standard 2: Total course length today is =-7 dos, or justified if longer *Standard 2: Total course length today is =-7 dos, or justified if longer *Standard 2: Total course length today is =-7 dos, or justified if longer *Standard 2: Total course length today is =-7 dos, or justified if longer *Standard 2: Total course length today is =-7 dos, or justified if longer *Standard 2: Total course length today is =-7 dos, or justified if longer *Standard 2: Total course length today is =-7 dos, or justified if longer *Standard 2: Total course length today is =-7 dos, or justified if longer *Standard 2: Total course length today is =-7 dos dos dos dos prescribed *Standard 2: Total course length today is =-7 do	Audit Entry		University Hospital Southampton
Control Net Action 11 Control Control Summary Control Contro Control Contr	Form on ABY, MARDI	Cascal	
Date audit recorded 13-NOV-2014	Potini on APA_nAPPI Patient IIo Forename Sumane Cons Ward *Care Group Specially 0 *Standard 1: Indication documented on start date *Standard 1: Note that was the outcome of the prescription review *Standard 5: Total course length today is <=7 days, or justified if longer Auditor i	General Surgery ORT Yes Off guideline-Not justified Ves Ves Change drug to broader spectrum Yes Kieran Hand	Standard 2: Antibiotic choice according to guideline X Was the empirical choice of antibiotic regimen according to UHS guideline or was a valid justification for off-guideline choice documented in the medical notes? Justification may include: culture and sensitivity report; risk of resistant pathogen (healthcare contact, co-morbidities, prior antibiotic exposure, travel history); recommendation by named medical microbiologist / ID doctor; treatment failure of guideline agents; contra-indication to guideline agent(s).
	Date audit recorded	13-NOV-2014	2014.11.05

Used with permission from Dr Kieran Hand - Southampton University Hospitals NHS Trust, Pharmacy and Microbiology Departments November 2014.

	Hospital Antimicrobial Prudent Prescribing Indicators (HAPPI) audit proforma											
Year	Month	Ward		Auditor	Name			Medical Notes – Prescribing Indicator Questions				
					1) Documentatio	2) Guideline prescribing or justified off-guideline Rx*		3) Duration to date				
	Drug Chart (complete one line below for <u>each</u> antimicrobial)					n of indication*	Choice of antimicrobial	Off- guideline prescribing	IV duration on audit day*	Total duration (IV + oral) on audit day for this indication*		
Date	Hospital number	Allergy box filled*	Antimicrobial name	Route	Course start date (IV/oral)	Review or stop date on chart*	Consultant team	Documented indication or provisional diagnosis? (on start date)	Guideline antimicrobial for indication	Valid reason documented on start date [§]	 IV duration currently ≤ 48h (surgical prophylaxis ≤ 24h) OR according to quideline 	 Total duration ≤ 7 days OR according to guideline
e.g.	1234567	Y	Flucloxacillin	Oral	23Jan09	Y	Dr Smith	Yes: Venflon infection	Y	N/A	Y	Y
e.g.	2345678	Y	Co-amoxiclav	IV	21Jan09	N	Dr Jones	Yes: Bronchiectasis	No guideline	N/A	N	Y
e.g.	3456789	N	Cefotaxime	IV	23Jan09	N	Dr Brown	No	Unknown	Reason (a)	Y	N/A (no guideline)

* Six prescribing standards. One point will be scored for each standard achieved (or N/A) for all antimicrobials prescribed for that patient.

[§]Valid reasons include: (a) contra-indication to guideline antimicrobials (e.g. allergy); (b) expert advice from named microbiology/infectious diseases doctor; (c) culture and sensitivity result (recent or previous) suggesting resistance to guideline antimicrobials; (d) patient risk factors for resistant pathogen (e.g. healthcare exposure, nursing/care home resident); (e) failure of reasonable trial of guideline therapy at adequate doses; (f) recent (within 2 weeks) exposure to guideline antimicrobials

Used with permission from Dr Kieran Hand - Southampton University Hospitals NHS Trust, Pharmacy and Microbiology Departments October 2009. Version 2.2 March 2010

Barking, Havering and Redbridge University Hospitals NHS Trust, Point Prevalence Study on Anti-infective Use Pharmacy Department

Q1 Date of Collection	Q2 Hospital Queens/ KGH	Q3 Ward	Q4a Allergies as written on chart	Q4b Is reaction of allergy stated? Y/N	Q5 Is Rx medical (m) or surgical (s)?	Q6 If surgical, is anti- microbial for <24hrs? Y/N/NA

	Antimicrobial Details														
							Route		Duration		Management code (API's)			Other	
	Q7 Antimicrobial	Q8 Dose & Frequency	Q9 Is indication on pt's drug chart/ medical notes at the point of prescribing? API 1 Y/N	Q10 Indication	Q12 Is antimicrobial prescribed acc. to Trust antimicrobial/ restricted guidelines? Y/N	Q11 Route (IV/PO/ Top)	Q13a If IV, is there a switch to PO within 72hrs? API 3 Y/N/NA	Q13b If IV- PO switch, Is total duration ≤7 days Y/N/NA	Q14 Is there a valid stop/review date or duration on the chart? API 2 Y/N/Na	Q15 If No for any API's is there a yellow sticker in pt's notes? Y/N/NA	Q16 If yellow sticker in notes has prescriber amended the prescription as req'd Y/N/NA	Q17 If no API sticker in notes is there appropriate p'cist endorsements on chart? Y/N/NA	Q18 If antimicrobial restricted, was Micro contacted where necessary? API 4 Y/N/Na	Q19 Is there DNO endorsement on chart? Y/N	
1															
2															
3															
4															

Antimicrobial Management Code – BHR Hospitals Used with permission from Antimicrobial Stewardship Group June 2011



÷	Site:		Ward:		Date:		Time taken:	
	Observation	Allergy Status Documented	Indication Documented	Stop/ Review Date Documented	Route Appropriate	Trust Guidelines Followed*	Comments	All Elements Achieved.
	1					Y / N / NA		
	2					Y / N / NA		
	3					Y / N / NA		
	4					Y / N / NA		
	5					Y / N / NA		
	6					Y / N / NA		
	7					Y / N / NA		
	8					Y / N / NA		
	9					Y / N / NA		
	10					Y / N / NA		
	TOTAL %							

Antimicrobial Care Bundle Audit Tool

*Trust guidelines followed if empirical treatment prescribed according to guidelines, antimicrobials prescribed according to culture and sensitivity results or following advice from Microbiology.

Data collection to ol developed by Allbhe Kavanach, Pre-registration Pharmacist and Claire Brandish, Lead Antimicrobial Pharmacist, October 2010

The Leeds Teaching Hospitals

NHS Trust

MONTHLY PRESCRIPTION AUDIT FORM

Date:		Compl	leted by:		Contact No
Main Specia	lity		Direct	orate:	Division:
Site (circle):	LGI	SJUH	CAH	WGH Name/Number of w	ard:

Antimicrobial standards: A duration or review date must be stated on the prescription chart An indication for the antimicrobial treatment must be stated on the prescription chart

Section 1: Prescribing of antimicrobials

	Space for notes (e.g. tally of patients)	Total number
How many antimicrobials do not have an indication on the chart?		
How many antimicrobials do not have a duration or review date stated on the chart?		
For how many antimicrobial prescriptions would you be unable to contact the prescriber if necessary?		

Section 2: Administration routes for antimicrobials

	Space for notes (e.g. tally of patients)	Total Number
Enteral (including oral)		
Parenteral (Infusion or injection)		
Number of parenterals given for greater than 48hrs		
In your opinion, how many of the parenteral antimicrobial prescriptions could have been given enterally? (Consider – is the patient's temperature normal, CRP normal, documented clinical improvement)		

Section 3: Sample size on the day of the audit

	Space for notes (e.g. tally of patients)	Total Number
Number of beds occupied		
Number of patients audited		
No of patients receiving antimicrobials		

NB. This audit is to be carried out on all the patients seen on the ward on one day each month, ideally within the first fortnight. This data is submitted to the Trust Board each month as a key performance indicator (KPI). Many thanks for your help.

C1Documents and Settings/amith/mi/Desktop/MonthlyAuditForm.doc

The Leeds Teaching Hospitals

MONTHLY PRESCRIPTION AUDIT FORM

				Antimicrobial M	edicines (Code Exce	ptions rep	orting for	m						
Pharm	acist Nam	ie:		Site: LGI/SJUH/CAH	WGH Wa	ard:	Date	e:		Monthly AB Med Code Audit: Y					
Pt initials	Unit number	Consultant	Specialty	Prescriber name, grade & contact number (if available from chart)	Indication missing (√)	Duration or review missing (*)	Prescriber name illegible (*)	Prescriber contact number missing (✓)	Prescriber spoken to about omission(s) (<)	Comments (eg on-call Dr, pt lodging on ward)					
					8										
		s.			<i>N</i>										
						0									
					0										
		S.			16										

Community Acquired Pneumonia Data Collection & Aggregation Form

Month_____Year____

Measures	Exar	nples					Pa	ntients					Total
To be done within 4hr of arrival at hospital	A	В	1	2	3	4	5	6	7	8	9	10	Data to be added to Extranet
Oxygen Therapy													
1) Oxygen saturation assessed?	Yes	Yes	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	1
2) Oxygen administered appropri	riately?												1
Choose target range													
a) Target range: 94-98% • O ₂ Sat < 94% • O ₂ administered	Yes	-	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	
 b) Target range: 88-92% O₂Sat < 88% O₂ administered 	-	Yes	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	
Severity Assessment	_	_	_	_		_		_		_		_	_
3) CURB65 score derived and documented?	Yes	Yes	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	Y/N	1
**Record CURB65 score	3	1											
Antibiotics if CURB65 ≥ 3													
IV antibiotics compliant with	policy												/
Route and Choice must be con	ect to be	e complia	nt										
macrolide)	Yes	NA	Y/N NA	Y/N NA	Y/N NA	Y/N NA	Y / N NA	Y/N NA	Y / N NA	Y/N NA	Y/N NA	Y/N NA	
Correct Route (Ⅳ)	Yes	NA	Y7N NA	Y7N NA	Y7N NA	Y7N NA	Y7N NA	Y7N NA	Y7N NA	Y7N NA	Y7N NA	Y/N	
Time to First Dose (CURB65 ≥ 3)												
to first antibiotic (hh : mm)	01:00	NA											

** Not for entry on Extranet Scottish Antimicrobial Prescribing Group

March 2011

ANTIMICROBIAL MONTHLY DATA COLLECTION

Antimicrobials - Collect data on ONE day for all patients on your ward. Document each antibiotic prescribed and whether a course length/review is recorded and whether added by medic or pharmacist. Do the same with the indication. Record if the Antimicrobial is policy or micro-approved. Mark each chart audited with an 'A' on the front top right hand corner to prevent double data collection. If the prescription is not compliant with policy, please add hospital number of patient.

[Name of Da	ta collector:				Ward: Date:							
[No. of patie	nts on ward:				No. of patients reviewed:							
+													
	Consultant	Antimicrobial		РО	Courselength/	by Pharm	Indication	by Pherm or	Policy or Micro	Patient's hospital no.			
			(~)	(\mathcal{O})	on Drug Chart	or medic (P or M)		Medic (Por M)	approve d ✓ x	with policy)			
Ī													
ſ													

CDI HEAT Target Empirical Prescribing Indicator Data Collection Form

<u>Example</u>

		Examp	ole Par	tients		Extended Date	Ré Compliance
Measures	A	В	С	D	E	Extranet Data	% compliance
Indication documented in patient's notes	Yes	Yes	No	No	Yes	3/5	60%
Antibiotics Compliant with Policy	No	Yes		-	Yes	2/3	66%

Month ______ Year _____ Ward _______

Week 1 Data	Patients						
Measures	1	2	3	4	5		
Indication documented in patient's notes	Y/N	Y/N	Y/N	Y/N	Y/N		
Antibiotics Compliant with Policy	Y/N	Y/N	Y/N	Y/N	Y/N		

Week 2 Data	Patients							
Measures	1	2	3	4	5			
Indication documented in patient's notes	Y/N	Y/N	Y/N	Y/N	Y/N			
Antibiotics Compliant with Policy	Y/N	Y/N	Y/N	Y/N	Y/N			

Week 3 Data	Patients				
Measures	1	2	3	4	5
Indication documented in patient's notes	Y/N	Y/N	Y/N	Y/N	Y/N
Antibiotics Compliant with Policy	Y/N	Y/N	Y/N	Y/N	Y/N

Week 4 Data	Patients				
Measures	1	2	3	4	5
Indication documented in patient's notes	Y/N	Y/N	Y/N	Y/N	Y/N
Antibiotics Compliant with Policy	Y/N	Y/N	Y/N	Y/N	Y/N

Total Monthly Extranet Data	Number Compliant	Total Patients
Indication documented in		
patient's notes		
Antibiotics compliant with		
policy		

Scottish Antimicrobial Prescribing Group

CDI HEAT Target Surgical Prophylaxis Data Collection Form

Example

	Example Patients						Rf Compliance	
Measures	A	В	С	D	E	Extranet Data	76 compliance	
Single dose	Yes	Yes	No	No	Yes	3/5	60%	
Antibiotics Compliant with Policy	No	Yes	No	Yes	Yes	3/5	60%	

Month _____ Year ____ Ward _____

Week 1 Data	Patients				
Measures	1	2	3	4	5
Single dose	Y/N	Y/N	Y/N	Y/N	Y/N
Antibiotics Compliant with Policy	Y/N	Y7N	Y/N	Y/N	Y/N

Week 2 Data	Patients				
Measures	1	2	3	4	5
Single dose	Y/N	Y/N	Y/N	Y/N	Y/N
Antibiotics Compliant with Policy	Y/N	Y/N	Y/N	Y/N	Y/N

Week 3 Data	Patients				
Measures	1	2	3	4	5
Single dose	Y/N	Y/N	Y/N	Y/N	Y/N
Antibiotics Compliant with Policy	Y/N	Y/N	Y/N	Y/N	Y/N

Week 4 Data	Patients				
Measures	1	2	3	4	5
Single dose	Y/N	Y/N	Y/N	Y/N	Y/N
Antibiotics Compliant with Policy	Y/N	Y/N	Y/N	Y/N	Y/N

Total Monthly Extranet Data	Number Compliant	Total Patients
Single dose		
Antibiotics compliant with policy		

Scottish Antimicrobial Prescribing Group

Antimicrobial Review Stickers

(NHS Tayside & Scottish Antimicrobial Prescribing Group)

Start date of antibiotic therapy	r: / / 2008	Date of review: / / 2008
Antibiotic plan	Agent:	Route of administration:
Dosage:	Dosing interval:	Planned total duration: days
If the patient receives iv antibi switch possible? □ Yes □ No □ Not app	iotic, is an oral plicable (not on iv)	Give the reason for that choice:

Pulcini et al <mark>JAC</mark>, 2008

Antimicrobial Management Code – Review Notice

The following antimicrobial prescription Trust's prescribing standards/Antimic	does	not fulfil the	
Antimicrobial details and date started			
 Please take the following action (s): API 1: specify the indication for a appropriate) API 2: Specify review date/stop of API 3: IV to oral review required API 4: Contact Medical Microbiol treatment (delete as appropriate) Please rewrite the prescription with a beyond 7 days 	antibiotics on medical notes date on drug chart within 72hours logy for restricted antibiotic specified duration on the	s/drug chart (delet c appro∨al/extende drug chart if you w	te as ed duration of /ish to continue
Pharmacist Name	Signature:	Date:	Time:
Approved by BHR Antimicrobia	al Stewardship Group and Drugs ar	d Therapeutics Commit	tee

Antimicrobial Management Code – BHR Hospitals Used with permission from Dr Diane Ashiru-Oredope - Barking Havering and Redbridge University Hospitals NHS Trust Antimicrobial Stewardship Group June 2011

Dedicated Antimicrobial Section on drug chart; separate sheets

ALLERGIES		NAME		BARCOD	E
		WARD	NHS/HOSPITAL	NUMBER	
SUGGESTED TIMES FOR ANTIMICROBIALS			TDS (8 hourly) –	6, 14, 22 : QDS (6	hourly) – 6, 12, 18, 24
	Antimi Tł Please en FC	icrobials should be provide the MEHT Antimicrobia sure the continued or ANTIMICRO	escribed in accordance Policy may be found of need for antimicro BIAL PRESCRI	with Hospital Formul on the Trust Intranet obials is reviewed PTIONS ONLY	ary DAILY
Drug			Date		ED
Clinical					OUIN
Route	Dose	Course length			S AN INUM
Signature		Bleep number			5 DAY
Date	Pharmacy screen	Pharmacy supply			WAT WICHCH
Additional Information					REVIE
Drug			Date 🗻 Times ¥		N OF
Clinical Indication					
Route	Dose	Course length			INUL IS RE
Signature		Bleep number			5 DAY
Date	Pharmacy screen	Pharmacy supply			W AT WICKO
Additional Information				REVIE	

Mid Essex Hospital Services NHS Trust Used with permission from Dr Louise Teare and Diane Ashiru-Oredope.

Impact of antimicrobial section on influencing prescribing published: http://www.pjonline.com/news/antimicrobials_section_in_hospital_prescribing_chart_improves_practices

INTRAVENUOS ANTEICOTOS ONLY Tick or insent times required 4		and the second second			that only enter	microsiology p	age of the Intran				1.00	Anti	biolic Guidelines	can be found or	n the Microb	nology <u>pad</u>	e or the <u>int</u>	ranet	
Tick or insert times required 1 Image: 1000 Image: 10000 Image: 1000 I			INTR	AVENOU	S ANTI	BIOTICS OF	NLY					and the second second	0	RAL ANTIB	IOTICS	ONLY			
Image: Date: Date: Date: V Partine at 48 hours with 1500 00 1 1 1 Partine at 48 hours with 1500 00 1 1 1 1 Benstlyther: YA Agenored 100 00 1 1 1 Benstlyther: YA Agenored 100 00 1 1 1 1 Benstlyther: YA Agenored 100 00 1 1 1 1 1 Benstlyther: YA Agenored 100 00 1 <			Tick or	r insert times	required J	1		No. of Concession, Name			-		Tick or insert tir	nes required L					
	Drug (Approved m	ame):			Date:						Drug (Appr	oved name):		Date:					
Ministry Bills Minis	Disei la	no chart	Otart data	Chan data:	0500						Dose:	Acute:	Start date:	0000-0000	-	-			
Percent at a brane with authors Bood (author) Percent (author) Perce	ing ing	IV	Start said.	Stop date:	1400			-	ASAP		Simo:	Oral	-	1800 - 1400					_
Jenusitives: YM Marcines: Marcine: Marcine: Marc	Silgn;		Review at -	48 hours with	1800				-		P:		Stop date:	1800 - 1800				15	
Instructure: Date:	P;	12	cul	hures .	2200	(Brown or other	-14-17				Indication:		Sensitivites: Y/N	10000 B					_
Under weiter Under weiter<	ndication:	Sen	isilivites; Y/N	approved:	E Y/N	advice:	_				approved:	Y/N	advice:			_			
Start date: Stort date:	rang (Abbuowed us	erner:			Date:			and the second se			rand (white	ovec name):		Date:		7.57		1.	
V Image: State of the set	Dose: Ro	outec	Start date:	Stop date:	1000						Dose:	Poute:	Start date:	0900-0600					_
Bestein at 48 hours with sensitivities: YN Magnetics advices: Magnetic Participation 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000		IV			1400				WEWE		Sign:	Utal	Stop date:	1200-1400					_
Sensitivities: VN Programed: Sensitivities: VN Bandcalor:	Sign:		Review at 4	48 hours with	1800				0	0	P:			2200-2400				3.	
Image: Image:<	P; indication:	le.	Cul	Micro	2200	Pharmecist				-	Micro	104	Sensitivites: Y/N Pharmacist	Constant of the local division of the local				-	_
Norm Norm <th< td=""><td>Drug (Approved or</td><td>amel:</td><td>and Plugs TITY</td><td>approved:</td><td>Date</td><td>advice:</td><td></td><td></td><td></td><td></td><td>approved: Drug (Appro</td><td>rin oved name'r</td><td>advice:</td><td>Data</td><td></td><td></td><td></td><td></td><td>_</td></th<>	Drug (Approved or	amel:	and Plugs TITY	approved:	Date	advice:					approved: Drug (Appro	rin oved name'r	advice:	Data					_
University Start data: Stop date: 1000 Image: Start data: Stop date: Stop	and a description				0600				1.000			and a manned.		Date;					
IV \$400 Image: 1000 Image: 10000 Image: 1000 Image: 1	Dose: Ro	oute:	Start date:	Stop date:	1000			1	CITER D		Dose:	Oral	Start date:	0600 0601					
Review 14 B hours with cultures 1800 Image: Construction Review 14 B hours with approved: YN Review 14 B hours with advice:		IV			1400	-					Sign:	erai	Stop date:	1600 - 1800					-
Sensitivities: Y/N Macro- approved: Prioring activities: V/N Prioring activities: Date: Date: <th< td=""><td>Bign:</td><td></td><td>Review at a</td><td>48 hours with nures</td><td>1800</td><td></td><td></td><td>-</td><td></td><td></td><td>P:</td><td></td><td>Sensibilian: Y/M</td><td>2200 - 2400</td><td></td><td></td><td></td><td></td><td>_</td></th<>	Bign:		Review at a	48 hours with nures	1800			-			P:		Sensibilian: Y/M	2200 - 2400					_
Image: Index of the second state is a second state second state second state is a second state is a second state is	indication:	Sen	silivites: Y/N	Micro	YN	Pharmacist					Micro	VIN	Pharmadist				1		
Openetic in the interval of the interva	Drug (Approved na	ame):		[approved:	Date:	advice:					Drug (Appril	oved name):	auvice:	Date:					
Uk: Start date: Nono Nono Nono Review at 48 hours with cultures 2000 Nono Nono Nono Nono Bensitivities: 2000 Nono Nono Nono Nono Nono W: Start date: Stop date: 1600 Nono Nono Nono V Bensitivities: Y/N Premasist advice: Nono Nono Nono Nono V Start date: Stop date: 1600 Nono Nono Nono Nono V Start date: Stop date: 1600 Nono Nono Nono Nono Nono V Start date: Stop date: 1600 Nono					0600						Doser	Boute	Start data:	CERTS - FORM					_
Berland Baues with cultures Study Stop date: Boot Start date: Stop date:	Dose: Ro	IV	Start date:	Stop date:	1000				ASAP			Oral	onart care:	1200 - 1400	-	-	-		
Industrie 2200 Pharmadia Bensitivities: YN Moro advice: mel: Date: advice: Pharmadia: ute: Date: advice: Pharmadia: V Find catalow: Stant date: (00)-500 Image: Pharmadia: Index: Find catalow: Stant date: (00)-500 Image: Pharmadia: Sign: Stant date: (00)-500 Image: Pharmadia:	Sion		Desting at .	10 hours with	1800				1000		Sign:		Stop date:	1600 - 1800					_
Bensitivites: YN More approved; VN Pre-model approved;	Pj	0.000	cul	Bures.	2200					1.1	Indication:		Sensitivities: Y/N	2200 - 2400					_
Date: Date: <th< td=""><td>indication:</td><td>Sen</td><td>sitivites: Y/N</td><td>Micro approved</td><td>YIN</td><td>Pharmacist advice:</td><td></td><td></td><td>0</td><td>0</td><td>Micro approved:</td><td>YAN</td><td>Pharmacist advice:</td><td></td><td></td><td></td><td></td><td></td><td>_</td></th<>	indication:	Sen	sitivites: Y/N	Micro approved	YIN	Pharmacist advice:			0	0	Micro approved:	YAN	Pharmacist advice:						_
Univ Start date: 000 Image: Start date: 000-500 000-500 000-500 000-500 <td>Drug (Approved na</td> <td>атте):</td> <td></td> <td></td> <td>Date:</td> <td></td> <td></td> <td></td> <td></td> <td></td> <td>Drug (Appro</td> <td>oved name):</td> <td></td> <td>Date:</td> <td></td> <td></td> <td>Martin F</td> <td></td> <td></td>	Drug (Approved na	атте):			Date:						Drug (Appro	oved name):		Date:			Martin F		
IV Instrument	Dose: In-	a dar	Giset data:	Stop date:	0600			-	man		Dose:	Route:	Start date:	0600-2900					
Image: Supervised of	House: Ho	IV	Sherr cane:	Stop date:	1000	-			Castal P			Oral	Concernance of	1200-1400			-		_
cultures 2200 Planmadit Sensitivities: YN More advice: More advice: Planmadit TIONS FOR INTRAVENOUS ANTIBIOTICS MUST BE REVIEWED AFTER 24 - 48 HOURS Planmadit More advice: Planmadit PRESCRIPTIONS FOR INTRAVENOUS ANTIBIOTICS MUST BE REVIEWED AFTER 24 - 48 HOURS PRESCRIPTIONS FOR ORAL ANTIBIOTICS ARE VALUE FOR 5 DAVS ONLY.	Gign:	1000	Review at 4	48 lours with	1800	-					F.		Slop date:	1600 - 1800					
Sanstiklies: YN approved: YN pharmadel advice: Pharmadel advice: Pharmadel advice: Pharmadel advice: TIONS FOR INTRAVENOUS ANTIBIOTICS MUST BE REVIEWED AFTER 24 - 48 HOURS PRESCRIPTIONS FOR ORAL ANTIBIOTICS ARE VALID FOR 5 DAYS ONLY.	P;		cu	itures	2200		1				Indication:		Sensitivites: Y/N						_
TIONS FOR INTRAVENOUS ANTIBIOTICS MUST BE REVIEWED AFTER 24 - 48 HOURS PRESCRIPTIONS FOR ORAL ANTIBIOTICS ARE VALUE FOR 5 DAVIS ONLY	indication:	Sen	istivites: Y/N	Approved:	YIN	Pharmadist advice:					approved:	Y/N	Phamacist advice:						
SWITCH TO ORAL AS SOON AS POSSIBLE	Dose: R: Gign: P: Indication: PRESCRIP	Sulle: IV Sen	Start data: Review at a cut stivites: Y/N OR INTRAVE SW/T	Biop date: Biours wills form approved ENQUS ANT TCH TO ORA	0600 1000 1408 2200 2200 YN IBIOTICS ALAS SOU	Pharmadist advice: MUST BE REVI ON AS POSSIBI	EWEDIAFTER 24	1-48 HOURS	131		Draw: Sign: P: Indication: Micro approved:	Poule: Oral Y/N PRI	Start dete: Stop dete: Bansifvitas: Y/N Pharmacist advice: ESCRIPTIONS FOI	0003-500 1230-500 1600-100 2230-3400 RORAL ANTIBIO	OTICS ARE	VALID FOR	5 DAV	(5.0)	'S ONLY
Date and Dessee Owning .	Date Ta	me		Ding	and Hease	an Omitted		signature			Date	Time		urug and i	season Umith	ed		Signa	ture
ne Drug and Reason Omitted Signature Date Time Drug and Reason Omitted Signature												-							
ne Drug and Reason Omitted Signature Date Time Drug and Reason Omitted Signature																			
Drug and Reason Omitted Signature Date Time Drug and Reason Omitted Signature Image: Ima											-							- 8	

Wrightington, Wigin and Leigh Foundation NHS Trust

Dedicated Antimicrobial Section on drug chart – added within regular prescriptions section

							PRES	CRIPTIC	DNS M	UST BE	REVIE	NED A	ND RE	WRITTE	EN EVE	RY TW	O WEE	KS OR SOONER
EN USE	TER D ONE	ROUTE (ainst tin Only for	AE REQ	UIRED) Y	REG	ULAR	PRE	5CRIP	TION	MC	NTH		YE/	AR		Medicines for Discharge
TIME		DOSE	DATE>															Dr's Sig:
TIVE		START	CHANGE	T														Date: Bleep:
DATE	E			DRUG	G (APPR	OVED	NAME)		OTHE	R INST	RUCTIO	NS	SIGNAT	URE & I	REG NO	PHAR	MACY	To be prescribed for Discharge
ROU	TE	S/C		1														Discharge
INITI	ALS			1	ENC	XAP	ARIN											N
																		Limited Duration
	08																	Days
	13																	e.g. Steroids, Antibiotics
	(18)																	Transcribed/
	22				0	N CO	MPL	TION	OF \	/TE R	ISK A	SSES	SMEN	T PR	ESCR	BE		Prescribed
				1	THR	OMB	OPRO	PHYL	AXIS	ABO	VE IF	NO 0	ONT	RAIN		IONS	5	Pharmacy date:
DATE	E			A	NTIMI(CROBI/	al nan	1E					SIGNAT	URE & I	REG NO	PHAR	MACY	To be prescribed fo
ROU	TE			1					INDIC/	ATION: .								Discharge
INITI	ALS			STATE C		ENCTL A			DURA	TION:								
				STATE C	OURSE L	Elwain A	AND REVI	CITY DATE										Limited Duration
<u> </u>	08			-		-		-										Days
<u> </u>	13					-												e.g. Steroids,
<u> </u>	18					-		<u> </u>										Transcribed/
<u> </u>	72					<u> </u>		<u> </u>								<u> </u>		Prescribed
<u> </u>	~~~					-		-						<u> </u>		<u> </u>		Pharmacy date:
DATI				A	NTIMI	CROBIA	AL NAN	1E					SIGNAT	URF & I	REG NO	PHAR	MACY	To be prescribed for
POL	TE			-					INDIC/	ATION: .								Discharge
INITI,	ALS			STATE C	OURSE L	ength A	ND REVI	ew date	DURA	ΠON:								
																		Limited Duration Days
	80																	
	13																	e.g. Steroids, Antibiotics
	18																	Transcribed/
	22																	Prescribeu
																		Pharmacy date:
DATE	E			DRUG	(APPR	OVED	NAME))	OTHE	R INST	RUCTIO	NS	SIGNAT	URE &	REG NO	PHAR	MACY	To be prescribed for Discharge
ROU	TE																	
INITI	ALS																	
																		Limited Duration Days
	08																	
	13																	e.g. steroids, Antibiotics
	18																	Prescribed
	22									1		1		1		1		1

CHECK ALLERGIES ON FRONT OF THE CHART BEFORE PRESCRIBING AND ADMINISTERING

Barking Havering and Redbridge University Hospitals NHS Trust Drug Chart June 2011: Used with permission from Antimicrobial Stewardship Group

United Lincolnshire Hospitals NHS NHS Trust **EXAMPLE ANTIMICROBIAL SECTIONS OF** PRESCRIPTION CHART Check allergy PATIENT NAME **IV ANTIBIOTIC PRESCRIPTIONS** status BEFORE

			pre	scribing/giving								
Antibiotic (approved name)					Date (d/m)	->					IV Dovi	
Dose	Route IV	Start	Stop	Pharmacy	Time	↓		W	ch		(please ti	ew ick)
					06			evie	wit		Switch to PO Prescribe	
Indication		Guidelines/ Micro						IT T(al s			
		approved?			12			not	OĽ		Continue IV	
Signature			Bleep	Pharm tech	18			8	V /		Stop	
								4	Ι		Signature	Date
Additional info					24							

DoB

PROLO))	ANTIBIO	TICS ON	Check allergy NLY status BEFORE escribing/giving	Cou	rse	ler	ngtl	1 / 1	rev	iew	da	te I	MU	ST	` be	sp	ecif	ïed	
Antibiotic (approved name)					Date (d/m)	÷														
Dose	Route	Start	Stop	Pharmacy	Time	\rightarrow														
Indication		Guidelines/ Micro			08															
		approved?			13															
Signature			Bleep	Pharm tech	18															

		22								
Additional info										

ORAL A	NTIBIO	TIC PRE	SCRIP BEFORE p	check allergy TIONS status rescribing/ giving	PATII	ENT	NAN	ИЕ		DoB	
Antibiotic (approved name)					Date (d/m)	<i>→</i>				5 DAV DEV	
Dose	Route Oral	Start	Stop	Pharmacy	Time	Ļ				5 DAY KEV (please tic	iE w :k)
										Stop	
Indication		Guidelines/ Micro			08					Continue (Re-prescribe)	
		approved?			13						
Signature			Bleep	Pharm tech	18					Total duration	
					22					Signature	Date
Additional info											

				le D-	aule	-	dice	tion									
Den	and need Bush with Cod	um Ohl		AS MO	quire		oica	son	aller	a desta		and a solar	Shee In			the state	
PTU -	and post riush with Sod	um cni	Doese	muşt	be gr	ALL W	in an	medic	auon	aurin	ISSUE 9			s av us	ious r	COLIN	-
und (abbio	ved namej		LOSe	Lakto	<u> </u>	<u> </u>	<u> </u>			<u> </u>		<u> </u>	<u> </u>		<u> </u>		⊢
				Tmo													⊢
Route	Minimal Interval / Instruc	sons	Max Dose Dally	Cose													L
				Route													
Start date	Indication		Pharmacy	_													Г
				- agen													
Sign		Stop da	ste / initials														F
-		· ·		Sign													
Oruc (appro	ved name)		Dose	Date		-	-			-		<u>├</u>	-		-		⊢
				1		-	-			-		├──	-		-	├	⊢
Deute	Minimal Internal / Inclusio	form	Max Doors		-	<u> </u>	-	-	-	-		├	-		<u> </u>		⊢
HOUSE .	Minimal Interval / Instruc	JOI D	Dally	LIDBO	<u> </u>	<u> </u>	<u> </u>			<u> </u>		<u> </u>	<u> </u>		<u> </u>		⊢
				Route								<u> </u>		L	<u> </u>	L	⊢
staft date	indication		Pharmacy	Sign													
Sign		Stop da	ste / Initials	Sen													[
Drug (approv	ved name)		Dose	Date													Γ
				Tmo													t
Route	Minimal Interval / Instruc	Sons	Max Dose	Does		-	-			-		<u>├</u>	-		-		⊢
			Dally	Death	-	<u> </u>	-			-		<u> </u>	-	<u> </u>	<u> </u>	├	⊢
Clast data	Indication		Desmark	10.00	-	<u> </u>	-			-	-	├	-		<u> </u>		⊢
	Indication		rianacy	Sign													
														L		L	⊢
Sign		Stop da	ste / Initials	Sign													
Drug (approv	ved name)		Dose	Date													Г
				Tmo													Г
Route	Minimal Interval / Instruc	Sons	Max Dose	Пово													F
			Dally	Route										\vdash		\vdash	t
Start date	Indication		Pharmacy		\vdash	-	\vdash			-		<u>├</u>	-	<u>├</u>	-	├──	⊢
	The second se			Sign													
		Dia d		<u> </u>	<u> </u>	<u> </u>	<u> </u>			-		<u> </u>	<u> </u>		<u> </u>		⊢
sign		stop of	ne / minais	Sign													
			1-														
Drug (approv	ved name)		Dose	Date													
				Time													
Route	Minimal Interval / Instruc	tions	Max Dose	Пове													
			Lay	Route													Γ
Start date	Indication		Pharmacy	_													F
			· · ·	Sign													
Sion	1	te / Initials	-		-					-	-	-	-	-	-	⊢	
	sign Stop da																1
Dava Jane	and manual		Denne	-	-	-			-			-	-	-		-	⊢
und (approv	rad name)		0050	(lade										<u> </u>		L	1
				Time													
Route	Minimal Interval / Instruct	Sons	Max Dose Dally	Com													
				Route													Γ
Start date	Indication		Pharmacy	-													Γ
				- Aller												1	
	1			-	-	<u> </u>	-					-	-	<u> </u>	-	-	⊢
Sign		Stop da	nte / Initials														

o drugs to be administered noise satiety box complete Chart Noof	In-Patien	t Medicatio	n Admi	nistra	tion R	ecord	Cour	ty Durha	am and	Dar	lingto	NHS
All ellergies / sensitivities - must be completed All ellergies / sensitivities - must be completed basis Data: basis Description of reaction: Data: Data: Data: Data: <t< td=""><td>No drugs to b</td><td>e adminstered</td><td>Charles</td><td></td><td></td><td>· ·</td><td>Nwaya use pi</td><td>tientaddr</td><td>e sa ogn</td><td>aph la</td><td>bel</td><td></td></t<>	No drugs to b	e adminstered	Charles			· ·	Nwaya use pi	tientaddr	e sa ogn	aph la	bel	
All ellergies / sensitivities must be completed Sumane: ione know: Signad: Date: Date of searchain: Signad: Date: Viergies / Sensitivities: Description of reaction: Signature Print name Print name Reap No. Reg. No. Signature Print name Viergies and mittals Calculated creat himp Complanes and mpp medication Calculated creat himp Signature Print name Viergies and mittals Calculated creat himp Camplanes and mpp medication Calculated creat himp Camplanes assessed Y/N Complanes assessed Y/N Community Pharmacy Name and Tot. Date of Initials Date of Initials Date of Initials Date of Initials Date of Initials Date of Initials Date of Initials Date of	uniessallerg)	y box complete	Charting	o o	DT TO	Hospital / N	HS Numbe	r				
Ione known: Signad: Data: Inable to accrtain: Signad: Data: Ister Signad: Data: Liter gles / Sensitivities: Description of reaction: Ward and Site Height (m) Prescriptor Details Consultant Signature Print name Print name Reap No. Right and in Intel	All allerg	ies / sensitivities	s – must be	o comple	hed	Sumame:						
Inside to ascertain: Signod: Date: Description of reaction: Description of reaction: Description of reaction: Date: Date: Date: Date: Date: Date: Date: Benergenergenergenergenergenergenergener	None known:	Signed:		Date:		First Name((s):					
Index: Date: Date: <th< td=""><td>Unable to asce</td><td>ertain: Signed:</td><td></td><td>Date:</td><td></td><td>Address:</td><td></td><td></td><td></td><td></td><td></td><td></td></th<>	Unable to asce	ertain: Signed:		Date:		Address:						
Ultergies / Sensitivities: Description of reaction :: Date of Birth:: Ward and Sile Height (pg) Date of Admission Age Weight (pg) Surface Area (m ²) Date of Admission Signature Print name Geep No. Reg No. Signature Print name Geep No. Reg No. Date of Initials Date of Admission Consultant Date of Birth: Date of Admission Consultant Signature Print name Geep No. Reg No. Date of Initials Date of Admission V/N Date of Birth: Date of Birth: Consultant Date of Admission Consultant Signature Print name Geep No. Reg No. Date of Initials Date of Dirth Consultant Date of Initials Date of Birth: Date of Birth: Date of Dirth Consultant Consultant V/N Date of Birth: Date of Dirth Consultant Date of Birth: Date of Birth: Date of Birth: Date of Dirth Consultant Date of Birth: Date of Birth: Date of Birth: Date of Dirth Consultant Date of Birth: Date of Birth: Date of Birth: Date of Dirth Consultant Date of	Yes: Signed:			Date:								
Prescriber Details Calculated creat in ine (inf) Admission Medication Signature Print name Bite p No. Reg. No. Paint's calculated creat in ine clearance (mis/min) Admission Medication Signature Print name Bite p No. Reg. No. Paint's calculated creat in ine clearance (mis/min) Admission Medication Signature Print name Bite p No. Reg. No. Paint's calculated creating ine clearance (mis/min) Paint's calculated creating ine clearance (mis/min) Paint's calculated creating ine clearance (mis/min) Signature Print name Bite p No. Reg. No. Paint's calculated creating ine clearance (mis/min) Paint's calculated creating ine clearance (mis/min) Date and Initials Date and Initials Date and Initials Modifies Recordination compiles V/N V/N Date and Initials Date and Initials Date and Initials Start date Stop date Stop date Start date Stop date Stop date Stop date Date and Initials Date and Initials Insulfit Chart Insulfit Chart Insulfit Chart Insulfit Chart Date and Initials Insulfit Chart Insulfit Chart Insulfit Chart Insulfit <	Allergies / Ser	sitivities: D	escription (of reaction	n:	Date of Birt	h:					
Age Weight (kg) Surface Area (m ²) Consultant (m ²) Each new prescher on biar down the discontent the detailshies including hises stepping medication signeture Calculate or each hing (elearance (mis/min)) Admission Medication Signeture Print name Bleep No. Reg. No. Date and Initials Date and Initials Other Charls in current use V/N Complance assessed V/N V/N Complance assessed V/N V/N Complance assessed V/N Complance assessed V/N V/N Complance assessed V/N V/N Complance assessed V/N Complance assessed V/N V/N Complance assessed V/N V/N Complance assessed V/N Complance assessed V/N V/N Complance assessed V/N V/N Date and Initials Date and Initials Date and Initials Date and Initials Date and Initials Date and Initials Date and Initials Date and Initials Date and Initials Date and Initials Date and Initials Date and Initials Date an						Ward and Sit	0	Height	(om)	Date	of Adr	nission
that details basice in foruing the asstopping market and the settopping market an						Age	Weight (kg)	Surface (m²)	Area	(Consult	lant
Celearance (mis/min) Dester Initials Signature Print name Bleep No. Pring No. Date and initials Date and initials Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distantiation Image distanti		Prescriber Det	ails		Calculate	d creatinine		Admissi	ion Me	dica	tion	
Signature Other name Signature Signature Prist name Bleep No. Reg. No. Date and Initials Date and Initials Chackod for use Y / N Compliance askings of the second list of t	Each new	prescriber on this chart	should comple	te .	clearance	ce (mis/min)					Delete	Initials
Segret ure Print name Deloy No. Neg. No. Image: Segret ure Image: Segret ure <td< td=""><td>their details b</td><td>and a protection of the s</td><td>ropping medici</td><td>an only</td><td></td><td></td><td>Patient's ow</td><td>in drugs br</td><td>ought in</td><td>1</td><td>Y/N</td><td>-</td></td<>	their details b	and a protection of the s	ropping medici	an only			Patient's ow	in drugs br	ought in	1	Y/N	-
Image: Index or of initials Medicines Reconciliation compilety Y / N Image: Index or of initials Compilance assessed Y / N Image: Index or of initials Date and initials Image: Initial Initia	Signature	Print name	bleep No.	Heg. No.	Data		Checked to	ruse		+	Y/N	
Image: Compliance assessed Y.N Image: Compliance assessed Image: Compliance assessed Image: Compliance assessed Image: Compliance assessed Image: Compliance assessed Image: Compliance assessed Image: Compliance assesses Image: Compliance assesses Image: Compliance assesses Image: Compliance assesses Image: Compliance assesses Image: Compliance assesses Image: Compliance a					Little and I	1025	Medicines P	Reconcillati	on comp	olata	Y/N	
Image: Start date Stop date Image: Start							Compliance	assessed			Y/N	
Image: Community Pharmacy Name and Tail. Image: Community Pharmacy Na							Compliance	ald in plac	9		Y/N	
Other Charts in current use Image: Start data Start data <t< td=""><td></td><td></td><td></td><td></td><td>Date and I</td><td>nitials</td><td>Community</td><td>Pharmacy</td><td>Name a</td><td>and Tel</td><td>L</td><td></td></t<>					Date and I	nitials	Community	Pharmacy	Name a	and Tel	L	
Image: Start date Start date Start date Start date Start date Stort date Image: Start date Image: Start date							01	her Cha	rts in o	urre	nt us	0
Image: Construction of the construction of										Start	date 5	Rop date
Image: Indiana in the indiana indina indiana indina indiana indina indiana indi							Epidural / P	CA			-	
Image: Second					Date and I	nitials	Insulin Cha	t				
Additional Information: Additional Information: Additional Information: Additional Information: Additional Information: Additional Information: Image: Second Secon							Other Rx ch	varts – spec	olly:			
Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system Image: Second system <t< td=""><td></td><td></td><td></td><td></td><td>Addition</td><td>al Information</td><td>1:</td><td></td><td></td><td></td><td>_</td><td></td></t<>					Addition	al Information	1:				_	
Image: Second secon												
Image: Second												
Image: Note of the second s												
Image: Constraint of the constr		+	+									
Date Medicine (approved name) Dose Route Time to be given Prescribe/r's Signature Phamacy Date Time Given by Checked by Image: State of the st												
Prescription for Once-only and Pre-anaesthetic Medication Date Medicine (approved name) Dose Route Time to be given Prescriber's Signature Phamacy Date Time diven given Checked by Image: Signature Phamacy Date Time diven given Image: Signature Phamacy Date Time diven given Checked by Image: Signature Image: Signature Phamacy Date Time diven by Checked by Image: Signature Image: Signature Phamacy Date Time diven by Checked by Image: Signature Image: Signature Phamacy Date Time diven by Checked by Image: Signature Image: Signature Phamacy Date Time diven by Checked by Image: Signature Image: Signature Phamacy Date Time diven by Checked by Image: Signature Image: Signature Phamacy Date Time divention Image: Signature Image: Signature Phamacy Date Time divention Image: Signature Image: Signature Image: Signature Phamacy Date Image: Signature Image: Signature Image: Signature Image: Signature Image: Signature Image: Si												
Date Medicine (approved name) Dose Route Time to be given Prescriber's Signature Phamacy Date Time to given Checked Image: Signature Image: Signature <t< td=""><td></td><td>Pre</td><td>e criation t</td><td>or Once</td><td>-only and</td><td>Pre-ana est h</td><td>ntic Medic</td><td>ation</td><td></td><td>_</td><td></td><td></td></t<>		Pre	e criation t	or Once	-only and	Pre-ana est h	ntic Medic	ation		_		
	Date Medi	cine (approved name) Dose	Route	Time to be given	Prescriber's	Signature	hamacy	Date	Time given	Given by	Checked
											-	
				+	+							
			_									
					1							
			+	<u> </u>	+							
					-					_		
				1	1							

LP55299

1

Pt Name: HOSP no:		Ward: Red:	DOA: Admitte Own ho Transfe Other D	ed From: me D Nursing/C r from another ho 1	are Home 🗆 Ispital 🗖	Site: DMH UHNI Date:	D		IVC & Relevant Current	t Admission Information:
DOB:			Allergie	6:		<u>Consultant</u>	:			
Previous Antibi	otic [)ose/rou	.te	START date	STOP date	Other I	nfor m:	ation		
Current Abx. Drug Name	Curren	nt Antibi	otic 1	Current	Antibiotic 2	Curren	t Antit	piotic 3		
Start Date Dose		<u> </u>					<u>7 7</u>			
Route Taking Oral Meds	IV D/ PO Yes / No	□/Oth (e.g	er 🗆	IV 🗆 / PO 🗆) Yes / No (e	.g	IV⊡/PO) Yes/Noi	□70t (e.g	her 🗆		
Course Length or Reviewdate	Yes□ ReviewD Length:_	No⊡)ate/Sto d	op Date ays	Yes D N Review Dat Length:	lo⊡ te/StopDate days	Yes Review D Length:	No□ ate/S	top Date days		
Where Rx		Reg	>10/7 🗖 Meds 🗆		>10/7 E Reg Meds I		Be	>10/7 □ g_Meds □		
Indication noted on <u>Kardex</u>										
Choice Appropriate If no state	Yes □ Micro □	No 🗆		Yes□ N Micro□	00	Yes□ Micro□	No 🗆			
PMH:										
									Current Diagnosis:	

County Durham and Darlington NHS Trust



Sepsis Audit Tool

				Sepsis Audi	t		
Date:				Patient Initials:			
Ward: Be	ed: Consulta	nt:		Hospital Numb	er:		
Admission Date:				DOB:			
Presenting Comp	aint:						
Is drug allergy bo	completed?	Y / N					
Antibiotic Informa	ation		At	0x1	Abx2		A
			Started:	Started:		Started:	bx 3
	N	lame:					
	Dose + Frequ	ency:					
	R	oute:					
	Indica	ation:	Y / N	Y	(/ N	Y / N	
Ston/review	date/ course le	ongth	 Y / N		/ / N	 Y / N	_
5(0)/10/10	pre	sent?					
	•						
Has the all	ergy check box comple	been eted?	Y / N	Y	(/ N	Y / N	
When was diagn	osis of sepsis m	nade?					_
When was the 1 ^s	^t dose of Abx g	iven?					_
ls treatme	nt in line with s	sepsis	Y / N	Ŷ	(/ N	Y / N	
	guid	eline:	Y / N	Y	(/ N	Y / N	
lf not – is reaso	n why documei	nted?					
					· / • ·		_
Is there an unus	ial dose/freque	ency?	Y/N Y/N	Y Y	(/ N / / N	Y/N Y/N	
lf so – is t	here a docume	ented	171			171	
	rea	ason?					
							_
Are ther	e any missed d	oses?	Y / N	<u> </u>	(/ N	Y / N	
Clinical Markers			Da	ate:		ADDITIONAL	
Pulse	/min		Lactate	n	nmol/I	COIVIIVIEINTS	-
RR	/min		INR				
BP	BP mmHg			n	nls kg-1 hr-1		
Temp	Temp °C						
wcc	WCC x10 ⁹ /L			x	: 10 ⁹ /l		
Neutrophills	eutrophills x10 ⁹ /L B			U	ımol/l		
CRP	mg/L	UTI ((✓)				
Cr				Dysuri	a		
CrCl	mL/min			> Frequenc	у		
BM	mmol/L		A	cute Incontinence	e	Reason for	

A~mental		Loin Pain	Delay?
state			
		LT Catheter	
		Urine Dipstick	
		Nitrate +ve.	
		Leucocyte est. +ve.	