



BSAC Methods for Antimicrobial Susceptibility Testing

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Abstract

There have been considerable changes to the format of the recommendations. The changes are as follows:

- The abstract contains a summary of the changes in the current version
- The footnotes to the tables have been removed and the notations added to the end column. It is hoped that this change will avoid confusion in interpretation.
- Antibiotics have been separated into groups e.g. β -lactams, aminoglycosides etc.
- UTI recommendations have been removed for most agents except for those that are administered solely for the treatment of uncomplicated UTIs or where there are limited recommendations for specific organisms e.g. trimethoprim. For agents that previously had dual recommendations, systemic recommendations remain and the intermediate category can be used for interpretation for UTIs because intermediate susceptibility infers that the infection may respond as the agent is concentrated at the site of infection. This change will also avoid errors in interpretation when an organism is isolated from multiple sites, for example blood and urine.
- The interpretative tables will be separated from the main document and will be available on the web site.

The changes that have been made to the previous version of the recommendations (version 7.1) are as follows:

MIC and zone diameter BPs for trimethoprim, fosfomycin and nitrofurantoin for UTIs (Table 6).

Colistin MIC BPs for *Pseudomonas* spp. (Table 8).

Co-trimoxazole MIC BPs for *Stenotrophomonas maltophilia* (Table 9).

Staphylococci: MIC and zone diameter BPs for clarithromycin, clindamycin, erythromycin, quinupristin/dalfopristin, trimethoprim UTI, nitrofurantoin UTI and rifampicin (Table 10).

Streptococcus pneumoniae: MIC and zone diameter BPs for azithromycin, clarithromycin, co-trimoxazole, erythromycin, linezolid, rifampicin and Telithromycin (Table 11).

Addition of streptomycin recommendations for enterococci (Table 12).

Enterococci: MIC and zone diameter BPs for quinupristin/dalfopristin, nitrofurantoin UTI and trimethoprim UTI (Table 12).

β -Haemolytic streptococci: MIC and zone diameter BPs for azithromycin, clarithromycin, erythromycin and telithromycin (Table 14).

Moraxella catarrhalis: MIC and zone diameter BPs for clarithromycin and erythromycin (Table 15).

Neisseria gonorrhoeae: MIC BPs for azithromycin (Table 16).

Neisseria meningitidis: MIC BPs for chloramphenicol and rifampicin (Table 17).

Haemophilus influenzae: MIC BPs for azithromycin (Table 18).

Bacteroides fragilis: MIC BPs for metronidazole (Table 22).

Bacteroides thetaiotaomicron: MIC BPs for metronidazole (Table 23).

Clostridium perfringens: MIC BPs for metronidazole (Table 24).

Susceptibility testing *Listeria* spp. (Additional Information Number 4)

The acceptable range for NCTC 25923 to a 10 µg tobramycin disc (Table 2 control of antimicrobial susceptibility testing).

Preface

Since the *Journal of Antimicrobial Chemotherapy* Supplement containing the BSAC standardized disc susceptibility testing method was published in 2001, there have been various changes to the recommendations and these have been posted on the BSAC website (<http://www.bsac.org.uk>). One major organizational change has been the harmonisation of MIC breakpoints in Europe.

In 2002 the BSAC agreed to participate with several other European national susceptibility testing committees, namely CA-SFM (Comité de l'Antibiogramme de la Société Française de Microbiologie, France), the CRG (Commissie Richtlijnen Gevoeligheidsbepalingen (The Netherlands), DIN (Deutsches Institut für Normung, Germany), NWGA (Norwegian Working Group on Antimicrobials, Norway) and the SRGA (Swedish Reference Group of Antibiotics, Sweden), in a project to harmonize antimicrobial breakpoints, including previously established values that varied among countries. This work is being undertaken by the European Committee on Antimicrobial Susceptibility Testing (EUCAST) with the support and collaboration of the national committees, and is funded by the European Union, the European Society for Clinical Microbiology and Infectious Diseases (ESCMID) and the national committees, including the BSAC. The review process includes application of more recent techniques, such as pharmacodynamic analysis, and current data, where available, on susceptibility distributions, resistance mechanisms and clinical outcomes as related to *in vitro* tests. There is extensive discussion between EUCAST and the national committees, including the BSAC Working Party on antimicrobial susceptibility testing, and wide consultation on proposals. In the interest of international standardization of susceptibility testing, and the need to update older breakpoints, these developments are welcomed by the BSAC.

The implication of such harmonization is that over time some MIC breakpoints will change slightly and these changes will be reflected, where necessary, in corresponding changes to zone diameter breakpoints in the BSAC disc diffusion method. It is appreciated that changes in the method require additional work for laboratories in changing templates and laboratory information systems, and that the wider use of 'intermediate' categories will add complexity. Nevertheless the benefits of international standardization are considerable, and review of some older breakpoints is undoubtedly warranted.

In line with the European consensus EUCAST MIC breakpoints are defined as follows:

- Clinically resistant: level of antimicrobial susceptibility which results in a high likelihood of therapeutic failure
- Clinically susceptible: level of antimicrobial susceptibility associated with a high likelihood of therapeutic success
- Clinically intermediate: a level of antimicrobial susceptibility associated with uncertain therapeutic effect. It implies that an infection due to the isolate may be appropriately treated in

body sites where the drugs are physically concentrated or when a high dosage of drug can be used; it also indicates a buffer zone that should prevent small, uncontrolled, technical factors from causing major discrepancies in interpretation.

The presentation of MIC breakpoints (mg/L) has also been amended to avoid the theoretical 'gap' inherent in the previous system as follows:

MIC \leq (as previously) MIC breakpoint concentration = organism is susceptible

MIC $>$ (previously \geq) MIC breakpoint concentration = organism is resistant

In practice, this does result in changes to breakpoint systems based on two-fold dilutions.

However, the appearance of the tables will change, e.g. R \geq 16, S \leq 8 will change to R $>$ 8, S \leq 8.

EUCAST MIC breakpoints have to date been agreed for the following agents and are available on the EUCAST web site (www.eucast.org):

Cephalosporins: cefazolin, cefepime, cefotaxime, ceftazidime, ceftriaxone, cefuroxime

Carbapenems: ertapenem, imipenem, meropenem

Monobactams: aztreonam

Fluoroquinolones: ciprofloxacin, levofloxacin, moxifloxacin, norfloxacin, ofloxacin

Aminoglycosides: amikacin, gentamicin, netilmicin, tobramycin

Glycopeptides: teicoplanin, vancomycin

Other agents: linezolid, daptomycin, tigecycline

Disc Diffusion Method for Antimicrobial Susceptibility Testing

1. Preparation of plates

- 1.1 Prepare Iso-Sensitest agar (ISA) (see list of suppliers) or media shown to have the same performance as ISA, according to the manufacturer's instructions. Supplement media for fastidious organisms with 5% defibrinated horse blood or 5% defibrinated horse blood and 20 mg/L β -nicotinamide adenine dinucleotide (NAD) as indicated in Table 1. Use Columbia agar with 2% NaCl for methicillin/oxacillin susceptibility testing of staphylococci.

Table 1: Media and supplementation for antimicrobial susceptibility testing of different groups of organisms

Organisms	Medium
Enterobacteriaceae	ISA
<i>Pseudomonas</i> spp.	ISA
<i>Stenotrophomonas maltophilia</i>	ISA
Staphylococci (tests other than methicillin/oxacillin)	ISA
<i>Staphylococcus aureus</i> (tests using cefoxitin to detect methicillin/oxacillin/cefoxitin resistance)	ISA
Staphylococci (tests using methicillin or oxacillin for the detection of methicillin/oxacillin/cefoxitin resistance)	Columbia agar (<i>see suppliers</i>) with 2% NaCl ¹
Enterococci	ISA
<i>Streptococcus pneumoniae</i>	ISA + 5% defibrinated horse blood ²
α -Haemolytic streptococci	ISA + 5% defibrinated horse blood + 20 mg/L NAD
β -Haemolytic streptococci	ISA + 5% defibrinated horse blood ²
<i>Moraxella catarrhalis</i>	ISA + 5% defibrinated horse blood ²
<i>Haemophilus</i> spp.	ISA + 5% defibrinated horse blood + 20 mg/L NAD
<i>Neisseria gonorrhoeae</i>	ISA + 5% defibrinated horse blood ²
<i>Neisseria meningitidis</i>	ISA + 5% defibrinated horse blood ²
<i>Pasteurella multocida</i>	ISA + 5% defibrinated horse blood + 20 mg/L NAD
<i>Bacteroides fragilis</i> , <i>Bacteroides thetaiotaomicron</i> , <i>Clostridium perfringens</i>	ISA + 5% defibrinated horse blood + 20 mg/L NAD
<i>Campylobacter</i> spp.	ISA + 5% defibrinated horse blood ²
Coryneform organisms	ISA + 5% defibrinated horse blood + 20 mg/L NAD

¹ See Section 8.

² ISA supplemented with 5% defibrinated horse blood + 20mg/L NAD may be used.

- 1.2 Pour sufficient molten agar into sterile Petri dishes to give a depth of 4 mm \pm 0.5 mm (25 mL in 90 mm diameter Petri dishes).
- 1.3 Dry the surface of the agar to remove excess moisture before use. The length of time needed to dry the surface of the agar depends on the drying conditions, e.g. whether a fan-assisted drying cabinet or 'still air' incubator is used, whether plates are dried before storage and storage conditions. **It is important that plates are not over dried.**
- 1.4 Store the plates in vented plastic boxes at 8-10°C prior to use. Alternatively the plates may be stored at 4-8°C in sealed plastic bags. Plate drying, method of storage and storage time should be determined by individual laboratories as part of their quality assurance programme. In particular, quality control tests should confirm that excess surface moisture is not produced and that plates are not over-dried.

2. Selection of control organisms

- 2.1 The performance of the tests should be monitored by the use of appropriate control strains (see section on control of antimicrobial susceptibility testing). The control strains listed (Tables 2a, 2b) include susceptible strains that have been chosen to monitor test performance and resistant strains that can be used to confirm that the method will detect a mechanism of resistance.
- 2.2 Store control strains at -70°C on beads in glycerol broth. Non-fastidious organisms may be stored at -20°C . Two vials of each control strain should be stored, one for an 'in-use' supply, the other for archiving.
- 2.3 Every week subculture a bead from the 'in-use' vial on to appropriate non-selective media and check for purity. From this pure culture, prepare one subculture on each of the following 5 days. For fastidious organisms that will not survive on plates for 5/6 days, subculture the strain daily for no more than 6 days.

Table 2a: Susceptible control strains or control strains with low-level resistance that have been chosen to monitor test performance of antimicrobial susceptibility testing

Organism	Strain		Characteristics
	Either	Or	
<i>Escherichia coli</i>	NCTC 12241 (ATCC 25922)	NCTC 10418	Susceptible
<i>Staphylococcus aureus</i>	NCTC 12981 (ATCC 25923)	NCTC 6571	Susceptible
<i>Pseudomonas aeruginosa</i>	NCTC 12934 (ATCC 27853)	NCTC 10662	Susceptible
<i>Enterococcus faecalis</i>	NCTC 12697 (ATCC 29212)		Susceptible
<i>Haemophilus influenzae</i>	NCTC 11931		Susceptible
<i>Streptococcus pneumoniae</i>	NCTC 12977 (ATCC 49619)		Low-level resistant to penicillin
<i>Neisseria gonorrhoeae</i>	NCTC 12700 (ATCC 49226)		Low-level resistant to penicillin
<i>Pasteurella multocida</i>	NCTC 8489		Susceptible
<i>Bacteroides fragilis</i>	NCTC 9343 (ATCC 25285)		Susceptible
<i>Bacteroides thetaiotaomicron</i>	ATCC 29741		Susceptible
<i>Clostridium perfringens</i>	NCTC 8359 (ATCC 12915)		Susceptible

Table 2b: Control strains with a resistance mechanism that can be used to confirm that the method will detect resistance.

Organism	Strain	Characteristics
<i>Escherichia coli</i>	NCTC 11560	TEM-1 β -lactamase-producer
<i>Staphylococcus aureus</i>	NCTC 12493	<i>MecA</i> positive, methicillin resistant
<i>Haemophilus influenzae</i>	NCTC 12699 (ATCC 49247)	Resistant to β -lactams (β -lactamase-negative)

3. Preparation of inoculum

The inoculum should give semi-confluent growth of colonies after overnight incubation. Use of an inoculum that yields semi-confluent growth has the advantage that an incorrect inoculum can easily be observed. A denser inoculum will result in reduced zones of inhibition and a lighter inoculum will have the opposite effect. The following methods reliably give semi-confluent growth with most isolates.

NB. Other methods of obtaining semi-confluent growth may be used if they are shown to be equivalent to the following.

3.1 Comparison with a 0.5 McFarland standard

3.1.1 Preparation of the 0.5 McFarland standard

Add 0.5 mL of 0.048 M $\text{BaCl}_2 \cdot 2\text{H}_2\text{O}$ (1.17% w/v $\text{BaCl}_2 \cdot 2\text{H}_2\text{O}$) to 99.5 mL of 0.18 M H_2SO_4 (1% w/v) with constant stirring. Thoroughly mix the suspension to ensure that it is even.

Using matched cuvettes with a 1 cm light path and water as a blank standard, measure

the absorbance in a spectrophotometer at a wavelength of 625 nm. The acceptable absorbance range for the standard is 0.08-0.13. Distribute the standard into screw-cap tubes of the same size and volume as those used in growing the broth cultures. Seal the tubes tightly to prevent loss by evaporation. Store protected from light at room temperature. Vigorously agitate the turbidity standard on a vortex mixer before use. Standards may be stored for up to six months, after which time they should be discarded. Prepared standards can be purchased (See list of suppliers), but commercial standards should be checked to ensure that absorbance is within the acceptable range as indicated above.

- 3.1.2 *Inoculum preparation by the growth method* (for non-fastidious organisms, e.g. Enterobacteriaceae, *Pseudomonas* spp. and staphylococci)
Touch at least four morphologically similar colonies (when possible) with a sterile loop. Transfer the growth into Iso-Sensitest broth or an equivalent that has been shown not to interfere with the test. Incubate the broth, with shaking at 35-37°C, until the visible turbidity is equal to or greater than that of a 0.5 McFarland standard.
- 3.1.3 *Inoculum preparation by the direct colony suspension method* (the method of choice for fastidious organisms, i.e. *Haemophilus* spp., *Neisseria gonorrhoeae*, *Neisseria meningitidis*, *Moraxella catarrhalis*, *Streptococcus pneumoniae*, α and β -haemolytic streptococci, *Clostridium perfringens*, *Bacteroides fragilis*, *Bacteroides thetaiotaomicron*, *Campylobacter* spp., *Pasteurella multocida* and Coryneform organisms). Colonies are taken directly from the plate into Iso-Sensitest broth (or equivalent) or sterile distilled water. The density of the suspension should match or exceed that of a 0.5 McFarland standard.

NB. With some organisms production of an even suspension of the required turbidity is difficult and growth in broth, if possible, is a more satisfactory option.

- 3.1.4 *Adjustment of the organism suspension to the density of a 0.5 McFarland standard*
Adjust the density of the organism suspension to equal that of a 0.5 McFarland standard by adding sterile distilled water. To aid comparison, compare the test and standard suspensions against a white background with a contrasting black line.

NB. Suspension should be used within 15 min.

- 3.1.5 *Dilution of suspension in distilled water before inoculation*

Dilute the suspension (density adjusted to that of a 0.5 McFarland standard) in distilled water as indicated in Table 3.

Table 3: Dilution of the suspension (density adjusted to that of a 0.5 McFarland standard) in distilled water

Dilute 1:100	Dilute 1:10	No dilution
β -Haemolytic streptococci	Staphylococci	<i>Neisseria gonorrhoeae</i>
Enterococci	<i>Serratia</i> spp.	<i>Campylobacter</i> spp.
Enterobacteriaceae	<i>Streptococcus pneumoniae</i>	
<i>Pseudomonas</i> spp.	<i>Neisseria meningitidis</i>	
<i>Stenotrophomonas maltophilia</i>	<i>Moraxella catarrhalis</i>	
<i>Acinetobacter</i> spp.	α -haemolytic streptococci	
<i>Haemophilus</i> spp.	<i>Clostridium perfringens</i>	
<i>Pasteurella multocida</i>	Coryneform organisms	
<i>Bacteroides fragilis</i>		
<i>Bacteroides thetaiotaomicron</i>		

NB. These suspensions should be used within 15 min of preparation.

3.2 Photometric standardization of turbidity of suspensions

A photometric method of preparing inocula was described by Moosdeen *et al* (1988)¹ and from this the following simplified procedure has been developed. The spectrophotometer must have a cell holder for 100 x 12 mm test tubes. A much simpler photometer would also probably be acceptable. The 100 x 12 mm test tubes could also be replaced with another tube/cuvette system if required, but the dilutions would need to be recalibrated.

3.2.1 Suspend colonies (touch 4-5 when possible) in 3 mL distilled water or broth in a 100 x 12 mm glass tube (note that tubes are not reused) to give just visible turbidity. It is essential to get an even suspension.

NB. These suspensions should be used within 15 min of preparation.

3.2.2 Zero the spectrophotometer with a sterile water or broth blank (as appropriate) at a wavelength of 500 nm and measure the absorbance of the bacterial suspension.

3.2.3 From table 4 select the volume to transfer (with the appropriate fixed volume micropipette) to 5 mL sterile distilled water.

3.2.4 Mix the diluted suspension to ensure that it is even

NB. Suspension should be used within 15 min. of preparation

Table 4: Dilution of suspensions of test organisms according to absorbance reading

Organisms	Absorbance reading at 500 nm	Volume (μ L) to transfer to 5 mL sterile distilled water
Enterobacteriaceae	0.01 - 0.05	250
Enterococci	>0.05 - 0.1	125
<i>Pseudomonas</i> spp.	>0.1 - 0.3	40
Staphylococci	>0.3 - 0.6	20
	>0.6 - 1.0	10
<i>Haemophilus</i> spp.	0.01 - 0.05	500
Streptococci	>0.05 - 0.1	250
Miscellaneous fastidious Organisms	>0.1 - 0.3	125
	>0.3 - 0.6	80
	>0.6 - 1.0	40

NB. As spectrophotometers may differ, it may be necessary to adjust the dilutions slightly to achieve semi-confluent growth with any individual set of laboratory conditions.

3.3 Direct antimicrobial susceptibility testing of urine specimens and blood cultures

Direct susceptibility testing is not advocated as the control of inoculum is very difficult. Direct testing is, however, undertaken in many laboratories in order to provide more rapid test results. The following methods have been recommended by laboratories that use the BSAC method and will achieve the correct inoculum size for a reasonable proportion of infected urines and blood cultures. If the inoculum is not correct (i.e. growth is not semi-confluent) or the culture is mixed, the test must be repeated.

3.3.1 Urine specimens

3.3.1.1 Method 1

Thoroughly mix the urine specimen, then place a 10 μ L loop of urine in the centre of the susceptibility plate and spread evenly with a dry swab.

3.3.1.2 Method 2

Thoroughly mix the urine specimen, then dip a sterile cotton-wool swab in the urine and remove excess by turning the swab against the inside of the container. Use the swab to make a cross in the centre of the susceptibility plate and spread evenly with another sterile dry swab. If only small numbers of organisms are seen in microscopy, the initial cotton-wool swab may be used to inoculate and spread the susceptibility plate.

3.3.2 Positive blood cultures

The method depends on the Gram reaction of the infecting organism.

3.3.2.1 Gram-negative bacilli.

Using a venting needle, place one drop of the blood culture in 5 mL of sterile water, then dip a sterile cotton-wool swab in the suspension and remove excess by turning the swab against the inside of the container. Use the swab to spread the inoculum evenly over the surface of the susceptibility plate.

3.3.2.2 Gram-positive organisms.

It is not always possible accurately to predict the genera of Gram-positive organisms from the Gram's stain. However, careful observation of the morphology, coupled with clinical information, should make an "educated guess" correct most of the time.

Staphylococci and enterococci.

Using a venting needle, place three drops of the blood culture in 5 mL of sterile water, then dip a sterile cotton-wool swab in the suspension and remove excess by turning the swab against the inside of the container. Use the swab to spread the inoculum evenly over the surface of the susceptibility plate.

Pneumococci, "viridans" streptococci and diptheroids.

Using a venting needle, place one drop of the blood culture in the centre of a susceptibility plate, and spread the inoculum evenly over the surface of the plate.

4. Inoculation of agar plate

Use the adjusted suspension within 15 min to inoculate plates by dipping a sterile cotton-wool swab into the suspension and remove the excess liquid by turning the swab against the side of the container. Spread the inoculum evenly over the entire surface of the plate by swabbing in three directions. Allow the plate to dry before applying discs.

NB. If inoculated plates are left at room temperature for extended times before the discs are applied, the organism may begin to grow, resulting in reduced zones of inhibition. Discs should therefore be applied to the surface of the agar within 15 min of inoculation.

5. Antimicrobial discs

Refer to interpretation tables 6-23 for the appropriate disc contents for the organisms tested.

5.1 Storage and handling of discs.

Loss of potency of agents in discs will result in reduced zones of inhibition. To avoid loss of potency due to inadequate handling of discs the following are recommended:

- 5.1.1 Store discs in sealed containers with a desiccant and protected from light (this is particularly important for some light-susceptible agents such as metronidazole, chloramphenicol and the quinolones).
- 5.1.2 Store stocks at -20°C except for drugs known to be unstable at this temperature. If this is not possible, store discs at <8°C.
- 5.1.3 Store working supplies of discs at <8°C.
- 5.1.4 To prevent condensation, allow discs to warm to room temperature before opening containers.
- 5.1.5 Store disc dispensers in sealed containers with an indicating desiccant.
- 5.1.6 Discard discs on the expiry date shown on the side of the container.

5.2 Application of discs

Discs should be firmly applied to the dry surface of the inoculated susceptibility plate. The contact with the agar should be even. A 90 mm plate will accommodate six discs without unacceptable overlapping of zones.

6. Incubation

If the plates are left for extended times at room temperature after discs are applied, larger zones of inhibition may be obtained compared with zones produced when plates are incubated immediately. Plates should therefore be incubated within 15 min of disc application.

6.1 Conditions of incubation

Incubate plates under conditions listed in table 5.

Table 5: Incubation conditions for antimicrobial susceptibility tests on various organisms

Organisms	Incubation conditions
Enterobacteriaceae	35-37°C in air for 18-20 h
<i>Acinetobacter</i> spp.	35-37°C in air for 18-20 h
<i>Pseudomonas</i> spp.	35-37°C in air for 18-20 h
<i>Stenotrophomonas maltophilia</i>	30°C in air for 18-20 h
Staphylococci (other than methicillin/oxacillin/cefoxitin)	35-37°C in air for 18-20 h
<i>Staphylococcus aureus</i> using cefoxitin for the detection of methicillin/oxacillin/cefoxitin resistance	35°C in air for 18-20 h
Staphylococci using methicillin or oxacillin to detect resistance	30°C in air for 24 h
<i>Moraxella catarrhalis</i>	35-37°C in air for 18-20 h
α -Haemolytic streptococci	35-37°C in 4-6% CO ₂ in air for 18-20 h
β -Haemolytic streptococci	35-37°C in air for 18-20 h
Enterococci	35-37°C in air for 24 h ¹
<i>Neisseria meningitidis</i>	35-37°C in 4-6 % CO ₂ in air for 18-20 h
<i>Streptococcus pneumoniae</i>	35-37°C in 4-6 % CO ₂ in air for 18-20 h
<i>Haemophilus</i> spp.	35-37°C in 4-6 % CO ₂ in air for 18-20 h
<i>Neisseria gonorrhoeae</i>	35-37°C in 4-6 % CO ₂ in air for 18-20 h
<i>Pasteurella multocida</i>	35-37°C in 4- 6% CO ₂ in air for 18-20 h
Coryneform organisms	35-37°C in 4-6% CO ₂ in air for 18-20 h
<i>Campylobacter</i> spp.	35-37°C in microaerophilic conditions for 18-20 h
<i>Bacteroides fragilis</i> , <i>Bacteroides thetaiotaomicron</i> , <i>Clostridium perfringens</i>	35-37°C in 10% CO ₂ /10% H ₂ /80% N ₂ for 18-20 h (anaerobic cabinet or jar)

¹It is essential that plates are incubated for at least 24 h before reporting a strain as susceptible to vancomycin or teicoplanin.

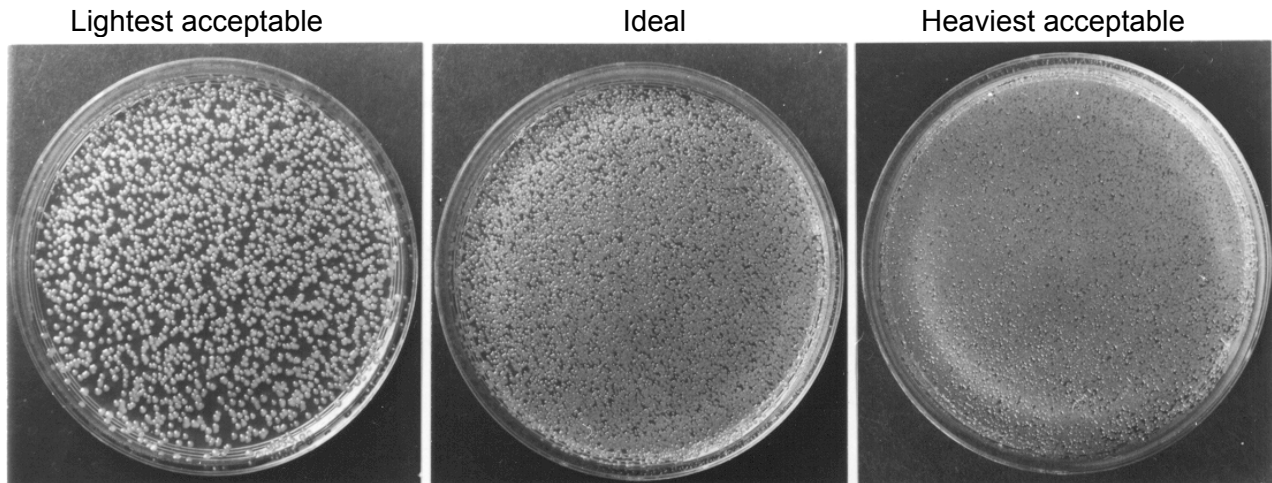
NB. Stacking plates too high in the incubator may affect results owing to uneven heating of plates. The efficiency of heating of plates depends on the incubator and the racking system used. Control of incubation, including height of plate stacking, should therefore be part of the laboratory's Quality Assurance programme.

7. Measuring zones and interpretation of susceptibility

7.1 Acceptable inoculum density

The inoculum should give semi-confluent growth of colonies on the susceptibility plate, within the range illustrated in Figure 1.

Figure 1: Acceptable inoculum density range for a Gram-negative rod



7.2 Measuring zones

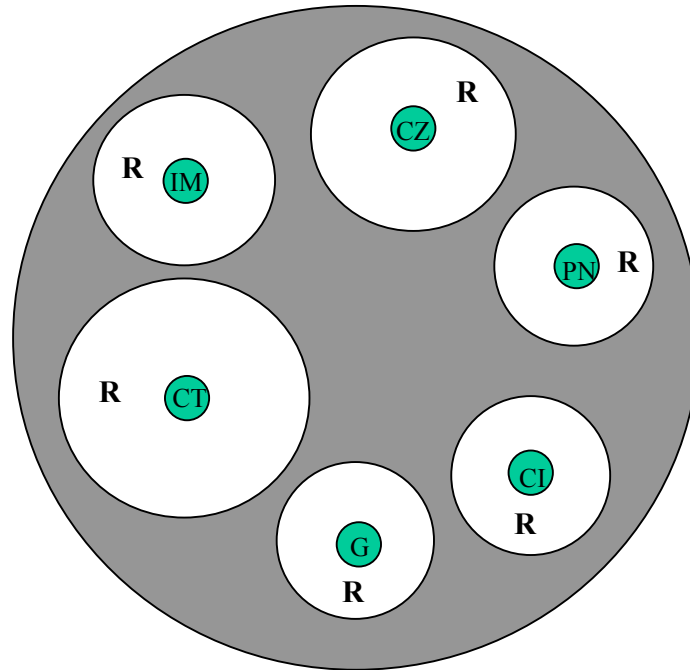
- 7.2.1 Measure the diameters of zones of inhibition to the nearest millimetre (zone edge should be taken as the point of inhibition as judged by the naked eye) with a ruler, callipers or an automated zone reader.
- 7.2.2 Tiny colonies at the edge of the zone, films of growth as a result of the swarming of *Proteus* spp. and slight growth within sulphonamide or trimethoprim zones should be ignored.
- 7.2.3 Colonies growing within the zone of inhibition should be subcultured and identified and the test repeated if necessary.
- 7.2.4 When using cefoxitin for the detection of methicillin/oxacillin/cefoxitin resistance in *S. aureus*, measure the obvious zone, taking care to examine zones carefully in good light to detect minute colonies that may be present within the zone of inhibition (see Figure 3)
- 7.2.5 Confirm that the zone of inhibition for the control strain falls within the acceptable ranges in Tables 20-23 before interpreting the test (see section on control of the disc diffusion method).

7.3 Use of templates for interpreting zone diameters

A template may be used for interpreting zone diameters (see Figure 2). A program for preparing templates is available from the BSAC (<http://www.bsac.org.uk>).

The test plate is placed over the template and the zones of inhibition are examined in relationship to the template zones. If the zone of inhibition of the test strain is within the area marked with an 'R', the organism is resistant. If the zone of inhibition is equal to or larger than the marked area, the organism is susceptible.

Figure 2: Template for interpreting zone diameters



8. Methicillin/oxacillin/cefoxitin testing of staphylococci

Methicillin susceptibility testing is difficult with some strains. Expression of resistance is affected by test conditions and resistance is often heterogeneous, with only a proportion of cells showing resistance. Adding NaCl or lowering incubation temperatures increases the proportion of cells showing resistance. Methicillin susceptibility testing of coagulase-negative staphylococci is further complicated as some strains do not grow well on media containing NaCl and are often slower-growing than *Staphylococcus aureus*. Detection of methicillin resistance in coagulase-negative staphylococci may require incubation for 48 h.

8.1 Method for detection of methicillin/oxacillin resistance in *S. aureus* and coagulase-negative staphylococci

8.1.1 Medium

Prepare Columbia (See list of suppliers) or Mueller-Hinton agar (See list of suppliers) following the manufacturer's instructions and add 2% NaCl. After autoclaving, mix well to distribute the sodium chloride. Pour plates to give a depth of 4 mm (\pm 0.5 mm) in a 90 mm sterile Petri dish (25 ml). Dry and store plates as previously described (section 1).

8.1.2 Inoculum

Prepare inoculum as previously described (section 3).

8.1.3 Control

Susceptible control strains (*Staphylococcus aureus* ATCC 25923 or NCTC 6571) test the reliability of disc content.

Staphylococcus aureus NCTC 12493 is a methicillin resistant strain and is used to check that the test will detect resistant organisms (although no strain can be representative of all the MRSA types in terms of their response to changes in test conditions).

8.1.4 Discs

Place a methicillin 5 µg or an oxacillin 1 µg disc on to the surface of inoculated agar. Discs should be stored and handled as previously described (section 5).

8.1.5 Incubation

Incubate plates for 24 h at 30°C.

8.1.6 Zone measurement

Measure zone diameters (mm) as previously described (section 7).

Examine zones carefully in good light to detect colonies, which may be minute, in zones. If there is suspicion that the colonies growing within zones are contaminants they should be identified and the isolate re-tested for resistance to methicillin/oxacillin if necessary.

8.1.7 Interpretation

For both methicillin and oxacillin interpretation is as follows:

Susceptible = ≥ 15 mm diameter, resistant = ≤ 14 mm diameter.

NB. Hyper-production of β -lactamase does not confer clinical resistance to penicillinase-resistant penicillins and such isolates should be reported susceptible to methicillin/oxacillin. Some hyper-producers of β -lactamase give zones within the range of 7-14 mm and, if possible, such isolates should be checked by a PCR method for *mecA* or by a latex agglutination test for PBP2a. Increase in methicillin/oxacillin zone size in the presence of clavulanic acid is not a reliable test for hyper-producers of β -lactamase as zones of inhibition with some MRSA also increase in the presence of clavulanic acid. Rarely, hyper-producers of β -lactamase give no zone in this test and would therefore not be distinguished from MRSA.

8.2 Detection of methicillin/oxacillin/cefoxitin resistance in *Staphylococcus aureus* by use of cefoxitin as the test agent

8.2.1 Medium

Prepare Iso-Sensitest agar as previously described (section 1).

8.2.2 Inoculum

Prepare inoculum as previously described (section 3).

8.2.3 Control

Use control strains as previously described (section 8.1.3).

8.2.4 Discs

Place a 10 µg cefoxitin disc on the surface of inoculated agar.

Discs should be stored and handled as previously described (section 5).

8.2.5 Incubation

Incubate plates at 35°C for 18-20 h.

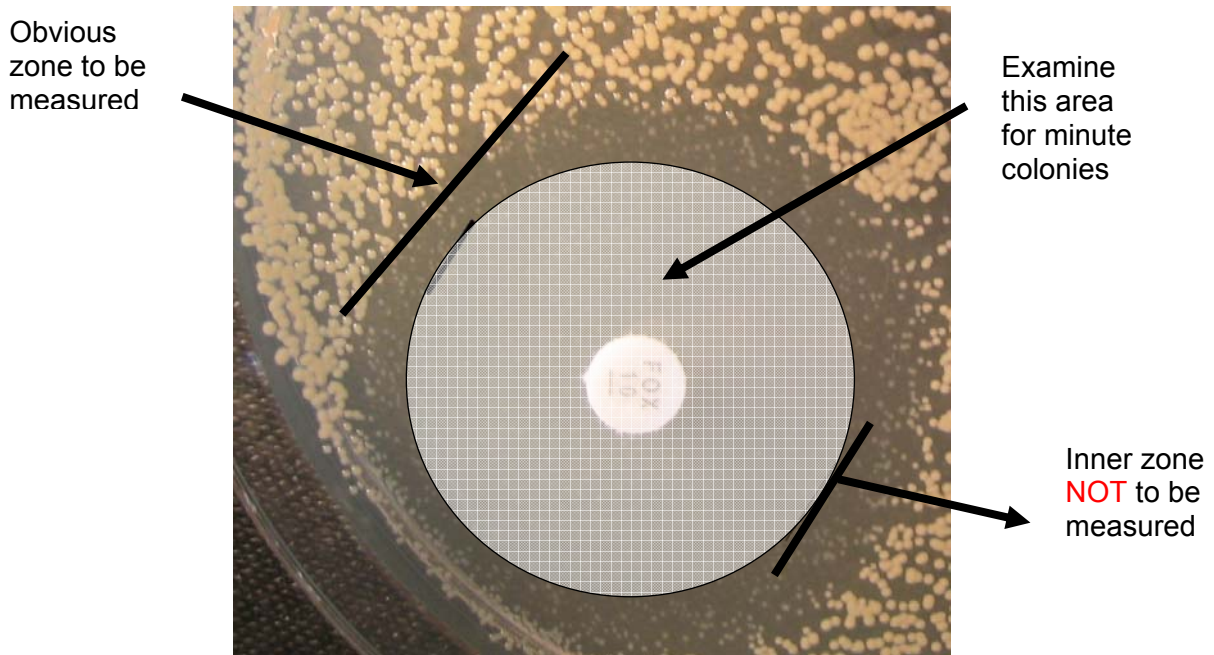
NB. It is important that the temperature does not exceed 36°C, as tests incubated at higher temperatures are less reliable.

8.2.6 Zone measurement

Measure zone diameters as previously described (section 7), reading the obvious zone edge (see Figure 3).

Examine zones carefully in good light to detect colonies, which may be minute, in zones. If there is suspicion that the colonies growing within zones are contaminants they should be identified and the isolate re-tested for resistance to cefoxitin if necessary.

Figure 3: Reading cefoxitin zones of inhibition with *Staphylococcus aureus*



8.2.7 Interpretation

Susceptible = ≥ 22 mm diameter, resistant = ≤ 21 mm diameter.

NB. Hyper-production of β -lactamase does not confer clinical resistance to penicillinase-resistant penicillins and such isolates should be reported susceptible to cefoxitin. Hyper-producers of β -lactamase give zones within the ranges of the susceptible population.

Table 6. MIC and zone diameter breakpoints for Enterobacteriaceae (including *Salmonella* and *Shigella* spp.)

Comments 1-6 relate to urinary tract infections (UTIs) only.

¹UTI recommendations are for organisms associated with uncomplicated urinary infections only. For complicated UTI systemic recommendations should be used.

²If an organism is isolated from multiple sites, for example from blood and urine, interpretation of susceptibility should be made with regard to the systemic site (e.g., if the blood isolate is resistant and the urine isolate susceptible, both should be reported resistant irrespective of the results obtained using interpretative criteria for urine isolates).

³For agents not listed, criteria given for systemic isolates may be used for urinary tract isolates. Intermediate susceptibility infers that the infection may respond as the agent is concentrated at the site of infection.

⁴Direct susceptibility tests on urine samples may be interpreted only if the inoculum gives semi-confluent growth.

⁵In the absence of definitive organism identification, use the recommendations most appropriate for the presumptive identification, accepting that on some occasions the interpretation may be incorrect. A more cautious approach is to use the systemic recommendations.

⁶Coliforms = On-line Medical Dictionary March 2000: "A common name for *E. coli* that is used as an indicator of faecal contamination of water, measured in terms of Coliform count. Occasionally used to refer to all lactose fermenting bacteria."

Table 6 Antibiotic	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Aminoglycosides								
Amikacin	16	16	8	30	15	16-18	19	<i>Salmonella</i> spp. should be reported resistant to these agents, irrespective of susceptibility testing result, as they are inactive against <i>Salmonella</i> spp. <i>in vivo</i> . Individual aminoglycoside agents must be tested; susceptibility to other aminoglycosides cannot be inferred from the gentamicin result and <i>vice versa</i> . For streptomycin, the zone diameter breakpoints are valid only for <i>Escherichia coli</i> , <i>Klebsiella</i> spp. and <i>Proteus mirabilis</i> .
Gentamicin	4	4	2	10	16	17-19	20	
Tobramycin	4	4	2	10	17	18-20	21	
Streptomycin	8	-	8	10	12	-	13	

Table 6	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Penicillins								
Amoxicillin	16	16	8	10	11	12-14	15	These interpretative standards apply only to <i>Escherichia coli</i> , <i>Salmonella</i> spp. and <i>Proteus mirabilis</i> . They do not apply to species that have chromosomal penicillinases (<i>Klebsiella</i> spp.) or those that typically have inducible AmpC enzymes (e.g. <i>Enterobacter</i> spp., <i>Citrobacter</i> spp. and <i>Serratia</i> spp.). The identification of Enterobacteriaceae to species level is essential before applying expert rules for the interpretation of susceptibility.
Ampicillin	16	16	8	10	11	12-14	15	
Co-amoxiclav	16	16	8	20/10	11	12-14	15	
Mezlocillin	16	-	16	75	21	-	22	These interpretative criteria are for <i>E. coli</i> , <i>Klebsiella</i> spp. and <i>P. mirabilis</i> only. Isolates of <i>Escherichia coli</i> and <i>Klebsiella</i> spp. that produce ESBLs often appear susceptible to mecillinam <i>in vitro</i> but clinical efficacy against these organisms is unproven.
Mecillinam UTI ¹⁻⁶	8	-	8	10	13	-	14	
Piperacillin	16	-	16	75	23	-	24	
Piperacillin/tazobactam	16	-	16	75/10	21	-	22	
Temocillin	8	-	8	30	19	-	20	The distribution of zone diameters for ESBL and AmpC producers straddles the breakpoint. Organisms that appear resistant by disc diffusion should have resistance confirmed by MIC determination.
Temocillin UTI ¹⁻⁶	32	-	32	30	11	-	12	These interpretative criteria are for <i>E. coli</i> , <i>Klebsiella</i> spp., <i>P. mirabilis</i> and coliforms only.
Ticarcillin/clavulanate	16	-	16	75/10	20	-	21	

Table 6	MIC breakpoint (mg/L)			Disc content (μg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S \leq		R \leq	I	S \geq	
Cephalosporins								
Cefaclor	1	-	1	30	34	-	35	
Cefalexin UTI ¹⁻⁶	32	-	32	30	15	-	16	These interpretative criteria are for <i>E. coli</i> and <i>Klebsiella</i> spp. only. Cefalexin results may be used to report susceptibility to cefadroxil.
Cefalexin UTI ¹⁻⁶	32	-	32	30	11	-	12	These interpretative criteria are for <i>P. mirabilis</i> only. Cefalexin results may be used to report susceptibility to cefadroxil.
Cefamandole	8	-	8	30	19	-	20	Zone diameter breakpoints are valid only for <i>Escherichia coli</i> , <i>Klebsiella</i> spp. and <i>Proteus mirabilis</i> . The MIC breakpoints have been adjusted to take account of the MIC distribution for the population lacking a mechanism of resistance.
Cefepime	8	2-8	1	30	26	27-31	32	
Cefixime	1	-	1	5	19	-	20	
Cefoperazone	4	-	4	30	24	-	25	Zone diameter breakpoints are valid only for <i>Escherichia coli</i> , <i>Klebsiella</i> spp. and <i>Proteus mirabilis</i> .
Cefotaxime	2	2	1	30	29	-	30	The zone diameters relate to an MIC breakpoint of 1 mg/L as no data for the intermediate category are currently available.
Cefotetan	4	-	4	30	23	-	24	Zone diameter breakpoints are valid only for <i>Escherichia coli</i> , <i>Klebsiella</i> spp. and <i>Proteus mirabilis</i> .
Cefoxitin	8	-	8	30	19	-	20	The MIC breakpoints have been adjusted to take account of the MIC distribution for the population lacking a mechanism of resistance.
Cefpirome	1	-	1	20	24	-	25	

Table 6	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Cephalosporins cont								
Cefpodoxime	1	-	1	10	19	-	20	For ESBL detection, all Enterobacteriaceae isolates should be tested with cefpodoxime or both cefotaxime (or ceftriaxone) and ceftazidime. Enterobacteriaceae with resistance to cefpodoxime, ceftriaxone, cefotaxime or ceftazidime should be tested for the presence of ESBLs. Organisms inferred to have ESBLs should be reported resistant to all penicillins (except temocillin) and cephalosporins, including the fourth-generation cephalosporins cefepime and ceftiprome. For serious infections, carbapenems (imipenem, meropenem, doripenem and ertapenem) are the treatment of choice. Organisms with cefpodoxime zone diameters of < 20 mm have a substantive mechanism of resistance. Organisms with zone diameters of 21-25 mm are uncommonly ESBL-producers and may require further investigation.
Ceftazidime	8	2-8	1	30	17	18-29	30	
Ceftibuten	1	-	1	10	27	-	28	
Ceftizoxime	1	-	1	30	29	-	30	
Ceftriaxone	2	2	1	30	23	24-27	28	
Cefuroxime (axetil)	1	-	1	30	24	-	25	<i>Salmonella</i> spp. should be reported resistant to these agents, irrespective of susceptibility testing result, as they are inactive <i>in-vivo</i> . For parenteral cefuroxime the breakpoint pertains to a dosage of 1.5 g three times a day and to <i>E. coli</i> , <i>Klebsiella</i> spp. and <i>P. mirabilis</i> only.
Cefuroxime (parenteral)	8	-	8	30	19	-	20	
Cefalothin	8	-	8	30	26	-	27	The MIC breakpoints have been adjusted to take account of the MIC distribution for the population lacking a mechanism of resistance.
Cefradine	8	-	8	30	11	-	12	

Table 6	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Carbapenems								
Doripenem	4	2-4	1	10	18	19-23	24	Detection of carbapenem resistance is best achieved by an MIC method on Mueller Hinton agar. <i>Proteus</i> spp. and <i>Morganella morganii</i> are considered poor targets for imipenem.
Ertapenem	1	1	0.5	10	15	16-27	28	
Imipenem	8	4-8	2	10	16	17-20	21	
Meropenem	8	4-8	2	10	19	20-26	27	
Other β-Lactams								
Aztreonam	8	2-8	1	30	22	23-27	28	The MIC breakpoint has been set to ensure that ESBL-producers with MIC values of 4 mg/L are not interpreted as susceptible to this agent.
Quinolones								
Ciprofloxacin	1	1	0.5	1	16	17-19	20	Isolates of <i>Escherichia coli</i> and <i>Klebsiella</i> spp. with ciprofloxacin MICs of 0.25 and 0.5 mg/L may be reported as resistant. These MICs are higher than those for the 'wild susceptible' populations for the species and may indicate a mechanism of resistance with clinical significance. For ciprofloxacin, there is clinical evidence to indicate a poor response in systemic infections caused by <i>Salmonella</i> spp. with reduced susceptibility to fluoroquinolones (ciprofloxacin MICs 0.125-1 mg/L). It is recommended that the ciprofloxacin MIC should be determined for all invasive <i>Salmonellae</i> infections.
Gatifloxacin	1	-	1	2	19	-	20	
Gemifloxacin	0.25	-	0.25	1	19	-	20	
Levofloxacin	2	2	1	1	13	14-16	17	
Moxifloxacin	1	1	0.5	1	16	17-19	20	
Nalidixic acid UTI ¹⁻⁶	16	-	16	30	17	-	18	These interpretative criteria are for <i>E. coli</i> , <i>Klebsiella</i> spp., <i>P. mirabilis</i> and coliforms only.
Norfloxacin UTI ¹⁻⁶	4	-	4	2	15	-	16	
Ofloxacin	1	1	0.5	5	25	26-28	29	

Table 6	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Miscellaneous antibiotics								
Azithromycin	-	-	-	-	-	-	-	Azithromycin has been used in the treatment of infections with <i>S. typhi</i> (MIC ≤16 mg/L for wild type isolates) and some enteric infections.
Chloramphenicol	8	-	8	30	20	-	21	
Colistin	4	-	4	25	14	-	15	Some strains of Enterobacteriaceae (particularly <i>Serratia</i> , <i>Providencia</i> , <i>Citrobacter</i> and <i>Enterobacter</i> spp.) produce clear zones of inhibition with small colonies around the colistin disc. These isolates are resistant as the MICs typically exceed 128 mg/L.
Co-trimoxazole	2	-	2	1.25/ 23.75	15	-	16	The MIC breakpoint is based on the trimethoprim concentration in a 1:19 combination with sulphamethoxazole. For advice on testing susceptibility to co-trimoxazole, see Appendix 1.
Sulfamethoxazole	32	-	32	100	13	-	14	
Trimethoprim	2	1-2	0.5	2.5	14	15-19	20	
Trimethoprim UTI ¹⁻⁶	4	4	2	2.5	13	14-16	17	These interpretative criteria are for <i>E. coli</i> , <i>Klebsiella</i> spp., <i>P. mirabilis</i> and coliforms only.
Doxycycline	1	-	1	30	28	-	29	
Fosfomycin UTI ¹⁻⁶	32	-	32	200/ 50	24	-	25	These interpretative criteria are for <i>E. coli</i> only. Disc content indicates fosfomycin/glucose-6-phosphate contents.
Fosfomycin UTI ¹⁻⁶	32	-	32	200/ 50	36	-	37	These interpretative criteria are for <i>P. mirabilis</i> only. The susceptibility of <i>Proteus</i> spp. that swarm up to the disc can be difficult to interpret. Disc content indicates fosfomycin/glucose-6-phosphate contents.

Table 6	MIC breakpoint (mg/L)			Disc content (μg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S \leq		R \leq	I	S \geq	
Miscellaneous antibiotics cont.								
Nitrofurantoin UTI ¹⁻⁶	64	-	64	200	16	-	17	These interpretative criteria are for <i>E. coli</i> only.
Tigecycline	2	2	1	15	19	20-23	24	<i>Morganella morganii</i> , <i>Providencia</i> spp. and <i>Proteus</i> spp. are considered inherently non-susceptible to tigecycline.

The information in bold is tentative. Breakpoints will remain tentative for one year from when published.

Table 7. MIC and zone diameter breakpoints for *Acinetobacter* species

Antibiotic	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Aminoglycosides								
Gentamicin	4	-	4	10	19	-	20	
Penicillins								
Piperacillin/tazobactam	16	-	16	75/10	21	-	22	
Carbapenems								
Doripenem	4	2-4	1	10	14	15-21	22	
Imipenem	8	4-8	2	10	13	14-24	25	
Meropenem	8	4-8	2	10	12	13-19	20	
Quinolones								
Ciprofloxacin	1	-	1	1	20	-	21	
Miscellaneous antibiotics								
Colistin	2	-	2	-	-	-	-	Disc diffusion susceptibility testing is unreliable because of the high rate of false susceptibility. An MIC method is therefore recommended.
Tigecycline	2	2	1	15	19	20-23	24	

The information in bold is tentative. Breakpoints will remain tentative for one year from when first published.

Table 8. MIC and zone diameter breakpoints for *Pseudomonas* spp.

Table 8	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Aminoglycosides								
Amikacin	16	16	8	30	15	16-18	19	
Gentamicin	4	-	4	10	17	-	18	
Netilmicin	4	-	4	10	13	-	14	
Tobramycin	4	-	4	10	19	-	20	
Penicillins								
Carbenicillin	128	-	128	100	12	-	13	
Piperacillin	16	-	16	75	23	-	24	
Piperacillin/tazobactam	16	-	16	75/10	21	-	22	
Ticarcillin	64	32-64	16	75	19	-	20	
Ticarcillin/clavulanate	64	32-64	16	75/10	19	-	20	
Cephalosporins								
Cefotaxime	1	-	1	30	26	-	27	
Cefpirome	1	-	1	20	19	20-24	25	
Ceftazidime	8	-	8	30	23	-	24	
Ceftriaxone	1	-	1	30	29	-	30	
Carbapenems								
Doripenem	4	2-4	1	10	24	25-31	32	The detection of resistance mediated by carbapenemases is difficult, particularly if resistance is not fully expressed. Consideration should be given to testing ceftazidime and carbapenem resistant isolates for the presence of carbapenemases.
Imipenem	8	8	4	10	16	17-22	23	
Meropenem	8	4-8	2	10	21	22-26	27	
Other β-Lactams								
Aztreonam	8	-	8	30	22	-	23	Relates only to isolates from patients with cystic fibrosis given high dosage therapy to treat <i>P. aeruginosa</i> infection.
Quinolones								
Ciprofloxacin	1	1	0.5	1	12	13-22	23	
Ciprofloxacin	1	1	0.5	5	19	20-29	30	
Gatifloxacin	1	-	1	2	19	-	20	
Gemifloxacin	0.25	-	0.25	5	19	-	20	

Table 8	MIC breakpoint (mg/L)			Disc content (μg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S \leq		R \leq	I	S \geq	
Quinolones cont.								
Levofloxacin	2	2	1	5	16	17-21	22	
Moxifloxacin	4	2-4	1	5	17	18-24	25	
Miscellaneous antibiotics								
Colistin	2	-	2	25	13	-	14	The MIC breakpoint has changed but a review of the data indicates that no adjustment of the zone diameter breakpoints is necessary.

The information in bold is tentative. Breakpoints will remain tentative for one year from when first published.

Table 9. MIC and zone diameter breakpoints for *Stenotrophomonas maltophilia*

Antibiotic	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Co-trimoxazole	4	-	4	1.25/23.75	19	-	20	<p>For <i>Stenotrophomonas maltophilia</i>, susceptibility testing is not recommended except for co-trimoxazole (see www.bsac.org.uk BSAC Standardized Susceptibility Testing Method, Additional Methodology, <i>Stenotrophomonas maltophilia</i>).</p> <p>The MIC breakpoint is based on the trimethoprim concentration in a 1:19 combination with sulfamethoxazole.</p> <p>The MIC breakpoint has changed but a review of the data indicates that no adjustment of the zone diameter breakpoints is necessary.</p>

The information in bold is tentative. Breakpoints will remain tentative for one year from when first published.

Table 10. MIC and zone diameter breakpoints for staphylococci

Comments 1-3 relate to urinary tract infections (UTI) only.

¹ These recommendations are for organisms associated with uncomplicated urinary tract infections only. For complicated infections and infections caused by *Staphylococcus aureus* and *Staphylococcus epidermidis*, which are associated with more serious infections, systemic recommendations should be used.

² If an organism is isolated from multiple sites, for example from blood and urine, interpretation of susceptibility should be made with regard to the systemic site (e.g., if the blood isolate is resistant and the urine isolate susceptible, both should be reported resistant irrespective of the results obtained using interpretative criteria for urine isolates).

³ Direct susceptibility tests on urine samples may be interpreted only if the inoculum gives semi-confluent growth.

Table 10	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Aminoglycosides								
Amikacin for <i>Staphylococcus aureus</i>	16	16	8	30	15	16-18	19	
Amikacin for coagulase-negative staphylococci	16	16	8	30	21	22-24	25	
Gentamicin	1	-	1	10	19	-	20	
Tobramycin for <i>Staphylococcus aureus</i>	1	-	1	10	20	-	21	
Tobramycin for coagulase-negative staphylococci	1	-	1	10	29	-	30	
Neomycin	-	-	-	10	16	-	17	For topical use only. The zone diameter breakpoint distinguishes the "wild type" susceptible population from isolates with reduced susceptibility.
β-Lactams								
								Staphylococci exhibiting resistance to methicillin/oxacillin/cefoxitin should be regarded as resistant to other penicillins, cephalosporins, carbapenems and combinations of β-lactam and β-lactamase inhibitors.
Ampicillin UTI ¹⁻³ <i>Staphylococcus saprophyticus</i>	32	-	32	25	25	-	26	

Table 10	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
β-Lactams cont.								
Cefoxitin <i>Staphylococcus aureus</i>	4	-	4	10	21	-	22	
Cefoxitin coagulase-negative staphylococci	4	-	4	10	21	-	27	
Co-amoxiclav	1	-	1	2/1	17	-	18	
Co-amoxiclav UTI ¹⁻³	32	-	32	20/10	27	-	28	
<i>Staphylococcus saprophyticus</i>								
Mecillinam UTI ¹⁻³	64	-	64	50	9	-	10	
<i>Staphylococcus saprophyticus</i>								
Methicillin	4	-	4	5	14	-	15	Recommendations for tests on Mueller-Hinton or Columbia agars with 2% NaCl: Some hyper-producers of β-lactamase give zones within the range of 7-14 mm and if possible, should be checked by a PCR method for <i>mecA</i> or a latex agglutination test for PBP2a. Increase in methicillin/oxacillin zone size in the presence of clavulanic acid is not a reliable test for hyper-producers of β-lactamase as zones of inhibition with some MRSA also increase in the presence of clavulanic acid. Rarely, hyper-producers of β-lactamase give no zone in this test and would therefore not be distinguished from MRSA.
Oxacillin	2	-	2	1	14	-	15	
								Oxacillin MIC breakpoint for coagulase-negative staphylococci is currently under review.
Penicillin	0.12	-	0.12	1 unit	24	-	25	With penicillin check for a heaped zone edge which equals resistant.
Quinolones								
Ciprofloxacin	1	-	1	1	13	-	14	MIC breakpoints relate to high-dose therapy (750 mg).
Ciprofloxacin UTI ¹⁻³	4	-	4	1	17	-	18	
<i>Staphylococcus saprophyticus</i>								

Table 10	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Quinolones cont.								
Gatifloxacin	1	-	1	2	19	-	20	
Gemifloxacin	0.25	-	0.25	1	19	-	20	
Moxifloxacin	1	1	0.5	1	15	16-19	20	
Ofloxacin	1	-	1	5	27	-	28	
Miscellaneous antibiotics								
Daptomycin	1	-	1	-	-	-	-	Strains with MIC values above the susceptible breakpoint are very rare or not yet reported. The identification and antimicrobial susceptibility tests on any such isolate must be repeated and if the result is confirmed the isolate sent to a reference laboratory. Until there is evidence regarding the clinical response for confirmed isolates with MIC above the current resistant breakpoint they should be reported resistant. Susceptibility testing by disc diffusion is not recommended. Susceptibility should be determined using a broth dilution method with Mueller- Hinton broth or by a gradient method on Mueller- Hinton agar.
Teicoplanin	8	8	4	30	14	-	15	Teicoplanin disc diffusion testing is not recommended for coagulase-negative staphylococci. An MIC method should be used to determine susceptibility. Glycopeptide intermediate <i>Staphylococcus aureus</i> (GISA) cannot be detected by this method or any other disc diffusion method. The Etest "macro-method" may be used to screen for GISA and GISA with heterogenous resistance to vancomycin (hetero-GISA) but positive results require confirmation. Population analysis is the most reliable method for confirming resistance and for distinguishing susceptible, hetero-GISA and GISA isolates. If, on clinical grounds, resistance to vancomycin is suspected, it is recommended that the organism be sent to a specialist laboratory, such as Southmead Hospital in Bristol or the Antibiotic Resistance Monitoring Reference Laboratory at Colindale.
Vancomycin	8	8	4	5	11	-	12	

Table 10	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Miscellaneous antibiotics cont.								
Azithromycin	1	-	1	15	19	-	20	
Clarithromycin	2	2	1	2	14	15-17	18	
Clindamycin	0.5	0.5	0.25	2	22	23-25	26	Organisms that appear resistant to erythromycin, but susceptible to clindamycin should be checked for the presence of inducible resistance (see www.bsac.org.uk/Susceptibility Testing/BSAC Standardized Disc Susceptibility Method/Additional Methods). Clindamycin should be used with caution (if at all) for organisms with inducible MLS _B resistance.
Erythromycin	2	2	1	5	16	17-19	20	
Quinupristin/dalfopristin	2	2	1	15	18	19-21	22	The presence of blood has a marked effect on the activity of quinupristin/dalfopristin. On the rare occasions when blood needs to be added to enhance the growth of staphylococci, susceptible = ≥15 mm, resistant ≤14 mm.
Telithromycin	0.5	-	0.5	15	26	-	27	
Chloramphenicol	8	-	8	10	14	-	15	
Co-trimoxazole	2	-	2	1.25/23.75	16	-	17	For advice on testing susceptibility to co-trimoxazole see Appendix 1. The MIC breakpoint is based on the trimethoprim concentration in a 1:19 combination with sulphamethoxazole.
Trimethoprim	1	-	1	5	19	-	20	Amended zone diameter breakpoints are microbiological breakpoints based on the MIC distribution for the wild type population. However, there is no clear evidence correlating these breakpoints with clinical efficacy.
Trimethoprim UTI ¹⁻³ <i>Staphylococcus saprophyticus</i>	4	4	2	2.5	12	13-14	15	
Doxycycline	1	-	1	30	30	-	31	
Minocycline	0.5	-	0.5	30	27	-	28	
Tetracycline	1	-	1	10	19	-	20	
Tigecycline	0.5	-	0.5	15	25	-	26	

Table 10	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Miscellaneous antibiotics cont.								
Fosfomycin UTI ¹⁻³ <i>Staphylococcus saprophyticus</i>	128	-	128	200/50	19	-	20	Disc content indicates fosfomycin/glucose-6-phosphate content.
Fusidic acid	1	-	1	10	29	-	30	
Linezolid	4	-	4	10	19	-	20	Information on clinical response in patients with serious staphylococcal infections is not yet available. In such patients an MIC determination might be appropriate.
Mupirocin	4	-	4	5	21	-	22	An Etest or other MIC method should be performed on any strain designated mupirocin resistant when tested with a 5 µg disc. The MIC will indicate whether the strain has low-level (MIC 8 – 256 mg/L) or high-level (MIC ≥512 mg/L) resistance. Isolates with low-level resistance to mupirocin (MICs 8-256 mg/L) may be eradicated more slowly than susceptible isolates.
Mupirocin	256	8-256	4	20	6	7-26	27	
Nitrofurantoin UTI ¹⁻³ <i>Staphylococcus saprophyticus</i>	64	-	64	200	19	-	20	A review of the data indicates that no adjustment of the zone diameter breakpoints is necessary with the change in MIC breakpoint.
Rifampicin	0.5	012-0.5	0.06	2	23	24-29	30	

The information in bold is tentative. Breakpoints will remain tentative for one year from when first published.

Table 11. MIC and zone diameter breakpoints for *Streptococcus pneumoniae*

Table 11	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Penicillins								<p>Reduced susceptibility to penicillin in <i>Streptococcus pneumoniae</i> is most reliably detected with an oxacillin 1 µg disc; confirm resistance with a penicillin MIC determination. Organisms with an MIC ≤ 1 mg/L are considered susceptible to β-lactam agents except in infections of the central nervous system. In addition, cefotaxime or ceftriaxone MIC determination is advised for isolates from meningitis or other invasive infections.</p> <p>For cefuroxime the zone diameter breakpoints relate to an MIC breakpoint of 0.5 mg/L as no data for the intermediate category are currently available.</p>
Penicillin	1	0.12-1	0.06	Oxacillin1	19	-	20	
Cephalosporins								
Cefaclor	1	-	1	30	24	-	25	
Cefixime	1	-	1	5	19	-	20	
Cefotaxime	2	1-2	0.5	5	20	21-24	25	
Cefpodoxime	1	-	1	1	21	-	22	
Ceftibuten	1	-	1	10	27	-	28	
Ceftizoxime	1	-	1	30	29	24-27	30	
Ceftriaxone	2	1-2	0.5	30	23	-	28	
Cefuroxime	1	1	0.5	5	24	-	25	
Cefadroxil	1	-	1	30	24	-	25	
Cefalexin	2	-	2	30	24	-	25	
Carbapenems								
Ertapenem	0.5	-	0.5	10	32	-	33	
Imipenem	4	-	4	10	24	-	25	
Meropenem	4	-	4	10	27	-	28	
Quinolones								
Ciprofloxacin	2	0.25-2	0.12	1	9	10-24	25	"Wild type" isolates (ciprofloxacin MICs 0.25-2 mg/L; ofloxacin MICs 0.25-4 mg/L) are considered intermediate in susceptibility.
Ofloxacin	4	0.25-4	0.12	5	15	16-27	28	
Gatifloxacin	1	-	1	2	19	-	20	
Gemifloxacin	0.25	-	0.25	1	19	-	20	
Levofloxacin	2	-	2	1	9	-	10	
Moxifloxacin	0.5	-	0.5	1	17	-	18	
Miscellaneous antibiotics								
Azithromycin	0.5	0.5	0.25	15	19	20-21	22	
Chloramphenicol	8	-	8	10	17	-	18	
Clarithromycin	0.5	0.5	0.25	2	19	20-21	22	

Table 11	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Miscellaneous antibiotics cont.								
Co-trimoxazole	2	2	1	1.25/23.75	16	-	17	For advice on testing susceptibility to co-trimoxazole see Appendix 1. The MIC breakpoint is based on the trimethoprim concentration in a 1:19 combination with sulphamethoxazole.
Erythromycin	0.5	0.5	0.25	5	19	20-21	22	
Linezolid	4	4	4	10	1	-	20	The MIC breakpoint has changed but a review of the data indicates that no adjustment of the zone diameter breakpoints is necessary.
Quinupristin/dalfopristin	2	-	2	15	19	-	20	
Rifampicin	0.5	0.12-0.5	0.06	5	20	21-22	23	
Telithromycin	0.5	0.5	0.25	15	28	-	29	Insufficient data are available to distinguish the intermediate category.
Tetracycline	1	-	1	10	19	-	20	
Vancomycin	4	-	4	5	12	-	13	

The information in bold is tentative. Breakpoints will remain tentative for one year from when first published.

Table 12. MIC and zone diameter breakpoints for enterococci

Comments 1-3 relate to urinary tract infections (UTIs) only.

¹ UTI recommendations are for organisms associated with uncomplicated urinary tract infections only. For complicated urinary tract infections, systemic recommendations should be used.

² If an organism is isolated from multiple sites, for example from blood and urine, interpretation of susceptibility should be made with regard to the systemic site (e.g., if the blood isolate is resistant and the urine isolate susceptible, both should be reported resistant irrespective of the results obtained using interpretative criteria for urine isolates).

³ Direct susceptibility tests on urine samples may be interpreted only if the inoculum gives semi-confluent growth.

NB. For isolates from endocarditis the MIC should be determined and interpreted according to national endocarditis guidelines (Elliott TS et al. Guidelines for the antibiotic treatment of endocarditis in adults: report of the Working Party of the British Society for Antimicrobial Chemotherapy. J Antimicrob Chemother. 2004; **54**: 971-81).

Table 12	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Aminoglycosides								
Gentamicin	128	-	128	200	14	-	15	High-level gentamicin-resistant enterococci usually give no zone or only a trace of inhibition around gentamicin 200 µg discs. Occasionally, however, the plasmid carrying the resistance gene may be unstable and the resistance is seen as a zone of inhibition with a few small colonies within the zone. Retesting of resistant colonies results in growth to the disc or increased numbers of colonies within the zone. Zones should be carefully examined to avoid missing such resistant organisms. If in doubt, isolates may be sent to a reference laboratory for confirmation.
Streptomycin	128	-	128	300	23	-	24	
Penicillins								
Ampicillin	8	-	8	10	19	-	20	
Ampicillin UTI ¹⁻³	32	-	32	25	19	-	20	
Co-amoxiclav UTI ¹⁻³	32	-	32	20/10	20	-	21	
Carbapenems								
Imipenem	8	8	4	10	16	17-18	19	Recommendations for <i>E. faecalis</i> only.
Meropenem	4	-	4	10	19	-	20	

Table 12	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Quinolones								
Ciprofloxacin UTI ¹⁻³	4	-	4	1	11	-	12	
Ciprofloxacin UTI ¹⁻³	4	-	4	5	15	-	16	
Nalidixic acid UTI ¹⁻³	16	-	16	30	17	-	18	
Norfloxacin UTI ¹⁻³	4	-	4	2	15	-	16	
Miscellaneous antibiotics								
Azithromycin	1	-	1	15	29	-	30	
Quinupristin/dalfopristin	4	2-4	1	15	11	12-19	20	Generally, <i>E. faecalis</i> are I or R and <i>E. faecium</i> are susceptible. The presence of blood has a marked effect on the activity of quinupristin/dalfopristin. On the rare occasions when blood needs to be added to enhance the growth of enterococci, breakpoints are ≥15 mm, ≤14 mm.
Fosfomycin UTI ¹⁻³	128	-	128	200/50	19	-	20	Disc content indicates fosfomycin/glucose-6-phosphate contents.
Linezolid	4	-	4	10	19	-	20	
Nitrofurantoin UTI ¹⁻³	64	-	64	200	19	-	20	
Teicoplanin	8	8	4	30	19	-	20	To ensure that microcolonies indicating reduced susceptibility to the glycopeptides are detected, it is essential that plates are incubated for at least 24 h before reporting a strain as susceptible to vancomycin or teicoplanin. Zone diameter breakpoints relate to the MIC breakpoint of 4 mg/L as no data for the intermediate category are currently available.
Vancomycin	8	8	4	5	12	-	13	
Tetracycline	1	-	1	10	25	-	26	
Tigecycline	0.5	0.5	0.25	15	20	-	21	There is no intermediate category for disc diffusion, as non-susceptible isolates are rare and were not available for testing.

Table 12	MIC breakpoint (mg/L)			Disc content (μ g)	Interpretation of zone diameters (mm)			Comment
	R >	I	S \leq		R \leq	I	S \geq	
Trimethoprim UTI ¹⁻³	1	0.06-1	0.03	2.5	25	26-32	33	There is conflicting evidence as to whether wild type Enterococci causing lower UTI respond to standard doses of trimethoprim. The present breakpoints allow laboratories to report these wild type isolates as susceptible or indeterminate depending on preference. It is more clear that Enterococci which test as resistance should not be treated with trimethoprim.

The information in bold is tentative. Breakpoints will remain tentative for one year from when first published.

Table 13. MIC and zone diameter breakpoints for α -haemolytic streptococci

N.B. For isolates from endocarditis the MIC should be determined and interpreted according to national endocarditis guidelines (Elliott TS et al. Guidelines for the antibiotic treatment of endocarditis in adults: report of the Working Party of the British Society for Antimicrobial Chemotherapy. J Antimicrob Chemother. 2004; **54**: 971-81).

Antibiotic	MIC breakpoint (mg/L)			Disc content (μ g)	Interpretation of zone diameters (mm)			Comment
	R >	I	S \leq		R \leq	I	S \geq	
Penicillins								
Amoxicillin	1	-	1	2	19	-	20	
Penicillin	0.12	-	0.12	1 unit	21	-	22	
Cephalosporins								
Cefotaxime	1	-	1	5	20	-	21	
Miscellaneous antibiotics								
Clindamycin	0.5	-	0.5	2	19	-	20	Organisms that appear resistant to erythromycin, but susceptible to clindamycin should be checked for the presence of inducible MLS _B resistance (see www.bsac.org.uk/Susceptibility Testing/BSAC Standardized Disc Susceptibility Method/Additional Methods). Clindamycin should be used with caution (if at all) for organisms with inducible MLS _B resistance.
Erythromycin	0.5	-	0.5	5	19	-	20	
Linezolid	2	-	2	10	19	-	20	
Teicoplanin	4	-	4	30	15	-	16	
Tetracycline	1	-	1	10	23	-	24	
Tigecycline	0.5	0.5	0.25	15	19	20-24	25	
Vancomycin	4	-	4	5	13	-	14	

Table 14. MIC and zone diameter breakpoints for β -haemolytic streptococci

Comments 1-3 relate to urinary tract infections (UTIs) only.

¹ UTI recommendations are for organisms associated with uncomplicated urinary tract infections only. For complicated urinary tract infections and infections systemic recommendations should be used.

² If an organism is isolated from multiple sites, for example from blood and urine, interpretation of susceptibility should be made with regard to the systemic site (e.g., if the blood isolate is resistant and the urine isolate susceptible, both should be reported resistant irrespective of the results obtained using interpretative criteria for urine isolates).

³ Direct susceptibility tests on urine samples may be interpreted only if the inoculum gives semi-confluent growth.

Table 14 Antibiotic	MIC breakpoint (mg/L)			Disc content (μ g)	Interpretation of zone diameters (mm)			Comment
	R >	I	S \leq		R \leq	I	S \geq	
Penicillins								
Amoxicillin	1	-	1	2	19	-	20	
Ampicillin UTI ¹⁻³ Group B Streptococci	32	-	32	25	25	-	26	
Co-amoxiclav UTI ¹⁻³ Group B Streptococci	32	-	32	20/10	27	-	28	
Penicillin	0.12	-	0.12	1 unit	19	-	20	
Cephalosporins								
Cefadroxil	1	-	1	30	24	-	25	
Cefalexin UTI ¹⁻³ Group B Streptococci	32	-	32	30	23	-	24	
Cefalexin	2	-	2	30	24	-	25	
Cefixime	1	-	1	5	19	-	20	
Cefotaxime	0.5	-	0.5	5	23	-	24	
Cefalothin	1	-	1	30	28	-	29	
Carbapenems								
Ertapenem	0.5	-	0.5	10	34	-	35	
Quinolones								
Ciprofloxacin UTI ¹⁻³ Group B Streptococci	4	-	4	1	12	-	13	
Ciprofloxacin UTI ¹⁻³ Group B Streptococci	4	-	4	5	18	-	19	
Miscellaneous antibiotics								
Azithromycin	0.5	0.5	0.25	15	19	20-21	22	
Clarithromycin	0.5	0.5	0.25	2	19	20-21	22	

Table 14	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Miscellaneous antibiotics cont.								
Clindamycin	0.5	-	0.5	2	16	-	17	Organisms that appear resistant to erythromycin, but susceptible to clindamycin should be checked for the presence of inducible MLS _B resistance (see www.bsac.org.uk/Susceptibility Testing/BSAC Standardized Disc Susceptibility Method/Additional Methods). Clindamycin should be used with caution (if at all) for organisms with inducible MLS _B resistance.
Erythromycin	0.5	0.5	0.25	5	19	20-21	22	
Co-trimoxazole	2	-	2	1.25/23.75	16	-	17	For advice on testing susceptibility to co-trimoxazole see Appendix 1. The MIC breakpoint is based on the trimethoprim concentration in a 1:19 combination with sulphamethoxazole.
Trimethoprim UTI ¹⁻³ Group B Streptococci	2	-	2	2.5	15	-	16	
Daptomycin	1	-	1	-	-	-	-	Strains with MIC values above the susceptible breakpoint are very rare or not yet reported. The identification and antimicrobial susceptibility tests on any such isolate must be repeated and if the result is confirmed the isolate sent to a reference laboratory. Until there is evidence regarding the clinical response for confirmed isolates with MIC above the current resistant breakpoint they should be reported resistant. Disc diffusion susceptibility testing is not recommended.
Linezolid	4	4	2	10	19	-	20	Zone diameter breakpoints relate to the MIC breakpoint of 2 mg/L as no data for the intermediate category are currently available.
Nitrofurantoin UTI ¹⁻³ Group B Streptococci	32	-	32	200	19	-	20	

Table 14	MIC breakpoint (mg/L)			Disc content (μg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S \leq		R \leq	I	S \geq	

Miscellaneous antibiotics cont.

Telithromycin	0.5	0.5	0.25	15	25	-	26	Zone diameter breakpoints relate to the "wild type" susceptible population as no data are available for the non-susceptible population.
Tetracycline	1	-	1	10	19	-	20	
Tigecycline	0.5	0.5	0.25	15	19	20-24	25	

The information in bold is tentative. Breakpoints will remain tentative for one year from when first published.

Table 15. MIC and zone diameter breakpoints for *Moraxella catarrhalis*

Table 15 Antibiotic	MIC breakpoint (mg/L)			Disc content (μg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S \leq		R \leq	I	S \geq	
Penicillins								
Ampicillin	1	-	1	2	29	-	30	Test for β -lactamase. β -Lactamase positive isolates of <i>Moraxella catarrhalis</i> are often slow to become positive in tests for β -lactamase production so tests must be examined after the longest recommended time before being interpreted as negative.
Co-amoxiclav	1	-	1	2/1	18	-	19	
Cephalosporins								
Cefaclor	1	-	1	30	22	-	23	
Cefuroxime	2	2	1	5	19	-	20	Zone diameter breakpoints relate to the MIC breakpoint of 1 mg/L as no data for the intermediate category are currently available.
Carbapenems								
Ertapenem	0.5	-	0.5	10	34	-	35	
Quinolones								
Ciprofloxacin	0.5	-	0.5	1	17	-	18	Quinolone resistance is most reliably detected with nalidixic acid discs. Isolates with reduced susceptibility to fluoroquinolones show no zone of inhibition with a 30 μg nalidixic acid disc.
Gatifloxacin	1	-	1	2	19	-	20	
Gemifloxacin	0.25	-	0.25	1	19	-	20	
Levofloxacin	1	-	1	1	19	-	20	
Moxifloxacin	0.5	-	0.5	1	17	-	18	
Nalidixic acid	-	-	-	30	-	-	-	Quinolone resistance is most reliably detected with nalidixic acid. Strains with reduced susceptibility to fluoroquinolones give no zone of inhibition with a 30 μg nalidixic acid disc.
Ofloxacin	0.5	-	0.5	5	34	-	35	
Miscellaneous antibiotics								
Chloramphenicol	2	-	2	10	22	-	23	
Clarithromycin	0.5	0.5	0.25	2	19	20-21	22	

Table 15 Antibiotic	MIC breakpoint (mg/L)			Disc content (μ g)	Interpretation of zone diameters (mm)			Comment
	R >	I	S \leq		R \leq	I	S \geq	
Miscellaneous antibiotics								
Erythromycin	0.5	0.5	0.25	5	27	-	28	Zone diameter breakpoints relate to the MIC breakpoint of 0.25 mg/L as no data for the intermediate category are currently available.
Telithromycin	0.5	-	0.5	15	29	-	30	
Co-trimoxazole	2	-	2	1.25/23.75	11	-	12	For advice on testing susceptibility to co-trimoxazole, see Appendix 1. The MIC breakpoint is based on the trimethoprim concentration in a 1:19 combination with sulphamethoxazole.
Linezolid	4	-	4	10	19	-	20	
Tetracycline	1	-	1	10	21	-	22	

The information in bold is tentative. Breakpoints will remain tentative for one year from when first published.

Table 16. MIC and zone diameter breakpoints for *Neisseria gonorrhoeae*

Table 16	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Penicillins								
Penicillin	1	0.12	0.06	1 unit	17	18-25	26	Test for β-lactamase.
Cephalosporins								
Cefixime	1	-	1	5	29	-	30	Resistance to ceftriaxone, cefotaxime and cefixime has not been described. Isolates with chromosomally encoded reduced susceptibility to penicillin have slightly reduced zones of inhibition with these agents but they remain susceptible. Results for isolates with reduced zones around ceftriaxone, cefotaxime and cefixime discs should be confirmed by MIC determinations. Although cefuroxime is not recommended for clinical use, it can be used as an indicator antibiotic to detect reduced susceptibility to other oxyimino cephalosporins.
Cefotaxime	0.12	-	0.12	5	29	-	30	
Ceftriaxone	0.12	-	0.12	5	34	-	35	
Cefuroxime	1	-	1	5	19	-	20	
Quinolones								
Ciprofloxacin	0.06	0.06	0.03	1	28	-	29	For ciprofloxacin the zone diameter breakpoints relate to the MIC breakpoint of 0.03mg/L as no data for the intermediate category are currently available. Quinolone resistance is generally reliably detected with nalidixic acid; however there are a few isolates that are resistant to ciprofloxacin yet susceptible to nalidixic acid in disc diffusion tests. The mechanism of resistance and the prevalence of these isolates in the UK is still under investigation. Isolates with reduced susceptibility to fluoroquinolones normally have no zone of inhibition with a 30 µg nalidixic acid disc. For organisms with nalidixic acid zone diameters 10-31 mm a ciprofloxacin MIC should be determined if the patient is to be treated with this agent.
Nalidixic acid	-	-	-	30	9	10-31	32	

Table 16	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Miscellaneous antibiotics								
Azithromycin	0.5	0.5	0.25	15	27	-	28	Zone diameter breakpoints relate to the MIC breakpoint of > 0.5 mg/L as disc diffusion testing will not reliably differentiate between the intermediate and susceptible populations.
Erythromycin	0.5		0.5	5	11	-	12	Erythromycin is no longer used for therapy, but may be tested for epidemiological purposes.
Rifampicin	1	-	1	2	20	-	21	
Spectinomycin	64	-	64	25	13	-	14	
Tetracycline	1	-	1	10	26	-	27	The tetracycline result may be used to infer susceptibility to doxycycline.

The information in bold is tentative. Breakpoints will remain tentative for one year from when first published.

Table 17. MIC and zone diameter breakpoints for *Neisseria meningitidis*

Antibiotic	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ?		R ?	I	S ?	
Penicillins								
Ampicillin	-	-	-	2	31	-	32	Ampicillin and amoxicillin are used as indicator antibiotics to detect reduced susceptibility to penicillin. The recommendations given are for this purpose only; ampicillin and amoxicillin should not be used therapeutically.
Amoxicillin	-	-	-	2	29	-	30	
Penicillin	0.06	-	0.06	1 unit	30	-	31	
Cephalosporins								
Cefotaxime	0.12	-	0.12	5	39	-	40	
Ceftriaxone	0.12	-	0.12	5	39	-	40	
Quinolones								
Ciprofloxacin	0.06	0.06	0.03	1	31	-	32	Quinolone resistance is most reliably detected in tests with nalidixic acid. Isolates with reduced susceptibility to fluoroquinolones have no zone of inhibition with 30 µg nalidixic acid discs. Zone diameter breakpoints relate to the MIC breakpoint of 0.03 mg/L as no data for the intermediate category are currently available.
Miscellaneous antibiotics								
Chloramphenicol	4	4	2	10	19	-	20	Zone diameter breakpoints relate to the MIC breakpoint of 2 mg/L as insufficient data to distinguish the intermediate category are currently available.
Erythromycin	0.5	-	0.5	5	26	-	27	
Rifampicin	0.25	-	0.25	2	29	-	30	Epidemiological breakpoint based on an MIC breakpoint of 0.25 mg/L.
Tetracycline	1	-	1	10	21	-	22	

Table 18. MIC and zone diameter breakpoints for *Haemophilus influenzae*

Table 18	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Penicillins								
Amoxicillin	1	-	1	2	16	-	17	Test for β-lactamase in isolates that appear resistant or have borderline susceptibility by disc testing.
Ampicillin	1	-	1	2	17	-	18	
Co-amoxiclav	1	-	1	2/1	16	-	17	
Cephalosporins								
Cefaclor	1	-	1	30	36	-	37	See Appendix 2.
Cefotaxime	0.12	-	0.12	5	24	-	25	
Ceftazidime	2	-	2	30	29	-	30	
Ceftriaxone	0.12	-	0.12	30	24	-	25	
Cefuroxime (parenteral)	2	-	1	5	16	-	17	Zone diameter breakpoints relate to the MIC breakpoint of 1 mg//L as no data for the intermediate category are currently available.
Carbapenems								
Ertapenem	0.5	-	0.5	10	32	-	33	
Imipenem	2	-	2	10	22	-	23	
Meropenem	2	-	2	10	22	-	23	
Quinolones								
Ciprofloxacin	0.5	-	0.5	1	27	-	28	Quinolone resistance is most reliably detected in tests with nalidixic acid. Strains with reduced susceptibility to fluoroquinolones give no zone of inhibition with a 30µg nalidixic acid disc.
Gatifloxacin	1	-	1	2	19	-	20	
Gemifloxacin	0.25	-	0.25	1	19	-	20	
Levofloxacin	1	-	1	1	19	-	20	
Moxifloxacin	0.5	-	0.5	1	17	-	18	
Nalidixic acid	-	-	-	30	-	-	-	
Ofloxacin	0.5	-	0.5	5	26	-	37	
Miscellaneous antibiotics								
Azithromycin	4	0.5-4	0.25	15	19	20-34	35	No resistant strains yet described.
Chloramphenicol	2	-	2	10	24	-	25	

Table 18	MIC breakpoint (mg/L)			Disc content (μg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S \leq		R \leq	I	S \geq	
Miscellaneous antibiotics cont.								
Clarithromycin	16	1-16	0.5	5	9	10-24	25	
Erythromycin	8	1-8	0.5	5	14	15-27	28	
Telithromycin	2	1-2	0.5	15	15	16-19	20	The mode telithromycin MIC for these organisms is 1 mg/L; therefore the majority of isolates will be interpreted as having intermediate susceptibility.
Co-trimoxazole	2	-	2	25	21	-	22	For advice on testing susceptibility to co-trimoxazole see Appendix 1. The MIC breakpoint is based on the trimethoprim concentration in a 1:19 combination with sulphamethoxazole.
Trimethoprim	0.5	-	0.5	2.5	20	-	21	
Tetracycline	1	-	1	10	21	-	22	

Table 19. MIC and zone diameter breakpoints for *Pasteurella multocida*

Antibiotic	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Penicillins								
Ampicillin	1	-	1	10	29	-	30	
Penicillin	0.12	-	0.12	1 unit	21	-	22	
Cephalosporins								
Cefotaxime	1	-	1	5	33	-	34	
Quinolones								
Ciprofloxacin	1	-	1	1	28	-	29	Quinolone resistance is most reliably detected in tests with nalidixic acid discs.
Nalidixic acid	-	-	-	30	27	-	28	
Miscellaneous antibiotics								
Tetracycline	1	-	1	10	25	-	26	

Table 20. MIC and zone diameter breakpoints for *Campylobacter* spp.

Antibiotic	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Quinolones								
Ciprofloxacin	1	1	0.5	1	17	-	18	Quinolone resistance is most reliably detected in tests with nalidixic acid discs. Strains with reduced susceptibility to fluoroquinolones give no zone of inhibition with a 30µg nalidixic acid disc. The zone diameters relate to an MIC breakpoint of 0.5 mg/L as no data for the intermediate category are currently available.
Nalidixic acid	-	-	-	30	-	-	-	
Miscellaneous antibiotics								
Erythromycin	0.5	-	0.5	5	19	-	20	

Table 21. MIC and zone diameter breakpoints for Coryneform organisms

Antibiotic	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Penicillins								
Penicillin	0.12	-	0.12	1 unit	19	-	20	
Quinolones								
Ciprofloxacin	1	1	0.5	1	11	12-16	17	The zone diameters relate to an MIC breakpoint of 0.5 mg/L as no data for the intermediate category are currently available.
Miscellaneous antibiotics								
Vancomycin	8	8	4	5	19	-	20	The zone diameters relate to an MIC breakpoint of 4 mg/L as no data for the intermediate category are currently available.

Table 22. MIC and zone diameter breakpoints for *Bacteroides fragilis*

<i>B. fragilis</i> is inherently resistant to penicillin.								
Antibiotic	MIC breakpoint (mg/L)			Disc content (μ g)	Interpretation of zone diameters (mm)			Comment
	R >	I	S \leq		R \leq	I	S \geq	
Penicillins								
Co-amoxiclav	8	-	8	30	27	-	28	
Piperacillin/tazobactam	16	-	16	75/10	26	-	27	The breakpoints are based on the "wild type" susceptible population as there are few clinical data relating MIC to outcome. Organisms that appear resistant in disc diffusion tests should have resistance confirmed by MIC determination and resistant isolates should be sent to the Anaerobe Reference Laboratory in Cardiff.
Carbapenems								
Meropenem	8	4-8	2	10	18	19-25	26	
Miscellaneous antibiotics								
Clindamycin	4	-	4	2	9	-	10	The breakpoints are based on the "wild type" susceptible population as there are few clinical data relating MIC to outcome. Organisms that appear resistant in disc diffusion tests should have resistance confirmed by MIC determination and resistant isolates should be sent to the Anaerobe Reference Laboratory in Cardiff.
Metronidazole	4	-	4	5	17	-	18	There is no evidence to change the epidemiological zone diameter breakpoint with the change in MIC breakpoint.

The information in bold is tentative. Breakpoints will remain tentative for one year from when first published.

Table 23. MIC and zone diameter breakpoints for *Bacteroides thetaiotaomicron*

Antibiotic	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Carbapenems								
Meropenem	8	4-8	2	10	18	19-25	26	
Miscellaneous antibiotics								
Clindamycin	4	-	4	2	9	-	10	The breakpoints are based on the "wild type" susceptible population as there are few clinical data relating MIC to outcome. Organisms that appear resistant in disc diffusion tests should have resistance confirmed by MIC determination and resistant isolates should be sent to the Anaerobe Reference Laboratory in Cardiff.
Metronidazole	4	-	4	5	17	-	18	

The information in bold is tentative. Breakpoints will remain tentative for one year from when first published.

NB. *B. thetaiotaomicron* is inherently resistant to penicillin.

There is a poor relationship between MIC and zone of inhibition for inhibitor combinations and therefore recommendations are not given for co-amoxiclav or piperacillin/tazobactam. If the recommendations for *B. fragilis* are used for *B. thetaiotaomicron* interpretation may be erroneous.

Table 24. MIC and zone diameter breakpoints for *Clostridium perfringens*

Antibiotic	MIC breakpoint (mg/L)			Disc content (µg)	Interpretation of zone diameters (mm)			Comment
	R >	I	S ≤		R ≤	I	S ≥	
Penicillins								
Co-amoxiclav	8	-	8	30	27	-	28	
Penicillin	0.12	-	0.12	1 unit	19	-	20	
Piperacillin/tazobactam	16	-	16	75/10	26	-	27	The breakpoints are based on the "wild type" susceptible population as there are few clinical data relating MIC to outcome. Organisms that appear resistant in disc diffusion test should have resistance confirmed by MIC determination and resistant isolates should be sent to the Anaerobe Reference Laboratory in Cardiff.
Carbapenems								
Meropenem	8	4-8	2	10	18	19-25	26	
Miscellaneous antibiotics								
Clindamycin	4	-	4	2	9	-	10	The breakpoints are based on the "wild type" susceptible population as there are few clinical data relating MIC to outcome. Organisms that appear resistant in disc diffusion test should have resistance confirmed by MIC determination and resistant isolates should be sent to the Anaerobe Reference Laboratory in Cardiff.
Metronidazole	4	-	4	5	17	-	18	There is no evidence to change the epidemiological zone diameter breakpoint with the change in MIC breakpoint.

The information in bold is tentative. Breakpoints will remain tentative for one year from when first published.

Appendix 1: Testing antimicrobial susceptibility to co-trimoxazole

Breakpoints for testing susceptibility to co-trimoxazole are provided. However, the following recommendations from the UK Committee on the Safety of Medicines (CSM) should be noted.

”Co-trimoxazole should be limited to the role of drug of choice in *Pneumocystis carinii* pneumonia; it is also indicated for toxoplasmosis and nocardiosis. It should now only be considered for use in acute exacerbations of chronic bronchitis and infections of the urinary tract when there is good bacteriological evidence of sensitivity to co-trimoxazole and good reason to prefer this combination to a single antibiotic; similarly it should only be used in acute otitis media in children when there is good reason to prefer it. Review of the safety of co-trimoxazole using spontaneous adverse drug reaction data has indicated that the profile of reported adverse reactions with trimethoprim is similar to that with co-trimoxazole; blood and generalised skin disorders are the most serious reactions with both drugs and predominantly have been reported to occur in elderly patients. A recent large post-marketing study has demonstrated that such reactions are very rare with co-trimoxazole; the study did not distinguish between co-trimoxazole and trimethoprim with respect to serious hepatic, renal, blood or skin disorders.”

Appendix 2: Efficacy of cefaclor in the treatment of respiratory infections caused by *Haemophilus influenzae*

Concerns have been expressed, particularly by laboratories moving from Stokes' method to the BSAC disc diffusion method, about the interpretation of susceptibility of *Haemophilus influenzae* to cefaclor. When using Stokes' method the majority of isolates appeared susceptible; but with the BSAC disc diffusion method most isolates are now reported resistant. The following comments explain the BSAC rationale for interpretation of cefaclor susceptibility.

Cefaclor pharmacokinetics

Cefaclor is dosed at 250-500 mg TDS po: 250 mg TDS is probably the most common dose but data is absent to confirm this. The expected C_{max} for 250 mg is 5-10 mg/L and 10-20 mg/l for 500 mg; the half life is 1 h; drug concentration in blood is <1 mg/L at 4 h and the protein binding is 25-50%. Tissue penetration is similar to other β -lactams.

*Cefaclor potency against *Haemophilus influenzae**

Data from the BSAC surveillance programme 2003-2004 (n= 899) indicates that the cefaclor MIC range is 0.12-128 mg/L; MIC₅₀ 2 mg/L; MIC₉₀ 8 mg/L.

Pharmacodynamics

An average patient with an *Haemophilus influenzae* infection will have a free drug Time>MIC of 25% with 250 mg dosing and 37% with 500 mg dosing. A conservative Time>MIC target for cephalosporins in community practice is 40-50%, but this is not achieved with cefaclor. Therefore, it is likely that cefaclor will have at best borderline activity against *Haemophilus influenzae*.

Conclusion

The pharmacodynamic data indicate that cefaclor has borderline activity against *Haemophilus influenzae*, even for community use. The outcome of infection will be difficult to predict and susceptibility testing is likely to be of limited value.

Acknowledgment

The BSAC acknowledges the assistance of the Swedish Reference Group for Antibiotics (SRGA) in supplying some breakpoint data for inclusion in this document.

References

1. Moosdeen, F., Williams, J.D. & Secker, A. (1988). Standardization of inoculum size for disc susceptibility testing: a preliminary report of a spectrophotometric method. *J. Antimicrob Chemother* **21**, 439-43.

Additional information

1. Susceptibility testing of *Helicobacter pylori*

Disc diffusion methods are not suitable for testing *Helicobacter pylori* as this species is slow growing and results may not be accurate. The recommended method of susceptibility testing is Etest (follow technical guide instructions).

Suspend colonies from a 2-3 day culture on a blood agar plate in sterile distilled water and adjust the density to equal a McFarland 3 standard.

Use a swab dipped in the suspension to inoculate evenly the entire surface of the plate. The medium of choice is Mueller-Hinton agar or Wilkins-Chalgren agar with 5-10% horse blood.

Allow the plate to dry and apply Etest strip.

Incubate at 35°C in microaerophilic conditions for 3-5 days. Read the MIC at the point of complete inhibition of all growth, including hazes and isolated colonies. Tentative interpretative criteria for MICs are given in Table 1.

Table 1: Tentative MIC breakpoints for *Helicobacter pylori*

Antimicrobial agent	MIC breakpoint (mg/L)	
	R >	S ≤
Amoxicillin	1	1
Clarithromycin	1	1
Tetracycline	2	2
Metronidazole	4	4

2. Susceptibility testing of *Brucella* species

Brucella spp. are Hazard Group 3 pathogens and all work must be done in containment level 3 accommodation. The antimicrobial agents most commonly used for treatment are doxycycline, rifampicin, ciprofloxacin, tetracycline and streptomycin and, from the limited information available, there is little or no resistance to these drugs. *Brucella* spp. are uncommon isolates and interpretative standards are not available. Since *Brucella* spp. are highly infectious, susceptibility testing in routine laboratories is not recommended.

3. Susceptibility testing of *Legionella* species

Legionella spp. are slow growing and have particular growth requirements. Disc diffusion methods for susceptibility testing are unsuitable. Susceptibility should be determined by agar dilution MICs on buffered yeast extract agar with 5% water-lysed horse blood. The antimicrobial agents commonly used for treatment are macrolides, rifampicin and fluoroquinolones. Validated MIC breakpoints are not established for *Legionella* spp. If results for test isolates are within range of the normal wild type distribution, given in table 2, clinical susceptibility may be assumed.

Table 2: MIC ranges for "wild type" *Legionella* spp.

Antimicrobial agent	MIC range for "wild-type" <i>Legionella</i> spp. (mg/L)
Erythromycin	0.06-0.5
Clarithromycin	0.004-0.06
Rifampicin	0.004-0.06
Ciprofloxacin	0.016-0.06

4. Susceptibility testing *Listeria* spp.

For susceptibility testing *Listeria* spp. an MIC determination is advised on Iso-Sensitest agar with incubation at 35-37°C in air. If a gradient method is used the test should be undertaken following the manufacturer's instructions. In Table 3 the MIC ranges and cut offs for "wild type" strains are shown and these can be used as an aid to interpreting susceptibility.

Table 3: MIC ranges for "wild type" *Listeria* spp.

Antimicrobial agent	MIC range (mg/L)	MIC cut off (mg/L)	Comment
Ampicillin	0.12-4	≤4	} No resistance described
Penicillin	0.015-2	≤2	
Daptomycin	1-4	≤4	
Erythromycin	0.12-1	≤1	Resistance very rare ≤ 0.5%
Gentamicin	0.06-1	≤1	Resistance rare 0%
Linezolid	1-4	≤4	
Tetracycline	0.06-1	≤1	
Trimethoprim	0.06-1	≤1	
Vancomycin	0.5-4	≤4	

5. Susceptibility testing of topical antibiotics

MIC breakpoints specifically for topical antibiotics are not given because there are no pharmacological, pharmacodynamic or clinical response data on which to base recommendations. [Relevant data would be gratefully received].

6. Development of MIC and zone diameter breakpoints

All breakpoints are subject to review in the light of additional data and any data relating to breakpoints, control zone ranges or any other aspect of antimicrobial susceptibility testing would be welcome (contact the Working Party secretary or any member listed at the front of this document).

The BSAC is part of the European Committee on Antimicrobial Susceptibility Testing (EUCAST) and is actively involved in the process of harmonization of MIC breakpoints in Europe. This process will undoubtedly lead to some small breakpoint adjustments, and these will be incorporated into the BSAC method as European breakpoints are agreed.

The BSAC has a mechanism to modify and publish changes to breakpoints on an annual basis via the BSAC www site (www.bsac.org.uk). Any changes will be dated.

Ad hoc modifications to breakpoints by users are not acceptable.

Control of Antimicrobial Susceptibility Testing

1. Control strains

Control strains include susceptible strains to monitor test performance (not for the interpretation of susceptibility), and resistant strains to confirm that the method will detect particular mechanisms of resistance, for example, *Haemophilus influenzae* ATCC 49247 is a β -lactamase negative, ampicillin resistant strain (see table 2 of Disc Diffusion Method). Tables 2-6 provide zone diameters for recommended control organisms under a range of test conditions.

Control strains can be purchased from the National Collection of Type Cultures (NCTC; HPA Centre for Infections, 61 Colindale Avenue, London NW9 5HT). Alternatively, some may be obtained commercially (see section on suppliers)

2. Maintenance of control strains

Store control strains by a method that minimises the risk of mutations, for example, at -70°C , on beads in glycerol broth. Ideally, two vials of each control strain should be stored, one as an "in-use" supply, the other for archiving. Every week a bead from the "in-use" vial should be subcultured on to appropriate non-selective media and checked for purity. From this pure culture, prepare one subculture for each of the following 7 days. Alternatively, for fastidious organisms that will not survive on plates for 7 days, subculture the strain daily for no more than 6 days.

3. Calculation of control ranges for disc diffusion tests

The acceptable ranges for the control strains have been calculated by combining zone diameter data from 'field studies' and from multiple centres supplying their daily control data, from which cumulative distributions of zones of inhibition have been prepared. From these distributions, the 2.5 and 97.5 percentiles were read to provide a range that would contain 95% of observations. If distributions are normal, these ranges correspond to the mean \pm 1.96 SD. The percentile ranges obtained by this method are, however, still valid even if the data do not show a normal distribution.

4. Frequency of routine testing with control strains

When the method is first introduced, daily testing is required until there are acceptable readings from 20 consecutive days (this also applies when new agents are introduced or when any test component changes). This provides sufficient data to support once weekly testing.

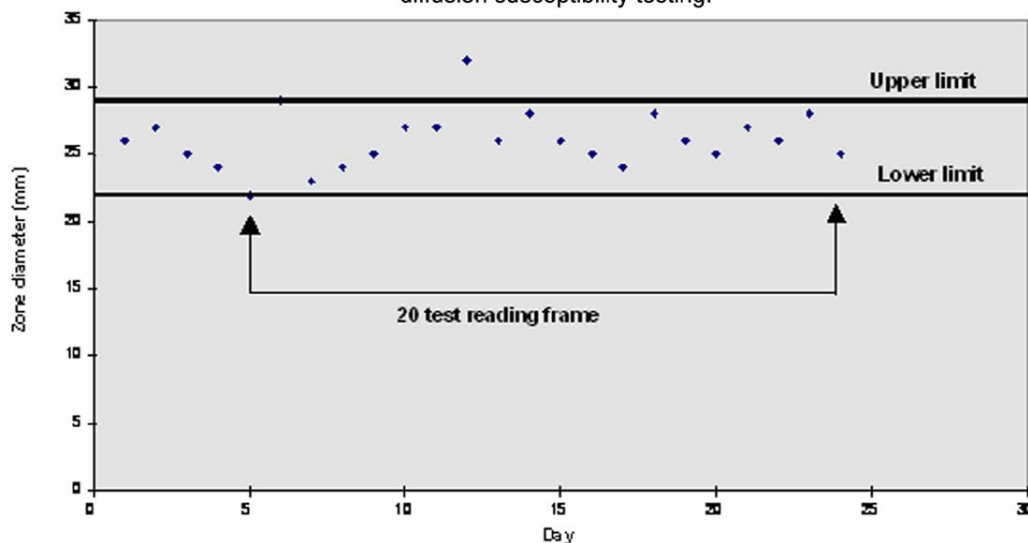
5. Use of control data to monitor the performance of disc diffusion tests

Use a reading frame of 20 consecutive results (remove the oldest result when adding a new one to make a total of 20) as illustrated in Figure 1. Testing is acceptable if no more than 1 in every 20 results is outside the limits of acceptability. If 2 or more results fall out of the acceptable range this requires immediate investigation.

Look for trends within the limits of acceptability e.g. tendency for zones to be at the limits of acceptability; tendency for zones to be consistently above or below the mean;

gradual drift in zone diameters. Quality Assurance will often pick up trends before the controls go out of range.

Figure 1. Use of control strain to monitor performance of disc diffusion susceptibility testing.



A free, supported QC programme is available from the following website:
<http://www.thehealthcarenet.com/shareware.htm>

6. Recognition of atypical results for clinical isolates

Atypical results with clinical isolates may indicate problems in testing that may or may not be reflected in zone diameters with control strains.

An organism with inherent resistance appears susceptible e.g. *Proteus* spp. susceptible to colistin or nitrofurantoin.

Resistance is seen in an organism when resistance has previously not been observed, e.g. penicillin resistance in Group A streptococci.

Resistance is seen in an organism when resistance is rare or has not been seen locally, e.g. vancomycin resistance in *Staphylococcus aureus*.

Incompatible susceptibilities are reported, e.g. a methicillin resistant staphylococcus reported susceptible to a β -lactam antibiotic.

In order to apply such rules related to atypical results it is useful to install an 'expert' system for laboratory reporting to avoid erroneous interpretation.

7. Investigation of possible sources of error

If the control values are found to be outside acceptable limits on more than one occasion during a reading frame of twenty tests, investigation into the possible source of error is required. Possible problem areas are indicated in table 1.

Table 1: Potential sources of error in disc diffusion antimicrobial susceptibility testing.

Possible source of error	Detail to check
Test conditions	Excessive pre-incubation before discs applied Excessive pre-diffusion before plates incubated Incorrect incubation temperature Incorrect incubation atmosphere Incorrect incubation time Inadequate illumination of plates when reading Incorrect reading of zone edges
Medium	Required susceptibility testing agar not used Not prepared as required by the manufacturer's instructions Batch to batch variation Antagonists present (e.g. with sulphonamides and trimethoprim) Incorrect pH Incorrect divalent cation concentration Incorrect depth of agar plates Agar plates not level Expiry date exceeded
Antimicrobial discs	Wrong agent or content used Labile agent possibly deteriorated Light sensitive agent left in light Incorrect storage leading to deterioration Disc containers opened before reaching room temperature Incorrect labelling of disc dispensers Expiry date exceeded
Control strains	Contamination Mutation Incorrect inoculum density Uneven inoculation Old culture used

8. Reporting susceptibility results when controls indicate problems

Microbiologists must use a pragmatic approach, as results from repeat testing are not available on the same day. If results with control strains are out of range the implications for test results need to be assessed.

Control results out of range

If control zones are below range but test results are susceptible, or control zones are above range but test results are resistant, investigate possible sources of error but report the test results. Otherwise it may be necessary to suppress reports on affected agents, investigate and retest.

Atypical results

If results are atypical with clinical isolates, the purity of the isolate and identification should be confirmed and the susceptibility repeated. Suppress the results for individual agents and retest.

Table 1: Acceptable zone diameter (mm) ranges for control strains on Iso-Sensitest agar, plates incubated at 35-37 °C in air for 18-20 h.

Antimicrobial agent	Disc content (µg unless stated)	<i>Escherichia coli</i>			<i>Pseudomonas aeruginosa</i>		<i>Staphylococcus aureus</i>		<i>Enterococcus faecalis</i>
		NCTC 10418	ATCC 25922	NCTC 11560 ¹	NCTC 10662	ATCC 27853	NCTC 6571	ATCC 25923	ATCC 29212
Amikacin	30	24-27	23-27	-	21-30	26-32	-	-	-
Ampicillin	10	21-26	16-22	-	-	-	-	-	26-35
Ampicillin	25	24-30	21-28	-	-	-	-	-	-
Aztreonam	30	39-44	36-40	-	27-30	26-30	-	-	-
Azithromycin	15	-	-	-	-	-	-	-	15-21
Cefixime	5	32-36	27-30	-	-	-	-	-	-
Cefoxitin	30	28-33	26-30	-	-	-	-	-	-
Cefotaxime	30	36-45	34-44	-	20-29	20-24	-	-	-
Ceftazidime	30	32-40	31-39	-	29-37	27-35	-	-	-
Cefuroxime	30	25-32	24-29	-	-	-	-	-	-
Cefalexin	30	21-28	16-21	-	-	-	-	-	-
Cefradine	30	19-25	16-22	-	-	-	-	-	-
Chloramphenicol	10	21-27	20-29	-	-	-	20-26	19-27	-
Ciprofloxacin	1	31-40	31-37	-	21-28	24-30	25-32	17-22	14-19
Ciprofloxacin	5	-	-	-	29-37	31-37	-	-	21-27
Clindamycin	2	-	-	-	-	-	30-35	26-33	No zone
Co-amoxiclav	3	-	-	-	-	-	-	27-32	-
Co-amoxiclav	30	18-31	20-26	12-18	-	-	-	-	-
Colistin	25	15-19	16-20	-	17-20	16-20	-	-	-
Doxycycline	30	-	-	-	-	-	35-40	33-37	-
Ertapenem	10	35-41	35-39	-	-	-	-	-	-
Erythromycin	5	-	-	-	-	-	22-31	22-29	-
Fusidic acid	10	-	-	-	-	-	32-40	30-37	-
Gentamicin	10	21-27	21-27	-	20-26	22-28	24-30	22-29	-
Gentamicin	200	-	-	-	-	-	-	-	22-27
Imipenem	10	32-37	33-37	-	20-27	23-28	-	-	28-32
Levofloxacin	1	30-33	28-34	-	-	-	-	-	-
Levofloxacin	5	-	-	-	22-29	23-29	-	-	-
Linezolid	10	-	-	-	-	-	26-33	26-30	24-29

Antimicrobial agent	Disc content (µg unless stated)	<i>Escherichia coli</i>			<i>Pseudomonas aeruginosa</i>		<i>Staphylococcus aureus</i>		<i>Enterococcus faecalis</i>
		NCTC 10418	ATCC 25922	NCTC 11560 ¹	NCTC 10662	ATCC 27853	NCTC 6571	ATCC 25923	ATCC 29212
Meropenem	10	38-42	27-39	-	26-33	32-39	-	-	22-28
Minocycline	30	-	-	-	-	-	34-39	33-36	-
Mupirocin	5	-	-	-	-	-	26-35	24-34	-
Mupirocin	20	-	-	-	-	-	30-38	27-35	-
Nalidixic acid	30	28-36	26-32	-	-	-	-	-	-
Neomycin	10	-	-	-	-	-	-	21-27	-
Netilmicin	10	22-27	22-26	-	17-20	20-24	-	22-28	-
Nitrofurantoin	200	25-30	23-27	-	-	-	21-25	20-26	-
Norfloxacin	2	34-37	32-36	-	-	-	-	-	-
Ofloxacin	5	31-37	31-38	-	18-26	18-25	-	-	-
Penicillin	1 unit	-	-	-	-	-	32-40	28-36	-
Piperacillin	75	30-35	27-32	-	27-35	27-34	-	-	-
Piperacillin/ tazobactam	85	30-35	26-31	-	28-35	28-35	-	-	26-32
Quinupristin/ dalfopristin	15	-	-	-	-	-	27-31	-	12-19
Rifampicin	2	-	-	-	-	-	27-39	29-36	-
Streptomycin	10	18-24	17-22	-	-	-	-	-	-
Teicoplanin	30	-	-	-	-	-	17-23	16-20	19-25
Tetracycline	10	23-29	22-28	-	-	-	31-40	26-35	-
Ticarcillin	75	32-35	27-30	-	24-28	23-27	-	-	-
Ticarcillin/ clavulanic acid	85	33-37	27-31	-	25-29	24-27	-	-	-
Tigecycline	15	29-32	28-32	-	-	-	29-34	27-30	26-31
Tobramycin	10	24-27	-	-	23-30	26-32	-	25-31	-
Trimethoprim	2.5	30-37	25-31	-	-	-	25-30	20-28	28-35
"	5	-	-	-	-	-	24-34	-	-
Vancomycin	5	-	-	-	-	-	14-20	13-17	13-19

26 β-Lactamase producing strain

Table 2: Acceptable zone diameter (mm) ranges for control strains on Iso-Sensitest agar supplemented with 5% defibrinated horse blood, with or without the addition of NAD, plates incubated at 35-37°C in air for 18-20 h.

Antimicrobial agent	Disc content (µg unless stated)	<i>Staphylococcus aureus</i>		Group A streptococci	
		NCTC 6571	ATCC 25923	NCTC 8198	ATCC 19615
Clindamycin	2	-	-	25-28	29-35
Erythromycin	5	22-29	23-29	-	-
Penicillin	1 unit	30-41	27-35	-	-
Tetracycline	10	30-38	28-36	-	-

Table 3: Acceptable zone diameter ranges for control strains for detection of methicillin/oxacillin/cefoxitin resistance in staphylococci (methicillin/oxacillin incubated at 30°C; cefoxitin incubated at 35°C).

Antimicrobial agent	Medium	Disc content (µg)	<i>Staphylococcus aureus</i>		
			NCTC 6571	ATCC 25923	NCTC 12493 ^a
Methicillin	Columbia/Mueller Hinton agar + 2% NaCl	5	18-30	18-28	No zone
Oxacillin	Columbia/Mueller Hinton agar + 2% NaCl	1	19-30	19-29	No zone
Cefoxitin	ISA	10	26-31	24-29	10-20

^a Methicillin/oxacillin/cefoxitin- resistant strain.

Table 4: Acceptable zone diameter (mm) ranges for control strains on Iso-Sensitest agar supplemented with 5% defibrinated horse blood and NAD, plates incubated at 35-37°C in 10% CO₂/10% H₂/80% N₂ for 18-20 h.

Antimicrobial agent	Disc content (µg unless stated)	<i>Bacteroides fragilis</i> NCTC 9343	<i>Bacteroides thetaiotaomicron</i> ATCC 29741	<i>Clostridium perfringens</i> NCTC 8359
Clindamycin	2	13-27	11-25	23-28
Co-amoxiclav	30	43-49	-	40-45
Meropenem	10	42-50	36-43	39-45
Metronidazole	5	34-43	26-40	11-23
Penicillin	1 unit	6	6	26-30
Piperacillin/tazobactam	75/10	41-48	-	37-43

Table 5: Acceptable zone diameter (mm) ranges for control strains on Iso-Sensitest agar supplemented with 5% defibrinated horse blood with or without the addition of NAD, plates incubated at 35-37°C in 4-6% CO₂ for 18-20 h.

Antimicrobial agent	Disc content (µg unless stated)	<i>Pasteurella</i>	<i>Neisseria</i>	<i>Staphylococcus</i>		<i>Haemophilus</i>		<i>Streptococcus</i>
		<i>multocida</i>	<i>gonorrhoeae</i> (with NAD)	<i>aureus</i>	<i>aureus</i>	<i>influenzae</i> (with NAD)	<i>pneumoniae</i>	
		NCTC 8489	NCTC 12700	NCTC 6571	ATCC 25923	NCTC 11931	ATCC 49247 ^a	ATCC 49619
Amoxicillin	2	-	-	29-34	-	-	-	-
Ampicillin	2	-	-	-	-	22-30	6-13	-
Ampicillin	10	32-37	-	-	-	-	-	-
Azithromycin	15	-	30-40	-	-	24-36	20-30	-
Cefixime	5	-	33-44	-	-	-	-	-
Cefotaxime	5	35-41	32-44	26-32	-	33-45	27-38	-
Ceftriaxone	5	-	33-47	-	-	-	-	-
Cefuroxime	5	-	23-32	22-29	24-29	22-28	6-16	-
Chloramphenicol	10	-	-	21-26	-	30-40	30-38	21-29
Ciprofloxacin	1	31-37	40-50	22-29	18-23	32-40	33-44	14-21
Clindamycin	2	-	-	21-25	-	-	-	-
Co-amoxiclav	3	-	-	-	-	20-27	10-20	-
Ertapenem	10	-	-	-	-	30-38	25-34	35-40
Erythromycin	5	-	20-29	25-29	-	12-23	9-16	23-36
Linezolid	10	-	-	22-26	-	-	-	-
Nalidixic acid	30	-	32-40	9-17	9-17	33-38	-	-
Oxacillin	1	-	-	-	-	-	-	8-16
Penicillin	1 unit	24-28	12-20	37-44	29-36	-	-	-
Quinupristin/ dalfopristin	15	-	-	-	-	-	-	21-29
Rifampicin	2	-	26-34	32-37	-	-	-	-
Rifampicin	5	-	-	-	-	-	-	28-35
Spectinomycin	25	-	17-23	-	-	-	-	-
Teicoplanin	30	-	-	14-19	-	-	-	-
Tetracycline	10	29-34	27-35	33-40	27-34	27-35	9-14	26-36
Tigecycline	15	-	-	27-30	24-28	-	-	26-30
Trimethoprim	2.5	-	-	-	-	30-40	28-36	-
Vancomycin	5	-	-	12-16	-	-	-	-

^a β-Lactamase-negative, ampicillin-resistant strain

9. Control of MIC determination

Tables 6-10 provide target MIC (mg/L) values for recommended control strains by BSAC methodology.^{1,2} MICs should be within one two-fold dilution of the target values.

Table 6: Target MICs (mg/L) for *Haemophilus influenzae*, *Enterococcus faecalis*, *Streptococcus pneumoniae*, *Bacteroides fragilis* and *Neisseria gonorrhoeae* control strains by BSAC methods

Antimicrobial agent	<i>Haemophilus influenzae</i>		<i>Enterococcus faecalis</i>	<i>Streptococcus pneumoniae</i>	<i>Bacteroides fragilis</i>	<i>Neisseria gonorrhoeae</i>
	NCTC	ATCC	ATCC	ATCC	NCTC	ATCC
	11931	49247	29212	49619	9343	49226
Amikacin	-	-	128	-	-	-
Amoxicillin	0.5	4	0.5	0.06	32	0.5
Ampicillin	-	-	1	0.06	32	-
Azithromycin	2	2	-	0.12	-	-
Azlocillin	-	-	-	-	4	-
Aztreonam	-	-	>128	-	2	-
Cefaclor	-	128	>32	2	>128	-
Cefamandole	-	-	-	-	8	-
Cefixime	0.03	0.25	-	1	64	-
Cefotaxime	-	0.25	32	0.06	4	-
Cefoxitin	-	-	-	-	4	-
Cefpirome	0.06	0.5	16	-	16	-
Cefpodoxime	0.12	0.5	>32	0.12	32	-
Ceftazidime	0.12	-	>32	-	8	-
Ceftriaxone	-	-	>32	0.06	4	-
Cefuroxime	2	16	>32	0.25	32	-
Cephadroxil	-	-	>32	-	32	-
Cefalexin	-	-	>32	-	64	-
Cephalothin	-	-	16	-	-	-
Chloramphenicol	-	-	4	4	4	-
Ciprofloxacin	0.008	0.008	1	1	2	0.004
Clarithromycin	8	4	-	0.03	0.25	0.5
Clindamycin	-	-	8	0.12	0.5	-
Co-amoxiclav	0.5	8	0.5	0.06	0.5	0.5
Cotrimoxazole	-	1	2	4	-	-
Quinupristin/ dalfopristin	-	-	1	0.5	16	-
Enoxacin	-	-	-	-	1	-
Ertapenem	0.12	0.5	-	0.12	0.25	-
Erythromycin	8	8	4	0.12	1	0.5
Faropenem	-	-	-	0.06	1	-
Fleroxacin	-	-	-	-	4	-
Flucloxacillin	-	-	-	-	16	-
Fucidic acid	-	-	2	-	-	-
Gatifloxacin	0.008	-	0.25	0.25	0.5	0.004
Gemifloxacin	0.12	-	0.03	0.03	0.25	0.002
Gentamicin	-	-	8	-	128	-
Grepafloxacin	-	0.004	-	0.25	-	-
Imipenem	-	-	0.5	-	0.06	-
Levofloxacin	0.008	0.015	1	0.5	0.5	0.008
Linezolid	-	-	-	2	4	-
Loracarbef	-	128	>32	2	>128	-
Mecillinam	-	-	>128	-	>128	-
Meropenem	-	-	2	-	0.06	-

Antimicrobial agent	<i>Haemophilus influenzae</i>		<i>Enterococcus faecalis</i>	<i>Streptococcus pneumoniae</i>	<i>Bacteroides fragilis</i>	<i>Neisseria gonorrhoeae</i>
	NCTC	ATCC	ATCC	ATCC	NCTC	ATCC
	11931	49247	29212	49619	9343	49226
Metronidazole	-	-	-	-	0.5	-
Moxalactam	-	-	-	-	0.25	-
Moxifloxacin	0.03	0.03	0.25	0.5	-	0.004
Naladixic acid	-	1	-	>128	64	-
Nitrofurantoin	-	-	8	-	-	-
Norfloxacin	-	-	2	-	16	-
Ofloxacin	-	-	2	-	1	-
Oxacillin	-	-	-	1	-	-
Pefloxacin	-	-	-	-	1	-
Penicillin	-	4	2	0.5	16	-
Piperacillin	-	-	2	-	2	-
Rifampicin	-	-	2	0.03	-	-
Roxithromycin	16	16	-	0.12	2	-
Rufloxacin	-	-	-	-	16	-
Sparfloxacin	-	0.002	-	0.25	1	-
Teicoplanin	-	-	0.25	-	-	-
Telithromycin	1	2	0.008	0.008	-	0.03
Tetracycline	-	16	16	0.12	0.5	-
Ticarcillin	-	-	-	-	4	-
Tigecycline	-	-	0.12	0.06	-	-
Tobramycin	-	-	16	-	-	-
Trimethoprim	-	-	0.25	4	16	-
Trovaflaxacin	0.008	0.002	0.06	0.12	0.12	-
Vancomycin	-	-	2	0.25	16	-

Table 7: Target MICs (mg/L) for *Escherichia coli*, *Pseudomonas aeruginosa* and *Staphylococcus aureus* control strains by BSAC methods

Antimicrobial agent	<i>Escherichia coli</i>		<i>Pseudomonas aeruginosa</i>		<i>Staphylococcus aureus</i>		
	NCTC 10418	ATCC 25922	NCTC 10662	ATCC 27853	NCTC 6571	ATCC 25923	ATCC 29213
Amikacin	0.5	1	2	2	1	-	2
Amoxicillin	2	4	>128	>128	0.12	0.25	0.5
Ampicillin	2	4	>128	>128	0.06	-	-
Azithromycin	-	-	-	-	0.12	0.12	0.12
Azlocillin	4	-	4	-	0.25	-	-
Aztreonam	0.03	0.25	4	2	>128	-	>128
Carbenicillin	2	-	32	-	0.5	-	-
Cefaclor	1	2	>128	>128	1	-	1
Cefamandole	0.25	-	>128	>128	0.25	-	-
Cefixime	0.06	0.25	16	-	8	8	16
Cefotaxime	0.03	0.06	8	8	0.5	-	1
Cefotetan	0.06	-	>128	>128	4	-	-
Cefoxitin	4	-	>128	>128	2	-	-
Cefpirome	0.03	0.03	4	1	0.25	-	0.5
Cefpodoxime	0.25	0.25	128	>128	1	4	2
Ceftazidime	0.06	0.25	1	1	4	-	8
Ceftizoxime	0.008	-	-	-	2	-	-
Ceftriaxone	0.03	0.06	8	8	1	-	2
Cefuroxime	2	4	>128	>128	0.5	1	1
Cephadroxil	8	8	>128	>128	1	-	2
Cefalexin	4	8	>128	>128	1	-	4
Cephaloridine	-	-	>128	>128	0.06	-	-
Cephalothin	4	8	>128	>128	0.5	-	0.25
Cephradine	-	-	>128	>128	2	-	-
Chloramphenicol	2	4	128	-	2	-	2
Ciprofloxacin	0.015	0.015	0.25	0.25	0.12	0.5	0.5
Clarithromycin	-	-	-	-	0.12	0.12	0.12
Clindamycin	-	-	-	-	0.06	0.12	0.06
Co-amoxiclav	2	4	>128	128	0.12	0.12	0.25
Colistin	0.5	-	2	-	128	-	-
Cotrimoxazole	0.25	0.25	-	-	-	-	2
Quinupristin/ dalfopristin	-	-	-	-	0.12	0.25	0.25
Dirythromycin	-	-	-	-	1	-	1
Doxycycline	-	-	-	-	0.06	0.12	-
Enoxacin	0.25	-	1	-	0.5	-	-
Ertapenem	0.008	0.015	-	-	-	-	-
Erythromycin	-	-	-	-	0.12	0.5	0.25
Farapenem	0.25	-	>128	>128	0.12	-	-
Fleroxacin	0.06	0.12	1	-	0.5	-	-
Flucloxacillin	-	-	>128	>128	0.06	-	-
Flumequine	2	-	>128	>128	-	-	-
Fosfomycin	4	-	>128	>128	8	-	-
Fusidic acid	>128	-	-	-	0.06	0.12	0.06
Gatifloxacin	0.015	0.015	1	1	0.03	0.12	0.12
Gemifloxacin	0.008	0.008	0.25	0.25	0.015	0.03	0.03
Gentamicin	0.25	0.5	1	1	0.12	0.25	0.25
Grepafloxacin	0.03	0.03	0.5	-	0.03	-	-
Imipenem	0.06	0.12	2	1	0.015	-	0.015
Kanamycin	1	-	1	-	2	-	-
Levofloxacin	0.03	0.03	0.5	0.5	0.12	0.25	0.25

Antimicrobial agent	<i>Escherichia coli</i>		<i>Pseudomonas aeruginosa</i>		<i>Staphylococcus aureus</i>		
	NCTC 10418	ATCC 25922	NCTC 10662	ATCC 27853	NCTC 6571	ATCC 25923	ATCC 29213
Linezolid	-	-	-	-	0.5	1	-
Lomefloxacin	-	-	-	-	0.5	-	-
Loracarbef	0.5	1	>128	>128	0.5	-	1
Mecillinam	0.12	0.12	8	-	8	-	64
Meropenem	0.015	0.008	2	0.25	0.03	-	0.06
Methicillin	-	-	>128	>128	1	2	2
Mezlocillin	2	-	8	-	0.5	-	-
Minocycline	-	-	-	-	0.06	0.06	-
Moxalactam	0.03	-	8	-	8	-	-
Moxifloxacin	0.03	0.03	2	2	0.06	0.06	0.06
Mupirocin	-	-	-	-	0.25	0.25	0.12
Nalidixic acid	2	4	>128	>128	>128	128	128
Neomycin	-	-	32	-	0.12	-	-
Netilmicin	-	-	1	0.5	-	-	-
Nitrofurantoin	4	8	-	-	8	-	16
Norfloxacin	0.06	0.06	1	1	0.25	-	1
Ofloxacin	0.06	0.03	1	1	0.25	-	0.5
Oxacillin	-	-	>128	>128	0.25	0.25	0.5
Pefloxacin	0.06	-	0.5	-	0.25	-	-
Penicillin	-	-	>128	>128	0.03	0.03	0.12
Piperacillin	0.5	2	4	2	0.25	-	1
Rifampicin	16	-	-	-	0.004	0.015	0.004
Roxithromycin	-	-	-	-	0.25	0.5	0.5
Rufloxacin	0.5	-	8	-	1	-	-
Sparfloxacin	0.015	0.015	0.5	0.5	0.03	-	-
Sulphonamide	16	-	>128	>128	64	-	-
Teicoplanin	-	-	-	-	0.25	1	1
Telithromycin	-	-	-	-	0.03	0.06	0.06
Temocillin	2	-	>128	-	128	-	-
Tetracycline	1	2	-	32	0.06	-	0.5
Ticarcillin	1	-	16	-	0.5	-	-
Ticarcillin/ 4mg/L clavulanate	-	-	32	16	-	-	-
Tigecycline	0.12	0.12	-	-	0.12	-	-
Tobramycin	0.25	0.5	0.5	0.5	0.12	-	0.5
Trimethoprim	0.12	0.25	32	-	0.25	-	0.5
Trovafoxacin	0.015	0.015	0.5	0.5	0.015	0.03	0.03
Vancomycin	-	-	-	-	0.5	0.5	1

Table 8: Target MICs (mg/L) for *Pasteurella multocida* control strain by BSAC methods

Antimicrobial agents	<i>Pasteurella multocida</i>	
	NCTC 8489	
Ampicillin	0.12	
Cefotaxime	0.004	
Ciprofloxacin	0.008	
Penicillin	0.12	
Tetracycline	0.25	

Table 9: Target MICs (mg/L) for anaerobic control strains by BSAC methods on Iso-Sensitest agar supplemented with 5% defibrinated horse blood and 20 mg/L NAD

Antimicrobial agent	<i>Bacteroides fragilis</i>	<i>Bacteroides thetaiotaomicron</i>	<i>Clostridium perfringens</i>
	NCTC 9343	ATCC 29741	NCTC 8359
Clindamycin	0.5	2	0.06
Co-amoxiclav (2:1 ratio)	0.5	0.5	≤ 0.06
Meropenem	0.06	0.12	≤ 0.015
Metronidazole	0.5	4	8
Penicillin	16	16	0.06
Piperacillin/tazobactam (fixed 4 mg/L tazobactam)	≤ 0.12	8	0.5

Table 10: Target MICs (mg/L) for Group A streptococci control strains by BSAC methods

Antimicrobial agent	Group A streptococci	
	NCTC 8198	ATCC 19615
Clindamycin	0.03	0.06

References

1. Andrews, J.M. Determination of minimum inhibitory concentrations. *Journal of Antimicrobial Chemotherapy*, Suppl S1 to Volume 48 July 2001.
2. Andrews, J. M., Jevons, G., Brenwald, N. and Fraise, A. for the BSAC Working Party on Sensitivity Testing. Susceptibility testing *Pasteurella multocida* by BSAC standardized methodology. *Journal of Antimicrobial Chemotherapy*.

Suppliers

Reagent	Suppliers (others may be available)
ISA	CM471, Oxoid, Basingstoke, UK
Columbia agar	CM331, Oxoid, Basingstoke, UK
Mueller Hinton agar	CM337, Oxoid, Basingstoke, UK
NAD	Mast Group, Merseyside, UK
McFarland turbidity standards	bioMérieux, Basingstoke, UK
Control strains	NCTC, Colindale, London Oxoid, Basingstoke, UK Mast Laboratories, Merseyside, UK Becton Dickinson, Oxford, UK TCS Biosciences Ltd. Buckingham, UK

Useful web sites

BSAC	British Society for Antimicrobial Chemotherapy	http://www.bsac.org.uk
SRGA	The Swedish Reference Group for Antibiotics	http://www.srga.org
CDC	Centre for Disease Control (Atlanta, USA)	http://www.cdc.gov
WHO	World Health Organisation (Geneva, Switzerland)	http://www.who.int
CLSI	Clinical and Laboratory Standards Institute	http://www.clsi.org
NEQAS	National External Quality Assessment Scheme	http://www.ukneqas.org.uk
NCTC	National Collection of Type Cultures	http://www.ukncc.co.uk
JAC	The Journal of Antimicrobial Chemotherapy	http://www.jac.oupjournals.org
EUCAST	European Committee on Antimicrobial Susceptibility Testing	http://www.eucast.org