

INFECTION PREVENTION AND CONTROL AUDIT REPORT AND ACTION PLAN

Practice Name: Arcadian Gardens NHS Medical Centre - F85034 Practice Code: F85034

GP Introduction

Introduction

The Health and Social Care Act (2008): code of practice on the prevention and control of infections and related guidance (revised 2015) outlines the role of infection prevention (including cleanliness) and optimising antimicrobial use and reducing antimicrobial resistance. This applies to NHS bodies and providers of independent healthcare and adult social care in England, including primary dental care, independent sector ambulance providers and primary medical care providers.

Effective prevention of infection must be part of everyday practice and be applied consistently by everyone. It is also a component of good antibiotic stewardship as preventing infections helps to reduce the need for antimicrobials. Good management and organisational processes are also crucial to ensure that high standards of infection prevention and control (including cleanliness) are set up and maintained Care Quality Commission (CQC) under the Health and Social Care Act 2008 (Regulated Activities) Regulations 2014: Regulation 12 ensures that the care people receive meets essential standards of quality and safety and encourages ongoing improvements by those who provide or commission care.

Providers must assess the risks to people's health and safety during any care or treatment and make sure that staff have the qualifications, competence, skills and experience to keep people safe. Where the responsibility for care and treatment is shared, care planning must be timely to maintain people's health, safety and welfare.

Providers must also ensure that the premises and any equipment used is safe, fit for purpose and where applicable, available in sufficient quantities. Medicines must be supplied in sufficient quantities, managed safely and administered appropriately to make sure people are safe.

Infection Prevention and Control in Primary Care General Practice

In primary care general practice there are a variety of ways where cross infection may be a potential risk, some of them may include:

- · Overcrowding.
- · Staff health
- · Poor hand hygiene.
- · Inadequate equipment cleaning / decontamination.
- \cdot Sharps management and transmission of blood borne viruses.
- · Specimen handling
- · Clinical waste segregation and management.
- \cdot Environmental cleaning and blood / bodily fluid spillages management.

All of these modes of transmission of infection are either avoidable or the risks can be minimised to low levels if the appropriate standard infection control procedures are followed. These measures include Hand hygiene, equipment management and decontamination, maintaining Aseptic Non-Touch Technique (ANTT), medicine management including cold chain system, environmental cleanliness and spillage management, wastes and sharps management, adhering to the correct usage of Personal Protective Equipment (PPE), staff health and management of infectious diseases. Hand hygiene is the single most effective means of preventing health care associated infections and should be a priority for implementation.

The Audit Process

NHS England London has placed increased emphasis on the use of audit to measure compliance to practice and the implementation of policies and procedures relating to infection control. The review of clinical practice through audit is a well-established means of monitoring and improving the quality of care and of supporting the implementation of change in practice.

As part of the process of ensuring that IPC standards are met, as well as ensuring that the quality of the infection control practice within primary care is of a high standard, NHS England London implements a continuing programme of practice visits, supported by experienced NEL Infection Prevention and control team.

The Audit Tool used is a consistent resource developed by NHS England London and NEL in collaboration with London Wide Local Medical Committees and Surrey and Sussex Local Medical Committees and is part of London region's Standard Operating Procedure, adopted by each Primary Care Commissioning Committee. The audit tool consists of several standards of IPC and defines acceptable criteria which minimises the risk of infection to patients, staff and relatives. The tool has recently been revised to take into account changes in standards, legislation and learning, where applicable to general medical practice, together with learning from more than 6 months of piloting and subsequent evaluation of the audit tool.

This new revised audit tool will present improved mechanisms for measuring compliance, identify best practice, and discover risks and vulnerabilities that may not have come to light using the current audit tool.

The audit tool standards and rating system have been updated to:

• More closely align with CQC registration compliance guidance where applicable

• provide clarification in terms of the requirements of primary medical contract holders related to the 'Health & Social Care Act 2008 code of practice on the prevention and control of infections', recognising that compliance with the act is a legal requirement, whereas the code of practice is considered guidance

• provide practices with a clearer understanding of the risks associated with non-compliance with each standard of the audit tool

- · assist practices with their own risk assessments
- · help practices to assure themselves that they are meeting infection control requirements in relation to CQC

The audit report and its recommendations help to ensure that practices improve their compliance to infection prevention and control in line with the Health & Social Care Act 2008 code of practice on the prevention and control of infections code of practice and other current national guidelines and should serve as a useful reference point.

Therefore it is essential that this report and its recommendations is given due consideration and that the agreed action plan which outlines how the practice plan to address the issues highlighted is completed and returned appropriately as advised.

Contractor Details

Practice Name	Arcadian Gardens NHS Medical Centi
Practice Contract Number	F85034
Email Address	1 Arcadian Gardens N22 5A
Telephone No	02088811957
Email address	savvakis.georgiou@nhs.ne
Practice Manager Name	Savvakis Georgiou
Principal GP Contractor	Dr Perera and Dr Karim
Name of IPC Lead for Practice	Pedro Daniel
Practice Nurse	Patricia Abebrese
Accompanying Practice Staff (One of whom	Say and Dadra
must be the IPC Lead for the Practice)	Sav and Pedro
Date audit completed	2024-12-11
Audit carried out by	Mehrunessa Eddoo
IUCD Fitting	No
	No
Does the practice undertake minor surgery	Level: None

Executive Summary

This is the report of infection prevention and control (IPC) audit carried out at Arcadian Gardens NHS Medical Centre - F85034 on 2024-12-11. The audit was completed with the assistance of Sav and Pedro.

The practice has 5 consulting rooms and 1 treatment room. The practice does not undertake minor surgery. The practice has 3 vaccine fridges.

Findings:

The practice shows full compliance on all standards of IPC. Overall the practice showed full compliance with all standards of IPC designated as Essential Quality Requirements.

No further findings.

Recommendations:

The practice is fully compliant, no action plan recommended. Advice was provided on the day of the audit on the importance of maintaining good IPC standards.

No further recommendations.

Risk assessment:

No risk assessment comments.

Next face-to-face audit: Next routine audit.

FURTHER CONSIDERATIONS

Thank you for colaborating with our audit.

Please complete this satisfaction survey to help us improve: <u>NHS England Satisfaction</u> <u>Survey</u> (click to open on a new window)

NHS England Contact Details

Practice Details

	Converted residential
Practice Building	Owned
No of Consulting Rooms	5
No of Treatment Rooms	1
	No
Minor surgery	None
IUCD Fitting	No
	Yes
	Quantity: 3
Vaccine Fridge	Location: Stock room and room 3
	Make & SN: LEC PESR47 10568, PSRC273UK WMSFR2 and
	Yes
Cleaners storage facilities	Limited space
Dirty Sluice Room	No
Clean Utility Room	No

Action Plan

The next section contains all details about the non-compliances found during the audit.

By now, you should have received an onboarding email to access MEG, where you will be able to manage the action plan and include the corrective actions.

If you haven't received any details or have any questions, please contact <u>england.phipc@nhs.net</u>

How to use MEG and access Action Plan (click to open on a new window)

IPC ACTION PLAN

No data yet!

Overall Compliance

100.0%

Compliance by Standard

General Management	Staff Health	Environment
100.0%	100.0%	100.0%
Hand Hygiene 100.0%	Personal protective equipment 100.0%	Prevention and management of spillages
Safe handling and disposal of sharps 100.0%	Waste Management Policy and Procedures 100.0%	Management of Specimens
Decontamination of medical devices	Clinical Rooms	Vaccine Storage and Cold Chain
Notification of infectious diseases and contamination 100.0%	Antimicrobial Stewardship (AMS)	Minor Surgery rooms No data yet!

Full Audit Tool

1- Management of IPC - General Management

12 0 0 0		12 / 12 - 100.0%		
Question	Issue	Comment	Attachments	Compliance
1. Is there a named clinical lead person in the practice for infection prevention and control? [M]	-	-	-	Compliant
2. Does the practice have infection prevention and control policies? [M]	-	-		Compliant
3. Is infection prevention and control included in all staff induction programmes? [M]	-	-	-	Compliant
4. Evidence to show that all clinical and non- clinical staff are up to date with IPC training specific to their roles? [M]	-	-	•	Compliant
5. Is there a process for internally recording/reporting untoward incidents in relation to infection prevention and control ? [M]	-	-	•	Compliant
6.1 Local IPC advice and support as needed [EQR & E]	-	-	-	Compliant
6.2 Local Hospital Consultant Microbiologists [EQR & E]	-	-	-	Compliant
6.3 Public Health England Health Protection teams [EQR & E]	-	-	-	Compliant
6.4 Local anti-microbial Pharmacy Lead [EQR & E]	-	•		Compliant
7 Does the practice have documentary evidence of IPC audits undertaken, evaluated and with actions taken to improve practice standards? [M]	-	-	•	Compliant
8 Has the Practice carried out a risk assessment for Legionella? [M]	-	The legionella risk assessment was completed by Envirisk Ltd in November 22. Another assessment has been undertaken and the report is awaited		Compliant
9 Does the practice have a written scheme for prevention of Legionella contamination in water pipes and other water lines? [M]	-	-	•	Compliant

Comments []

2- Management of IPC - Staff Health

4 0 0 0		4 / 4 - 100.0%						
Question	Issue	Comment	Attachments	Compliance				
1. Have all staff at risk been immunised against hepatitis B? [M]	-	-	-	Compliant				
2. Are all staff routinely advised regarding immunisation against seasonal influenza? [EQR & E]	-			Compliant				
3. Does the practice have access to Occupational Health service or access to appropriate occupational health advice? [M]	-	Occupational health services are provided by Bart's Health.	•	Compliant				
4. Has the issue of immunity to Measles, Rubella and Varicella in clinical staff been considered in the practice and a risk assessment undertaken? [M]	-	-		Compliant				

Comments I1

3- Environment

Q		
Č	0	

8 / 8 - 100.0%

Question	Issue	Comment	Attachments	Compliance
1. Are all areas including clinical areas and equipment visibly clean and free from extraneous items? [M]	-	-		Compliant
2. Are there comprehensive written specifications for cleaning the environment and equipment in the practice? [M]	-	-	•	Compliant
3. Are there up to date cleaning schedules which includes regular cleaning of clinical, admin and sanitary areas? [M]	-	-	•	Compliant
4. Are walls in all areas in good condition, intact and have smooth easy-to-clean surfaces? [EQR]	-	•	•	Compliant
5. Is flooring in all areas that are accessible to patients in a good state of repair and easy-to-clean? [EQR]	-	The steps to Dr Perera's room is damaged and plans are in place to have it replaced. There are plans for the wooden laminate in the waiting area to be replaced with wipeable Lino.		Compliant
6. Are furniture in clinical areas and other areas accessible to patients impermeable / washable / suitable for its use? [EQR]	-	Plans are for cork boards to be replaced.		Compliant
7. Are cleaning equipment and materials for cleaning colour coded, suitable for use and stored appropriately? [EQR]	-	•	•	Compliant
8. Is the area for storing cleaning equipment well ventilated, clean and tidy (no clutter) and is it of an adequate size? [EQR]	-	-		Compliant

Comments I1

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4- Hand Hygiene

13	0	0	0

13 / 13 - 100.0%

Question	Issue	Comment	Attachments	Compliance
1. Does the practice has a Hand Hygiene Policy? [M]	-	-	-	Compliant
2. Is the hand hygiene technique displayed as a laminated poster adjacent to the hand washbasin or is it featured on the soap dispenser? [EQR]	-	-		Compliant
3.Does your practice policy demonstrate an awareness of the DH uniform policy? [EQR]	-	-		Compliant
4. Are there wash basins dedicated to hand hygiene in each clinical and consulting room which can be easily accessed? [EQR]	-	-		Compliant
5. Do all hand wash basins for use in connection with clinical procedures have elbow or wrist operated mixer taps? [EQR]	-	-		Compliant
6. Is the hot water thermostatically controlled? [EQR]	-	-		Compliant
7. Are taps at all clinical hand wash basins free from swan neck type taps? [EQR]	-	-	-	Compliant
8. Are all hand wash basins free from plugs? [EQR]	-	-	-	Compliant
9. Are all hand wash basins in clinical and consulting rooms free from an overflow? [EQR]	-	-	-	Compliant
10. Are hand hygiene facilities clean and free from clutter? [EQR]	-	-	-	Compliant
11. Are hand hygiene facilities free from damage? [EQR]	-	-	-	Compliant
12. Is the tap off-set from the waste outlet? [EQR]	-	-	-	Compliant
13. Is liquid soap dispensed from single use cartridges or bottles? [EQR]	-	-	-	Compliant

4 0 0 0

4 / 4 - 100.0%

Question	Issue	Comment	Attachments	Compliance
14. Is alcohol-based hand rub available for use when required, including use during domiciliary visit? [EQR]	-	-		Compliant
15. Are paper hand towels available? [EQR]	-	-	-	Compliant
16. Are hand wash basins free from nail brushes? [EQR]	-	-	-	Compliant
17. Are there separate arrangements available to dispose of waste materials other than using the hand washbasin? [EQR]	-	-		Compliant

Comments 11

5- Personal Protective Equipment (PPE)

9 / 9 - 100.0%

Question	Issue	Comment	Attachments	Compliance
1. Does the practice have a policy on the appropriate use of PPE? [M]	-	-	-	Compliant
2.1 Gloves (sterile/non-sterile) are appropriate for use, i.e, latex & latex free nitrile? [EQR]	-	-		Compliant
2.2 Disposable aprons available? [EQR]	-	-	-	Compliant
2.3 Disposable face and eye protection? [EQR]	-	-	-	Compliant
3. Are staff aware of the principles of wearing and disposing of PPE? [EQR]	-	-	-	Compliant
4. Are PPE items worn as single use items? [EQR]	-	-	-	Compliant
5. Where required are aprons and gloves changed between different episodes of care on the same patient? [EQR]	-	-	-	Compliant
6. Are gloves removed and hand hygiene performed after every clinical activity? [EQR]	-	-	-	Compliant
7. Are staff aware on the decontamination process required for re-usable goggles? [EQR]	-	-	-	Compliant

Comments 11

Adequate supply of PPE is available

6- Prevention and management of spillages

4 0 0 0 4/	4 - 100.0%				
Question	I	ssue	Comment	Attachments	Compliance
1. Does the practice have a policy for managing spillages in healthcare [[EQR]	premises? -		-	-	Compliant
2. Are all staff aware of the procedure for dealing with spillages of blood fluids? [EQR]	l or other body -	-	-	-	Compliant
3. Are spillage kits available for dealing with spillages of blood/body flui	ds? [M] -	-	-		Compliant
4. Are disposable cloths or mop heads available for cleaning blood or of spillages? [EQR]	ther body fluid -	-	-		Compliant

Comments 1

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7- Safe handling and disposal of sharps

11	0	0	0
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11 / 11 - 100.0%

Question	Issue	Comment	Attachments	Compliance
1. Does the practice have a policy on safe handling & disposal of sharps? [M]	-	-	-	Compliant
2. Are sharps containers that conform to BS 7320 and UN3291 available in every clinical/consulting area? [M]	-	-	-	Compliant
3. Are sharps containers discarded when two thirds full and stored in a secure facility away from public access until collected for disposal? [EQR]	-	-	-	Compliant
4. Is blood sampling undertaken by using a 'Sharp Safe' single-use vacuum blood collection system? [EQR]	-	-	-	Compliant
5. Is Aseptic Non-Touch Technique (ANTT) used when performing venepuncture? [EQR]	-	-	-	Compliant
6. Sharps used for taking blood from patients at home/care home disposed of in to an appropriate sharps container? [EQR]	-	-	-	Compliant
7. Is there evidence that the practice has undertaken a review of sharps management? [EQR]	-	-	-	Compliant
8. Are sharps containers assembled according to manufacturer's instructions? [M]	-	-	-	Compliant
9. Are staff encouraged to wear gloves when undertaking venepuncture? [EQR & E]	-	-	-	Compliant
10. Are staff aware of the correct procedure to follow after a needle stick injury, other sharps or blood splash exposure? [M]	-	-	-	Compliant
11. Are posters available which show staff the emergency algorithm to follow in case of a sharp injury and is it up to date? [EQR]	-	-	-	Compliant

Comments I1

8- Waste Management Policy and Procedures

9	0	0	0

9 / 9 - 100.0%

Question	Issue	Comment	Attachments	Compliance
1. Does the practice have a policy on waste management? [M]	-	-	-	Compliant
2. Is there documentary evidence to show that all clinical waste is disposed of by a registered waste collection company? [M]	-	-	•	Compliant
3. Are records of waste transfer and disposal arrangements kept and stored in accordance with the EPA 1990? [EQR]	-	-		Compliant
4. Are there easily accessible and compliant foot-operated & fully enclosed clinical waste bins available in each clinical area? [EQR]	-	-	-	Compliant
5. Is clinical and domestic waste correctly segregated? [EQR]	-	-	-	Compliant
6. Are clinical waste bags marked with the practice code when securing for disposal? [EQR]	-	-	-	Compliant
7. Are waste bags less than 2/3 full and securely tied? [EQR]	-	-	-	Compliant
8. Where clinical waste is not collected directly from clinical areas, is it stored in a separate, secure area for waste? [EQR]	-	-	-	Compliant
9. Are staff encouraged to report all incidents to the designated infection control lead at the practice? [EQR]	-	-	-	Compliant

Comments 11

Clinical waste is collected by Grundon Waste Contractors

9- Management of Specimens

3 0 0 1	3 / 3 - 100.0%				
Question		Issue	Comment	Attachments	Compliance
1. Does the practice have a policy or procedure for specimen handlin	ıg? [M]	-	-	-	Compliant
2. Where applicable are specimens stored in a dedicated refrigerator vaccines or medicines)? [EQR]	(not with food,	-	-	-	N/A
3. Are arrangements for specimen testing appropriate in consulting r	ooms? [EQR]	-	-	-	Compliant
4. Are staff aware of the appropriate way to handle and transport spe E]	cimens? [EQR &	-	-		Compliant

Comments 11

The practice provides a phlebotomy service and samples are collected once a day to be processed at the North Middlesex Hospital

10- Decontamination of medical devices

6	0	0	1

6 / 6 - 100.0%

Question	Issue	Comment	Attachments	Compliance
1. Does the practice have a policy which outlines the decontamination processes the GP Practices use for all medical devices? [M]	-	-	-	Compliant
2. Does the practice use single use surgical instruments? [EQR]	-	-	-	Compliant
3. Does the practice use an accredited external sterile supply service? [EQR]	-	-	-	N/A
4. Are all medical devices stored appropriately and above floor level to avoid contamination? [EQR]	-	-		Compliant
5. Are all items of sterile equipment within their use-by date? [EQR]	-	-	-	Compliant
6. All items of equipment that come into contact with patients cleaned or decontaminated according to guidelines or are disposed of after each use? [M]	-	-		Compliant
7. Is there a cleaning schedule/check list maintained for all items requiring cleaning? [EQR]	-	-		Compliant

Comments I1

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11- Clinical Rooms

8 0 0 0

8 / 8 - 100.0%

Question	Issue	Comment	Attachments	Compliance
1. Are all clinical rooms and all work surfaces clean and free from extraneous items? [M]	-	-		Compliant
2. Is flooring impervious to liquids, non-slip, intact and clean? [EQR]	-	-	-	Compliant
3. Does the flooring form a coved skirting OR is the gap between the floor and the skirting sealed and is the seal maintained? [EQR]	-	-		Compliant
4. Are walls and ceilings clean, dry and free from visible defects and have smooth easy to clean surfaces? [EQR]	•	-	-	Compliant
5. Is there an examination couch with an intact, impervious cover and single use roller paper available for use? [EQR]	•	-	-	Compliant
6. Is the examination couch fitted with a paper roll holder? [E]	-	-	-	Compliant
7. Are there sufficient work surfaces and dressing trolleys of smooth, impervious and cleanable material? [EQR]	-	-		Compliant
8. Are all treatment surfaces in the room cleaned every working day? [EQR]	-	-	•	Compliant

Comments 11

12- Vaccine Storage and Cold Chain

11	0	0	0	
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11 / 11 - 100.0%

Question	Issue	Comment	Attachments	Compliance
1. Does the practice have an up to date cold chain policy as per the standards of the Green Book and is this accessible to all staff? [M]	-	-		Compliant
2. Is there a designated person/s in the practice responsible for the ordering, delivery and storage of vaccines? [EQR]	-	-		Compliant
3. Are vaccines monitored for their expiry dates and the close to expiry stocks clearly labelled? [EQR]	-	-		Compliant
4. Is the refrigerator specialised for the storage of vaccines? [M]	-	-	-	Compliant
5. Are vaccines correctly stored to allow good air flow within the vaccine refrigerator? [EQR]	-	-		Compliant
6. Are there measures in place to prevent the fridge from being turned off? [EQR]	-	-	-	Compliant
7. Is/Are the vaccine fridge/s located in a well-ventilated area? [EQR]	-	-	-	Compliant
8. Is the temperature of the vaccine fridge continually monitored with a min/max thermometer and the temperatures are recorded each working day to ensure vaccines are maintained at 2-80C? (Min, max and actual fridge temperatures are recorded)? [M]	-	-	•	Compliant
9. Does the practice has a maintenance contract that allows for at least yearly servicing, calibration of the temperature gauge? [EQR]	-	-		Compliant
10. Is a second min/max thermometer or Data Logger temperature recording device available and used? [EQR]	-	-		Compliant
11. Is the fridge either self-defrosting or is it defrosted monthly or sooner if needed? [EQR]	-	-	•	Compliant

3	0	0	0

3 / 3 - 100.0%

Question	Issue	Comment	Attachments	Compliance
12. Is there a process in place for safe disposal of expired, damaged or surplus vaccines? [EQR]	-	-		Compliant
13. Does the practice have records of vaccines received? [EQR]	-	-	-	Compliant
14. Is there accessible written guidance on what staff should do in the event of a power cut or a temperature reading outside the required range? [EQR]	-	-		Compliant

Comments I1

The cold chain lead is Pedro. A service contract is in place with Williams Medical Ltd for the calibration and maintenance of the vaccine fridges.

13- Notification of infectious diseases and contamination

6 0 0 0	6 / 6 - 100.0%				
Question		Issue	Comment	Attachments	Compliance
1. Does the practice have a policy on managing patients with comm [M]	unicable diseases?	-	-	-	Compliant
2. Does the practice notify all reportable infectious disease on susp officer at the local authority? [M]	icion to the proper	-	-	•	Compliant
3. Does the practice have access to notification forms? [EQR]		-	-	-	Compliant
4. Does the practice notifying gastro intestinal disease (food poisor [EQR]	ning) on suspicion?	-	-	-	Compliant
5. Does the practice notify Gastro intestinal disease (food poisoning specimen results are received from the microbiology laboratory? [E		-	-	-	Compliant
6. Is the practice aware of the new requirements to notify cases of c other diseases? [EQR]	ontamination and	-	-	-	Compliant

Comments 11

Notification forms are accessible online

14- Antimicrobial Stewardship (AMS)

13	0	0	0

13 / 13 - 100.0%

Question	Issue	Comment	Attachments	Compliance
1. Are GP prescribers in the practice aware of the TARGET toolkit? [EQR]	-	-	-	Compliant
2. Have all GP prescribers completed the Antimicrobial Stewardship Self-Assessment Checklist available in TARGET? [EQR]	-	-		Compliant
2.1 Give number of GP prescribers using the self-assessment checklist against those who do not [EQR]	-	-	-	Compliant
3. Is the document "Antimicrobial prescribing and stewardship competencies" available and/or has it been read by prescribers in the practice? [EQR]	-	-	-	Compliant
3.1 Give number of prescribers who are aware of this document against those who are not [EQR]	-	-	-	Compliant
4. Are all the prescribers in the Practice aware of the Public Health England AMR local indicators? [EQR]	-	-		Compliant
5. Is the practice aware of how they may access their antibiotic prescribing data online? [EQR]	-	-	-	Compliant
6. Are all prescribers in the practice aware of the NICE guidelines on AMS [EQR]	-	-	-	Compliant
7. Do all prescribers give information to their service users of how they should correctly use antimicrobial medicines and the dangers associated with their overuse and misuse? [EQR]	-	-		Compliant
8. Are all prescribers aware of the UK's 5-year Antimicrobial Resistance Strategy? [E]	-	-	-	Compliant
9. Does the practice actively participates in the European Antibiotic Awareness Day/Week (EEAD) held in November each year? [E]	-	-	-	Compliant
10. Are all clinical staff in the Practice aware of the PHE Antibiotic Guardian campaign? [E]	-	-	-	Compliant
11. Does the practice have an identified sepsis lead / link? [E]	-	-	•	Compliant

3 0 0 0

3 / 3 - 100.0%

Question	Issue	Comment	Attachments	Compliance
12. Are clinicians/GPs aware of and/or have received training in identifying sepsis? [EQR]	-	-	-	Compliant
13. Does the practice promote the use of the GRASP Fever Audit tool? [E]	-	-	-	Compliant
14. Does the practice make use of the National Early Warning Signs (NEWS) tool and the Paediatric Early Warning Signs (PEWS) tool? [EQR]	-	-	•	Compliant

Comments 11

The sepsis lead is Dr Karim.

15- Minor Surgery rooms

0 0 0 13

0/0--

Question	Issue	Comment	Attachments	Compliance
1. Is there is a dedicated room for minor surgery or is the clinical room of sufficient standard for undertaking level 2 minor surgery? [M for Level 2&3 minor surgery]	-	-		
2. Are all sterile packs and other equipment stored appropriately and is there adequate storage space? [M for Level 2&3 minor surgery]	-	-		
3. Is the practice using disposable single use instruments, supplied by a recognised manufacturer of sterile disposable instruments? [M for Level 2&3 minor surgery]	-	-	-	
4. Where applicable, is an accredited external sterile supply service used for re-usable surgical instrument that need to be sterile at the point of use? [M for Level 2&3 minor surgery]	-	-	-	
5. Is the minor surgery room clean and free from extraneous items? [M]	-	-	-	
6. Are walls in good condition, intact and have smooth easy-to-clean surfaces? [M for Level 2&3 minor surgery]	-	-		
7. Is flooring impermeable, intact and have continuous edging coved up the walls? [M for Level 2&3 minor surgery]	-	-	-	
8. Are ceilings intact and free from visible cracks or visible defects? [M for Level 2&3 minor surgery]	-	-	-	
9. Are ceiling lights protected / enclosed from potential contamination? [M for Level 2&3 minor surgery]	-	-	-	
10. Does the room have adequate ventilation - natural or mechanical (no desktop fans)? [M for Level 2&3 minor surgery]	-	-	-	
11. Is the heat source and pipe work in the room enclosed to prevent accumulation of dust and dirt? [M for Level 2&3 minor surgery]	-	-		
12. Is the treatment couch intact and is protected with disposable paper towel that is changed after each patient? [M for Level 2&3 minor surgery]	-	-	•	
13 Are all work surfaces intact, smooth, and impervious easy to clean and are able to withstand cleaning with chemical disinfectants? [M for Level 2&3 minor surgery]	-	-	•	

0 0 0 11

Question	Issue	Comment	Attachments	Compliance
14. Are all wall cabinets intact with doors? [EQR]	-	-	-	
15. Does the clinical hand wash basin conforms to current recommended guidance (HTM 00-10; HBN 00-09)? [M for Level 2&3 minor surgery]	-	-	-	
16. Are there wall mounted dispensers for liquid soap and is the liquid soap dispensed via non-refillable cartridges? [M for Level 2&3 minor surgery]	-	-		
17. Are there wall mounted dispensers with good quality disposable paper hand towels? [M for Level 2&3 minor surgery]	-	-	-	
18. Is the clinical hand wash basin free from re-usable nail brushes? [M for Level 2&3 minor surgery]	-	-		
19. Are single use sterile and non-sterile gloves available in latex and non-latex nitrile material? [M for Level 2&3 minor surgery]	-	-		
20. Is there is a designated stainless steel trolley available for use in this room only? [M for Level 2&3 minor surgery]	-	-	-	
21. Is there a clean clinical waste bin with a foot pedal with waste bag fully enclosed and is it in good operating condition? [M for Level 2&3 minor surgery]	-	-		
22. Do all staff use recommended PPE when splashing of body fluids is anticipated? [M for Level 2&3 minor surgery]	-	-		
23. Are disposable sterile drapes available and used for level 2 and level 3 minor surgeries? [M for Level 2&3 minor surgery]	-	-		
24. Does the GP Practice audit post-operative wound infections and are records kept? [M for Level 2&3 minor surgery]	-	-	-	

Comments I1

-

Signatures

Mehrunessa Eddoo:



APPENDIX

Key - Infection Prevention & Control Questionnaire

- **Mandatory** [M]: either required on a statutory and/or contractual basis and aims to align with relevant CQC requirements that result in CQC reports specifying that a practice 'must do...'
- An Essential Quality Recommendation [EQR]: the minimum expected standards for compliance as detailed in the Health and Social Care Act 2008 (Hygiene Code). EQRs will typically align with relevant CQC requirements that result in CQC reports specifying a practice 'should do...'
- Educational [E]: these are best practice standards, which align with what practices should 'know about'. Contractors are advised to record that there has been a discussion about these standards within the practice.

Section 1: The Management of Infection Prevention and Control (General Management)

Standard: Infection prevention and control is managed effectively and complies with the Health and Social Care Act 2008: Code of practice on the prevention and control of infection and related guidance (July 2015)

		M= Mandatory	
#	Questions	EQR= EQR	Risk Level
		E= Educational	
1	Is there a named clinical lead person in the practice for infection prevention and control?	M This is a requirement of Criterion 2 of the Health & Social Care Act 2008 (Amended 2015), which states that: The IPC Lead should: • be responsible for the organisation's infection prevention (including cleanliness) management and structure and the establishment of a water safety group; • oversee local prevention of infection policies and their implementation; • report directly to the registered provider; • have the authority to challenge inappropriate practice, if appropriate, including antimicrobial prescribing practice; • have the authority to set and challenge standards of cleanliness; • assess the impact of all existing and new policies on infections and make recommendations for change; • be an integral member of the organisation's governance, water safety group, and safety teams and structures where they exist;	Moderate
		and • produce an annual statement with regard to compliance with practice on infection prevention and cleanliness and make it available on request.	
2	Does the practice have infection prevention and control policies?	M Criterion 9 of The Health & Social Care Act 2008 (Updated 2015) requires registered provider to: Have and adhere to policies, designed for the individual's care and provider organisations that will help to prevent and control infections. 9 A registered provider should, in relation to preventing, reducing and controlling the risks of infections, have in place the appropriate policies concerning the matters mentioned in a) to y) below. All policies should be clearly marked with a review date and the review date adhered to. 9 Any registered provider should have policies in place relevant to the regulated activity it provides. Each policy should indicate ownership (i.e. who commissioned and retains managerial responsibility), authorship and by whom the policy will be applied. Implementation of policies should be monitored and there should be evidence of a rolling programme of audit and a date for revision stated.	2008 (Amended 2015) Criterion 9 th
3	Is infection prevention and control included in all staff induction programmes?	M The Health and Social Care Act 2008 (Amended 2015) Criterion 6, section 6.2 states that 'Infection prevention would need to be included in the job descriptions and be included in the induction programme and staff updates of all employees (including volunteers)'.	Moderate
4	Does the practice have evidence to show that all clinical and non-clinical staff (including cleaning staff) are up to date with infection prevention and control training specific to their roles?	M Criterion 6 of The Health and Social Care Act 2008 (Amended 2015) requires providers to have systems to ensure that all care workers (including contractors and volunteers) are aware of and discharge their responsibilities in the process of preventing and controlling infection.	Moderate Criterion 6 of The Health and Social (Amended 2015) requires providers to to ensure that all care workers (includ and volunteers) are aware of and di responsibilities in the process of pro controlling infection.

		М	
5	Is there a process for internally recording/reporting untoward incidents in relation to infection prevention and control (e.g. sharps and body fluid splashes)?	 Under 'Health and Safety (Sharp Instruments in Healthcare) Regulations 2013 Guidance for employers and employees' Recording and investigating the incident – regulation 7(1) Employers must make a record of the sharps injury when they are notified of it, whoever provides that notification. They must investigate the circumstances and causes of the incident and take any action required. The injured person is required to provide sufficient information to their employer to allow them to carry out this investigation. The record of the injury should include who was injured, and when and where the incident occurred. If possible, the summary record should contain sufficient detail to identify what type of sharp was involved, at what stage of a procedure or post-procedure/disposal of the sharp the injury occurred, and the severity of the injury. If the employer has an existing accident book or other recording system, it will be appropriate to use this for the record of sharps injuries. Under the EH 40/2005 Workplace Exposure Limit, it is required that those responsible for controlling exposure to hazardous substances at work are not breaching the Health and Safety at Work Act 1974. 	
6	Does the practice have a recorded process in place that includes access to:		
6.1	Local IPC advice and support as needed.	EQR & E	Low
6.2	Local Hospital Consultant Microbiologists.	EQR & E	Low
6.3	Public Health England Health Protection teams	EQR & E	Low
6.4	Local anti-microbial Pharmacy Lead	EQR & E	Low
7	Does the practice have documentary evidence of infection prevention and control audits undertaken, evaluated and with actions taken to improve practice standards?	M Criterion 1 of The Health & Social Care Act 2018 (Amended 2015) requires that providers have 'Systems to manage and monitor the prevention and control of infection. These systems use risk assessments and consider the susceptibility of service users and any risks that their environment and other users may pose to them'.	Moderate
8	Has the Practice carried out a risk assessment for Legionella under the Health & Safety Executives "Legionella' disease – the control of Legionella bacteria in water systems: Approved code of practice & Guidance" (also known as L8)	M Legionella disease – The control of legionella in water system: Approved Code of Practice (ACOP): A suitable and sufficient assessment must be carried out to identify and assess the risk of exposure to legionella bacteria from work activities and water systems on the premises and any precautionary measures needed. The duty-holder is responsible for ensuring the risk assessment is carried out.	A detailed risk assessment will inform any areas of non-compliance such a dirty water tanks and any remedial ac to minimise the risk of legionella prolifi groups of patients such as immunoc
9	Does the practice have a written scheme for prevention of Legionella contamination in water pipes and other water lines?	M Legionella disease – The control of legionella in water system: Approved Code of Practice (ACOP): A suitable and sufficient assessment must be carried out to identify and assess the risk of exposure to legionella bacteria from work activities and water systems on the premises and any precautionary measures needed. The duty-holder is responsible for ensuring the risk assessment is carried out.	A detailed risk assessment will inform any areas of non-compliance such a dirty water tanks and any remedial ac to minimise the risk of legionella prolifi groups of patients such as immunoc

Section 2: The Management of Infection Prevention and Control (Staff Health)

Standard: Infection prevention and control is managed effectively and complies with the Health and Social Care Act 2008: Code of practice on the prevention and control of infection and related guidance (July 2015)

1	1	I			
		M= Mandatory			
#	Questions	EQR= EQR	Risl		
		E= Educational			
		м			
		Criterion 9 of The Health & Social Care Act 2008 (Updated 2015) requires registered provider to:			
	Have all staff at risk been immunised	Have and adhere to policies, designed for the individual's care and provider organisations that will help to prevent and control infections.			
	against hepatitis B and have they had their response to vaccination confirmed by serology for anti HBs antibodies? It is	f. Section F: Prevention of occupational exposure to blood-borne viruses (BBVs) including prevention of sharps injuries	I		
1	recommended that practices keep a copy	Measures to avoid exposure to BBV's (hepatitis B and C and HIV) should include:	Low level of immunity o		
	of the hepatitis B levels. {At risk staff are those who may have direct contact with patient's blood or blood stained body	 immunisation against hepatitis B, as set out in <i>Immunisation against infectious</i> disease, better known as 'The Green Book' (published by Public Health England); 	may place HCWs at risk in cases of ino		
	fluids (including cleaning staff)}	 the wearing of gloves and other protective clothing; 			
		 the safe handling and disposal of sharps, including the provision of medical devices that incorporate sharps protection where there are clear indications that they will provide safe systems of working for staff; and 			
		 measures to reduce risks during surgical procedures 			
2	Are all staff routinely advised regarding immunisation against seasonal influenza?	EQR & E	Mc Some groups of patient: infection which may be wc		
		М			
		Under the Health and Safety Act 1974, the Occupational Health are required to provide advice on health issues relevant to the working environment.			
		The Health & Social Care Act 2008			
		C Criterion 10, Section 10.3 Occupational health services in respect of BBVs should include:			
3	Does the practice have access to Occupational Health service or access to appropriate occupational health advice? (This may include pre-employment checks to ensure appropriate immunisations have been given.)	 having arrangements for identifying and managing healthcare staff infected with hepatitis B or C or HIV and advising about fitness for work and monitoring as necessary, in line with Department of Health guidance; 	Ma		
		 liaising with the UK Advisory Panel for Healthcare Workers Infected with Blood-borne Viruses when advice is needed on procedures that may be carried out by BBV-infected care workers, or when advice on patient tracing, notification and offer of BBV testing may be needed; 			
		 a risk assessment and appropriate referral after accidental occupational exposure to blood and body fluids; and 			
		 management of occupational exposure to infection, which may include provision for emergency and out-of-hours treatment, possibly in conjunction with accident and emergency services and on-call infection prevention and control specialists. 			

		М	
		Criterion 9 of The Health & Social Care Act 2008 (Updated 2015) requires registered provider to:	
		Have and adhere to policies, designed for the individual's care and provider organisations that will help to prevent and control infections.	
	Has the issue of immunity to Measles,	f. Section F: Prevention of occupational exposure to blood-borne viruses (BBVs) including prevention of sharps injuries	Ma
4	Rubella and Varicella in clinical staff been considered in the practice and a	Measures to avoid exposure to BBV's (hepatitis B and C and HIV) should include:	Some groups of patient: Rubella and Varicelli
	risk assessment undertaken?	• immunisation against hepatitis B, as set out in Immunisation against infectious	transmitted by I
		<i>disease</i> , better known as 'The Green Book' (published by Public Health England);	
		 the wearing of gloves and other protective clothing; 	
		 the safe handling and disposal of sharps, including the provision of medical devices that incorporate sharps protection where there are clear indications that they will provide safe systems of working for staff; and 	
		measures to reduce risks during surgical procedures	

Section 3: Environment

Standard: The environment is designed and managed to minimise reservoirs for microorganisms and reduce the risk of cross-infection to patients, staff and visitors.

M= Mandatory						
#	Questions	EQR= EQR	Risk Level	Rem		
		E= Educational				
		M				
1	Are all areas including clinical areas and equipment visibly clean and free from extraneous items?	Criterion 2 of The Health and Social Care Act 2008 (Amended 2015) requires all parts of the premises from which it provides care are suitable for the purpose, kept clean and maintained in good physical repair and condition.	Moderate – High This depends on whether it is a consulting room or treatment room where invasive procedures such as coil fitting is undertaken.	All areas in items. Clut ensure surfaces is		
		Μ				
2	Are there comprehensive written specifications for cleaning the environment and equipment in the practice?	Criterion 2 of The Health and Social Care Act 2008 (Amended 2015) requires providers to maintain a clean and safe environment to maintain patient and staff safety.	Moderate	All health ca		
		м				
3	Are there up to date cleaning schedules which includes regular cleaning of clinical, admin and sanitary areas (e.g. toilets, fans, air conditioners, high areas, curtains, blinds, toys, computer keyboards, telephones and desks)?	Criterion 2 of The Health and Social Care Act 2008 (Amended 2015) requires providers to maintain a clean and safe environment to maintain patient and staff safety.	Moderate	Up to date and sanitar toys, co		
				Wall surf		
4	Are walls in all areas in good condition (no cracked or peeling paintwork), intact and have smooth easy-to-clean surfaces?	EQR	Moderate – High This depends on the condition of the walls. For example walls in a clinical room affected by damp and mould will pose a higher risk of cross infection if the room is being used for certain invasive procedures.	manipie je		
		EQR				
5	Is flooring in all areas that are accessible to patients (including corridors, staircase leading to consulting rooms, consulting rooms in a good state of repair and easy-to-clean? (Carpets are not recommended)	Where carpet flooring is available in consulting rooms, these should be in a good state of repairs and steam cleaned on a regular basis. There should be a plan in place for carpet flooring in all patient associated areas to be replaced at the next practice refurbishment with flooring which are impervious to fluids, seamless and without joints and the edges with the walls should be either sealed or continuous to the walls with a coved edge.	Moderate – High This depends on whether invasive procedures such as coil fitting is being undertaken in consulting rooms.	Carpets patients c these sho basis. Ther areas to b imperviou shoul		
		Flooring for clinical/ treatment rooms is covered in Section 11 of the audit tool.				
	Are furniture (e.g. chairs, couches, pillows etc) in	EQR	Moderate	Funniture.		
6	clinical areas and other areas accessible to patients impermeable / washable / suitable for its use?	Furniture such as seating for patient in clinical rooms / waiting areas, should be impervious and easy to clean as well as compatible with detergents and disinfectants.	HTM 03-01 section 10.21 requires impervious surfaces for easy clean and to avoid the build- up of dust.	Furniture s imperv		

7	Are cleaning equipment and materials for cleaning colour coded, suitable for use and stored appropriately?	EQR	Moderate	Mops and b & Yellow) tr define To ensure t used. Fur and ensu multiplica heads t
8	Is the area for storing cleaning equipment well ventilated, clean and tidy (no clutter) and is it of an adequate size?	EQR	Moderate Mops and cleaning equipment need to be protected from risk of contamination. A well ventilated cleaning storage area will allow for mop heads to dry.	To ensure

Section 4: Hand Hygiene

Standard: The practice has a clear mechanism to ensure effective implementation of hand hygiene procedures are in place and hand hygiene is practiced at all times to reduce the potential for cross infection between staff, patients, the environment and equipment.

		M= Mandatory		
#	Questions	EQR= EQR	Risk Level	
		E= Educational		
		М		
	Does the practice has a Hand Hygiene Policy?	Criterion 9 of The Health & Social Care Act 2008 (Updated 2015) requires registered provider to:		
		Have and adhere to policies, designed for the individual's care and provider organisations that will help to prevent and control infections.		
		a. A. Standard infection prevention and control precautions		
		Preventing infections reduces the overall need to use antimicrobials and helps to reduce selection pressure for the development of antimicrobial resistance.		
1		 Policy should be based on evidence-based guidelines, including those on hand hygiene at the point of care and the use of personal protective equipment; 	Moderate	
		 Policy should be easily accessible and be understood by all groups of staff, service users and the public. 		
		Compliance with the policy should be audited		
		 Provisions on regular refresher training, support for patients to clean their hands, and products for staff with occupational dermatitis are among the issues that should be covered in the hand hygiene policy. 		
		The NHS Outcome Framework enforces the 'treating and caring for people in a safe environment and protecting them from avoidable harm'		
	Is the hand hygiene technique displayed as a laminated		Low	
2	poster adjacent to the hand washbasin or is it featured on the soap dispenser?	EQR	This serves as a reminder for staff t the correct techniques.	
		EQR	Moderate	
3	Does your practice policy demonstrate an awareness of the DH uniform policy particularly in treatment rooms and minor surgery room? (E.g. bare below the elbows).		Clinical staff should be free fror jewellery, wrist watches, long na artificial nails and nail varnish, as t will prevent adequate hand hygier clinical environment.	
4	Are there wash basins dedicated to hand hygiene in each clinical and consulting room which can be easily		Moderate	
4	accessed?	EQR	To comply with HBN 00-09 requirem	
5	Do all hand wash basins for use in connection with	EQR	Moderate	
J	clinical procedures have elbow or wrist operated mixer taps?		To comply with HBN 00-09 requirem	
6	Is the hot water thermostatically controlled?	EQR	Moderate To comply with HBN 00-09 requirem	
	An town at all aligned been described to the first form		Moderate	
7	Are taps at all clinical hand wash basins free from swan neck type taps?	EQR	To comply with HBN 00-09 requirem	
8	Are all hand wash basins free from plugs?	EQR	Moderate To comply with HBN 00-09 requirem	

9	Are all hand wash basins in clinical and consulting rooms free an overflow?	EQR	Moderate To comply with HBN 00-09 requirem
10	Are hand hygiene facilities clean and free from clutter (check wash basins, taps, splash-backs, soap and paper- towel dispensers)?	EQR	Moderate To prevent contamination and to ad to Criterion 2 of The Health & Soc Care Act 2008.
11	Are hand hygiene facilities free from damage?	EQR	Low To prevent the risks of multiplicatio micro-organisms in grooves/ dama surfaces which are not easily clear
12	Is the tap off-set from the waste outlet?	EQR	Moderate This is a requirement for clinical ha washing sinks as detailed on page the HBN 00-09.
13	Is liquid soap dispensed from single use cartridges or bottles? (I.e. no bar soap or refillable containers)?	EQR	Low Liquid soap dispensers should be v mounted at all wash-hand basins ar designed to be operated withou contamination from the user's har coming into direct contact with th dispensing mechanism.
14	Is alcohol-based hand rub available for use when required, including use during domiciliary visit?	EQR	Low Guidance from WHO
15	Are paper hand towels available? (I.e. no cloth towels in use).	EQR	Moderate As detailed in HBN 00-09. The use paper towels in rolls should be discouraged; they are difficult to tea without contaminating the remaining
16	Are hand wash basins free from nail brushes?	EQR	Low
17	Are there separate arrangements available to dispose of waste materials (e.g. urine) other than using the hand washbasin?	EQR	Moderate This can be a source of cross contamination.

Section 5: Personal Protective Equipment (PPE)

		M= Mandatory	
#	Questions	EQR= EQR	Risk Level
		E= Educational	
		М	
		Criterion 9 of The Health & Social Care Act 2008 (Updated 2015) requires registered provider to:	
		Have and adhere to policies, designed for the individual's care and provider organisations that will help to prevent and control infections.	
		Section f. Prevention of occupational exposure to blood-borne viruses (BBVs) including prevention of sharps injuries	
1	Does the practice have a policy on the appropriate use of PPE?	M Measures to avoid exposure to BBV's (hepatitis B and C and HIV) should include:	Moderate
		 immunisation against hepatitis B, as set out in Immunisation against infectious disease, better known as 'The Green Book' (published by Public Health England); 	
		 the wearing of gloves and other protective clothing; 	
		 the safe handling and disposal of sharps, including the provision of medical devices that incorporate sharps protection where there are clear indications that they will provide safe systems of working for staff; and 	
		measures to reduce risks during surgical procedures	
2 2.1	Are the following PPE available for staff?		
	Gloves (sterile/non-sterile) are appropriate for use, i.e, latex & latex free nitrile?	EQR	Moderate
	Vinyl gloves are not recommended for clinical activities were blood/body fluid may be anticipated.		Health and Safety Act 1
2.2	Disposable aprons available?	EQR	Moderate
			Health and Safety Act 19
2.3	Disposable face and eye protection (to be worn by staff if splashing of blood, body fluids or chemicals is anticipated)?	EQR	Moderate Infection: Prevention and c of healthcare associated in in primary and community (2012)
			Moderate
3	Are staff aware of the principles of wearing and disposing of PPE? (i.e. disposable gloves, aprons masks and goggles)	EQR	Training of staff in the use of to minimise risks of contamination.

Standard: Protective clothing is available/worn for all aspects of care which may involve contact with blood/body fluids or where asepsis is required

4	Are PPE items worn as single use items?	EQR	Moderate To minimise risks of crc contamination/transmiss
5	Where required are aprons and gloves changed between different episodes of care on the same patient?	EQR	Moderate To minimise risks of crc contamination/transmiss
6	Are gloves removed and hand hygiene performed after every clinical activity?	EQR	Moderate Compliance with the WH moments of Hand Hygie
7	Are staff aware on the decontamination process required for re- usable goggles (if available)?	EQR	Low

Section 6: Prevention and management of spillages of blood & high risk body fluids

Standard: Equipment appropriate for cleaning blood or other body fluid is available specifically for dealing with such incidents safely.

#	Questions	M= Mandatory EQR= EQR E= Educational	Risk Level
1	Does the practice have a policy for managing spillages in healthcare premises?	EQR	Moderate COSHH regulation/Health and Safety
2	Are all staff aware of the procedure for dealing with spillages of blood or other body fluids?	EQR	Moderate COSHH regulation/Health and Safety
3	Are spillage kits available for dealing with spillages of blood/body fluids, i.e, separate kits for dealing with blood spillages and a separate kit for dealing with urine/vomit spillages?	M COSHH regulation/Health and Safety Act 1974 requirements.	Moderate COSHH regulation/Health and Safety
4	Are disposable cloths or mop heads available for cleaning blood or other body fluid spillages?	EQR	Moderate COSHH regulation/Health and Safety

Section 7: Safe handling and disposal of sharps

Standard: Sharps are managed safely to reduce the risk of inoculation injury.

		M= Mandatory	
#	Questions	EQR= EQR	Risk Level
		E= Educational	
		M	
		Criterion 9 of The Health & Social Care Act 2008 (Updated 2015) requires registered provider to:	
		Have and adhere to policies, designed for the individual's care and provider organisations that will help to prevent and control infections.	
		Safe handling and disposal of sharps	
	Doop the practice have a policy on	Relevant considerations include:	
1	Does the practice have a policy on safe handling & disposal of sharps?	 risk management and training in the management of mucous membrane exposure and sharps injuries and incidents; 	Moderate
		 provision of medical devices that incorporate sharps protection mechanisms where there are clear indications that they will provide safe systems of working for staff; 	
		 a policy that is easily accessible and understood by all groups of staff; 	
		 safe use, secure storage and disposal of sharps; and 	
		 auditing of policy compliance 	
		M	
		Criterion 9 of The Health & Social Care Act 2008 (Updated 2015) requires registered provider to:	
		Have and adhere to policies, designed for the individual's care and provider organisations that will help to prevent and control infections.	
	Are sharps containers that conform to	Safe handling and disposal of sharps	
	BS 7320 and UN3291 available in	Relevant considerations include:	
2	every clinical/consulting area and are they positioned safely; out of reach of vulnerable people?	 risk management and training in the management of mucous membrane exposure and sharps injuries and incidents; 	Moderate - High
		 provision of medical devices that incorporate sharps protection mechanisms where there are clear indications that they will provide safe systems of working for staff; 	
		 a policy that is easily accessible and understood by all groups of staff; 	
		 safe use, secure storage and disposal of sharps; and 	
		auditing of policy compliance	
3	Are sharps containers discarded when two thirds full and stored in a secure facility away from public access until collected for disposal?	EQR	Moderate
4	Is blood sampling undertaken by using a 'Sharp Safe' single-use vacuum blood collection system?	EQR	Moderate
5	Is Aseptic Non-Touch Technique (ANTT) used when performing venepuncture?	EQR	Moderate
6	Are sharps used for taking blood from patients at home/care home disposed of in to an appropriate sharps container which is returned to the surgery for safe disposal?	EQR	Moderate

7	Is there evidence that the practice has undertaken a review of sharps management within the practice and employed 'safer sharps' techniques where applicable?	EQR	Moderate Health and Safety (Sharp Instr Healthcare) Regulations Guidance for employers and e Use safer sharps (incorporating mechanisms) – regulation 5 The employer must substitute t unprotected medical sharps wii sharp' where it reasonably practi so. The term 'safer sharp' mear sharps that incorporate featu mechanisms to prevent or minimis accidental injury. For example, syringes and needles are now ava shield or cover that slides or pivo the needle after use.
			Moderate
8	Are sharps containers assembled according to manufacturer's instructions and labelled in accordance with legal requirements?	M Criterion 9 of The Health & Social Care Act 2008 (Updated 2015) requires registered provider to: Have and adhere to policies, designed for the individual's care and provider organisations that will help to prevent and control infections. Safe handling and disposal of sharps	As detailed in HTM 07-0 Each container must be labe accordance with the details of the legal requirer transporting and packaging the v
		Council Directive 2010/32/EU (2010)	container should be tagged or la manner that identifies the inc producer).
9	Are staff encouraged to wear gloves when undertaking venepuncture?	EQR & E	Moderate
10	Are staff aware of the correct procedure to follow after a needle stick injury, other sharps or blood splash exposure?	M Criterion 9 of The Health & Social Care Act 2008 (Updated 2015) requires registered provider to: Have and adhere to policies, designed for the individual's care and provider organisations that will help to prevent and control infections. Section g. Management of occupational exposure to BBVs and post-exposure prophylaxis Management should ensure: • that any member of staff who has a significant occupational exposure to blood or body fluids is aware of the immediate action required and is referred appropriately for further management and follow-up; • provision of clear information for staff about reporting potential occupational exposure – in particular the need for prompt action following a known or potential exposure to HIV or hepatitis B; and • arrangements for post-exposure prophylaxis for hepatitis B and HIV	Moderate
11	Are posters available which show staff the emergency algorithm to follow in case of a sharp injury and is it up to date?	EQR	Moderate

Section 8: Waste Management Policy and Procedures

Standard: Waste is managed safely and in accordance with legislation to minimise the risk of infection or injury to patients, staff and the public.

		M= Mandatory
#	Questions	EQR= EQR
		E= Educational
		м
		Criterion 9 of The Health & Social Care Act 2008 (Updated 2015) requires registered provider to:
		Have and adhere to policies, designed for the individual's care and provider organisations that will help to preve nt and control infections.
		Section P: Safe handling and disposal of waste
		e The risks from waste disposal should be properly controlled. In practice, in relation to waste, this involves:
		assessing risk;
		developing appropriate policies;
		 putting arrangements in place to manage risks;
		 monitoring, auditing and reviewing the way in which arrangements work; and
		being aware of statutory requirements and; legislative change and managing compliance
		Precautions in connection with handling waste should include:
		 training and information (including definition and classification of waste);
		personal hygiene;
1	Does the practice have a policy on waste	 segregation and storage of waste;
	management?	the use of appropriate personal protective equipment;
		• immunisation;
		 appropriate procedures for handling such waste;
		 appropriate packaging and labelling;
		suitable transport on-site and off-site; clear procedures for dealing with accidents, incidents and spillages; and
		appropriate treatment and disposal of such waste
		Systems should be in place to ensure that the risks to service users from exposure to infections caused by waste present in the environment are properly managed, and that duties under environmental law are discharged. The most important of these are:
		 duty of care in the management of waste;
		 duty to control polluting emissions to the air;
		 duty to control discharges to sewers;
		 obligations of waste managers;
		collection of data and obligations to complete and retain documentation including record keeping; and
		requirement to provide contingency plans and have emergency procedures in place
	Is there documentary evidence to show that all	M
2	clinical waste (including sharps containers) is	HTM 07-01: Safe Management of Healthcare Wastes.
-	disposed of by a registered waste collection company?	Waste can only be handed to such authorised persons as registered carriers, permit/ licence holders or someone who is exempt frc either being a registered carrier or operating under a permit/ licence.
	Are records of waste transfer and disposal	
3	arrangements kept and stored in accordance with the EPA 1990?	EQR
4	Are there easily accessible and compliant foot- operated &fully enclosed clinical waste bins, with the appropriate colour coded bag (yellow or orange) available, in each clinical area? (E.g. the foot operation is in working order)?	EQR
5	Is clinical and domestic waste correctly segregated (clinical waste in yellow or orange bags, according to waste regulations and domestic waste in black bags)?	EQR

6	Are clinical waste bags marked with the practice code when securing for disposal?	EQR
7	Are waste bags less than 2/3 full and securely tied?	EQR
8	Where clinical waste is not collected directly from clinical areas, is it stored in a separate, secure area for waste which is kept clean and tidy and secure from vermin and/or other inappropriate/extraneous items?	EQR
9	Are staff encouraged to report all incidents (including near misses) to the designated infection control lead at the practice?	EQR

Section 9: Management of Specimens

Standard: All specimens will be collected packaged and transported safely in approved containers in line with recognised standards – Packaging Instruction 650 and 621 and requirements of UN3373 or UN3291 to minimise the risk of cross infection.

		M= Mandatory	
#	Questions	EQR= EQR	
		E= Educational	
		М	
		Criterion 9 of The Health & Social Care Act 2008 (Updated 2015) requires registered provider to:	
		Have and adhere to policies, designed for the individual's care and provider organisations that will help to prevent and control infections.	
1	Does the practice have a policy or procedure for specimen handling?	<i>q</i> .	
		P Section Q: Packaging, handling and delivery or laboratory specimens	
		Biological samples, cultures and other materials should be transported in a manner that ensures that they do not leak in transit and are compliant with current legislation. Staff who handle samples must be aware of the need to correctly identify, label and store samples prior to forwarding to laboratories. In addition, they must be aware of the procedures needed when the container or packaging becomes soiled with body fluids.	
2	Where applicable are specimens stored in a dedicated refrigerator (not with food, vaccines or medicines)?	EQR	
3	Are arrangements for specimen testing appropriate in consulting rooms?	EQR	
4	Are staff aware of the appropriate way to handle and transport specimens?	EQR & E	

Section 10: Decontamination of medical devices

Standard: All medical devices are decontaminated in a safe and appropriate manner to minimise the risk of infection and cross-infection.

Note: Medical devices include not only surgical instruments but a wide variety of other equipment such as dressing trolleys, BP cuffs and baby scales. A risk assessment needs to be carried out on each medical device to ensure that the appropriate level of decontamination is carried out. For those in the high or medium risk categories cleaning and sterilisation must be carried out (e.g. autoclaving). For those in the lowest risk category cleaning or cleaning plus disinfection are needed depending on circumstances

		M= Mandatory	
#	Questions	EQR= EQR	Risk Lev
		E= Educational	
1	Does the practice have a policy which outlines the decontamination processes the GP Practices use for all medical devices?	M Criterion 9 of The Health & Social Care Act 2008 (Updated 2015) requires registered provider to: Have and adhere to policies, designed for the individual's care and provider organisations that will help to prevent and control infections. <i>j. Decontamination of reusable medical devices</i> Decontamination involves a combination of processes and includes cleaning, disinfection and sterilisation, according to the intended use of the device. This aims to render a reusable item safe for further use on service users and for handling by staff; Effective decontamination of reusable medical devices is an essential part of infection risk control and is of special importance when the device comes into contact with service users or their body fluids. There should be a system to protect service users and staff that minimises the risk of transmission of infection from medical devices. This requires that the device or instrument set can be clearly linked in a traceable fashion to the individual process cycle that was used to decontaminate it, such that the success of that cycle in rendering the device safe for reuse can be verified; Reusable medical devices should be decontaminated in accordance with manufacturers' instructions and current national or local best practice guidance. This must ensure that the device complies with the 'Essential Requirements' provided in the Medical Devices Regulations 2002 where applicable. This requires that the device should be clean and, where appropriate, sterilised at the end of the decontamination process and maintained in a clinically satisfactory condition up to the point of use; Reusable medical devices employed in invasive procedures, for example, endoscopes and surgical instruments have to be either individually identifiable or identified to a set of which they are a consistent member, throughout the use and decontamination cycle in order to ensure subsequent traceability; Systems should also be implemented to enable the identification of service users on whom the medical devices have	Moderate
2	Does the practice use single use surgical instruments?	EQR	Moderate
3	Does the practice use an accredited external sterile supply service for re- usable surgical instruments and devices that need to be sterile at the point of use?	EQR	Moderate - F
4	Are all medical devices stored appropriately and above floor level to avoid contamination?	EQR	Moderate

			High
			All sterile instruments should t patient safe
5	Are all items of sterile equipment within their use-by date?	EQR	Single use instruments shu immediately after use. In the instrument packs are damage immediately discarded, beca their sterile prop
			A strict protocol should be ma sterile instruments and equipi date. Those that have exceec should be disposed of
	Are all items of equipment that come into	M	
6	contact with patients cleaned or decontaminated according to guidelines or are disposed of after each use?	Criterion 9 of The Health & Social Care Act 2008 (Updated 2015) requires registered provider to:	Moderate – F This depends on the re-usab shared, the decontamination procedure that these instrum
		Have and adhere to policies, designed for the individual's care and provider organisations that will help to prevent and control infections. <i>j. Decontamination of reusable medical devices</i>	for.
7	Is there a cleaning schedule/check list maintained for all items requiring cleaning?	EQR	Moderate

Section 11: Clinical Rooms

Standard: The environment is designed and managed to minimise reservoirs for micro-organisms and reduce the risk of cross infection to patients, staff and visitors.

		M= Mandatory	
	Questiens	EQR= EQR	Risk Lev
#	Questions		NISK LEW
		E= Educational	
		М	
1	Are all clinical rooms and all work surfaces clean and free from extraneous items?	Criterion 2 of The Health and Social Care Act 2008 (Amended 2015) requires all parts of the premises from which it provides care are suitable for the purpose, kept clean and maintained in good physical repair and condition.	Moderate - H
2	ls flooring impervious to liquids, non-slip, intact and clean?	EQR	Moderate
3	Does the flooring form a coved skirting (i.e. uplifted at the edges on to the walls) OR is the gap between the floor and the skirting sealed and is the seal maintained?	EQR	Moderate
4	Are walls and ceilings clean, dry and free from visible defects (no cracks, peeling paintwork) and have smooth easy to clean surfaces?	EQR	Moderate – H Walls in clinical areas should cracks or joints and easy to textured wall paper in clinical r be removed and the walls ma room affected by damp and higher risk of cross infection i used for certain invasive
5	Is there an examination couch with an intact, impervious cover and single use roller paper available for use?	EQR	Moderate
6	Is the examination couch fitted with a paper roll holder?	E	Low
7	Are there sufficient work surfaces and dressing trolleys of smooth, impervious and cleanable material?	EQR	Low
8	Are all treatment surfaces in the room cleaned every working day with hot water and detergent or detergent wipes in accordance with written practice cleaning schedules?	EQR	Moderate

Section 12: Vaccine Storage and Cold Chain

Standard: Vaccines a	ire stored and trar	sported safely.
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		M= Mandatory		
#	Questions	EQR= EQR	Risk Level	Ren
		E= Educational		
1	Does the practice have an up to date cold chain policy (reviewed within the last two years) as per the standards of the Green Book and is this accessible to all staff?	M Criterion 9 of The Health & Social Care Act 2008 (Updated 2015) requires registered provider to: Have and adhere to policies, designed for the individual's care and provider organisations that will help to prevent and control infections.	High	To ensu years) shou
2	Is there a designated person/s (at least two recommended) in the practice responsible for the ordering, delivery and storage of vaccines?	EQR	Moderate As detailed in Chapter 3 of The Green Book.	To ensur
3	Are vaccines monitored for their expiry dates and the close to expiry stocks clearly labelled?	EQR	Moderate As detailed in Chapter 3 of The Green Book.	To ensur
4	Is the refrigerator specialised for the storage of vaccines (eg. the refrigerator has wire shelves/baskets or shelves capable of allowing air ventilation, there are no vaccines stored in enclosed plastic trays at bottom of refrigerator, domestic type refrigerators are not recommended)?		Moderate As detailed in Chapter 3 of The Green Book.	To ensu (DH,
5	Are vaccines correctly stored to allow good air flow within the vaccine refrigerator? (eg. vaccines are not stored against the back plates, touching the side of the fridge, at bottom of fridge or in vegetable bins and not stored in containers that are not webbed baskets)?	EQR	Moderate As detailed in Chapter 3 of The Green Book.	The air f Vaccine Good a cooling f the side plastic tra
6	Are there measures in place to prevent the fridge from being turned off (switch-less socket or warning label on plug)?	EQR	Moderate As detailed in Chapter 3 of The Green Book.	The vaco sup
7	Is/Are the vaccine fridge/s located in a well-ventilated area. (eg. not located near any heat source, ie radiator, or direct sunlight)?	EQR	Moderate As detailed in Chapter 3 of The Green Book.	To ensi
8	Is the temperature of the vaccine fridge continually monitored with a min/max thermometer and the temperatures are recorded each working day to ensure vaccines are maintained at 2-8 ⁰ C? (Min, max and actual fridge temperatures are recorded)? (<i>It is best practice to record the temperatures twice daily</i>)	M As detailed in Chapter 3 of The Green Book.	Moderate As detailed in Chapter 3 of The Green Book.	Tempera Vaccine t of the the To ensure To ensur
9	Does the practice has a maintenance contract that allows for at least yearly servicing, calibration of the temperature gauge?	EQR	Moderate As detailed in Chapter 3 of The Green Book.	To ensure calibratio
10	Is a second min/max thermometer or Data Logger temperature recording device, independent of mains electricity supply available and used?	EQR	Moderate As detailed in Chapter 3 of The Green Book.	A secon
11	Is the fridge either self-defrosting or is it defrosted monthly or sooner if needed and a validated cool box is then used to maintain the cold chain?	EQR	Moderate As detailed in Chapter 3 of The Green Book.	To ensu

12	Is there a process in place for safe disposal of expired, damaged or surplus vaccines?	EQR	Moderate It is a requirement of the HTM 07-01: Safe management of Healthcare waste.	To ensure
13	Does the practice have records of vaccines received, batch numbers, expiry dates, fridge temperatures, servicing and defrosting of the fridge?	EQR	Moderate As detailed in Chapter 3 of The Green Book.	To ensu fridç
14	Is there accessible written guidance on what staff should do in the event of a power cut or a temperature reading outside the required range?	EQR	Moderate As detailed in Chapter 3 of The Green Book.	To ensı event o

Section 13: Notification of infectious diseases and contamination

Standard: All notifiable diseases are reported on suspicion, within the time frames set out in the Health Protection (Notification) Regulations 2010

		M= Mandatory	
#	Questions	EQR= EQR	Risk Level
		E= Educational	
1	Does the practice have a policy on managing patients with communicable diseases?	M Criterion 9 of The Health & Social Care Act 2008 (Updated 2015) requires registered provider to: Have and adhere to policies, designed for the individual's care and provider organisations that will help to prevent and control infections. Section d. Isolation of service users with an infection (see also criterion 7) • The isolation policy should be evidence based and reflect local risk assessment; ¹ • Indications for isolation should be included in the policy, as should procedures for the infection prevention and control management of service users in isolation; • Information on isolation should be easily accessible and understood by	Moderate
2	Does the practice notify all reportable infectious disease on suspicion to the proper officer at the local authority?	all groups of staff, service users and the public M Criterion 9 of The Health & Social Care Act 2008 (Updated 2015) requires registered provider to: Have and adhere to policies, designed for the individual's care and provider organisations that will help to prevent and control infections. Section d. Isolation of service users with an infection (see also criterion 7).	Moderate
3	Does the practice have access to notification forms?	EQR	Moderate
4	Does the practice notifying gastro intestinal disease (food poisoning) on suspicion?	EQR	Moderate
5	Does the practice notify Gastro intestinal disease (food poisoning) when stool specimen results are received from the microbiology laboratory?	EQR	Moderate
6	Is the practice aware of the new requirements to notify cases of contamination and other diseases which may have public health significance that are not listed in the regulations?	EQR	Moderate

Section 14: Antimicrobial Stewardship (AMS)

Standard: Prescribers are aware of the relevant guidelines and regularly audit their own, and discuss in practice meetings, their antibiotic prescribing patterns. GPs are aware of TARGET: Treat Antibiotics Responsibly. Guidance, Education, Tool.

		M= Mandatory	
#	Questions	EQR= EQR	Risk Level
		E= Educational	
		EQR	
		H&SC Act 2008 – Criterion 3	
1	Are GP prescribers in the practice aware of the	Ensure appropriate antimicrobial use to optimise patient outcomes and to reduce the risk of adverse events and	Madaata
1	TARGET toolkit?	antimicrobial resistance. Antimicrobial prescribing should follow local policies and national guidance such as PHE primary care guidance: Managing common infections; guidance for primary care and TARGET. Evidence to demonstrate adoption and adherence to policies and guidelines should be available to commissioners.	Moderate
	Have all GP prescribers completed the Antimicrobial		
	Stewardship Self-Assessment Checklist available in TARGET?		
2	(Give number of GP prescribers using the self-	EQR	Moderate
	assessment checklist against those who do not)		
	/		
	Is the document "Antimicrobial prescribing and stewardship competencies" available and/or has it		
3	been read by prescribers in the practice? (Give number of prescribers who are aware of this document against	EQR	Low
	those who are not)		
	1		
4	Are all the prescribers in the Practice aware of the Public Health England AMR local indicators?	EQR	Moderate
5	Is the practice aware of how they may access their antibiotic prescribing data online?	EQR	Low
6	Are all prescribers in the practice aware of the NICE guidelines on AMS.	EQR	Low
7	Do all prescribers give information to their service users of how they should correctly use antimicrobial medicines and the dangers associated with their overuse and misuse?	EQR	Low
8	Are all prescribers aware of the UK's 5-year Antimicrobial Resistance Strategy?	E	Low
9	Does the practice actively participates in the European Antibiotic Awareness Day/Week (EEAD) held in November each year?	E	Low
10	Are all clinical staff in the Practice aware of the PHE Antibiotic Guardian campaign?	E	Low
11	Does the practice have an identified sepsis lead / link?	E	Moderate
12	Are clinicians/GPs aware of and/or have received training in identifying sepsis?	EQR	Moderate
13	Does the practice promote the use of the GRASP Fever Audit tool?	E	Moderate
14	Does the practice make use of the National Early Warning Signs (NEWS) tool to assess adults for sepsis and the Paediatric Early Warning Signs (PEWS) tool to assess children?	EQR	Moderate

Section 15: Minor Surgery rooms

Standard: The environment is designed and managed to minimise reservoirs for micro-organisms and reduce the risk of cross infection to patients, staff and visitors

		M= Mandatory	
#	Questions	EQR= EQR	
		E= Educational	
1	Is there is a dedicated room for minor surgery (for practices performing level 3 minor surgery) or is the clinical room of sufficient standard for undertaking level 2 minor surgery?	H&SC Act 2008 – All parts of the premises from which it provides care are suitable for the purpose,	The risk minor st
		M for Level 2 & 3 minor surgery	
		Criterion 9 of The Health & Social Care Act 2008 (Updated 2015) requires registered provider to:	
		Have and adhere to policies, designed for the individual's care and provider organisations that will help to prevent and control infections.	
		j. Decontamination of reusable medical devices	
		 Decontamination involves a combination of processes and includes cleaning, disinfection and sterilisation, according to the intended use of the device. This aims to render a reusable item safe for further use on service users and for handling by staff; 	
2	there adequate storage space?	 Effective decontamination of reusable medical devices is an essential part of infection risk control and is of special importance when the device comes into contact with service users or their body fluids. There should be a system to protect service users and staff that minimises the risk of transmission of infection from medical devices. This requires that the device or instrument set can be clearly linked in a traceable fashion to the individual process cycle that was used to decontaminate it, such that the success of that cycle in rendering the device safe for reuse can be verified; Reusable medical devices should be decontaminated in accordance with manufacturers' instructions and current national or local best practice guidance. This must ensure that the device complies with the 'Essential Requirements' provided in the Medical Devices Regulations 2002 where applicable. This requires that the device should be clean and, where appropriate, sterilised at the end of the decontamination process and maintained in a clinically satisfactory condition up to the point of use; 	
		Management systems should ensure adequate supplies of reusable medical devices, particularly where specific devices are essential to the continuity of care;	
		Reusable medical devices employed in invasive procedures, for example, endoscopes and surgical instruments have to be either individually identifiable or identified to a set of which they are a consistent member, throughout the use and decontamination cycle in order to ensure subsequent traceability;	
		Systems should also be implemented to enable the identification of service users on whom the medical devices have been used;	
		Decontamination of single-patient use devices, i.e. that equipment designated for use only by one patient, should be subject to local policy and manufacturer's instructions	

		M for Level 2 & 3 minor surgery	
		Criterion 9 of The Health & Social Care Act 2008 (Updated 2015) requires registered provider to:	
		Have and adhere to policies, designed for the individual's care and provider organisations that will help to prevent and control infections.	
		j. Decontamination of reusable medical devices	
	Is the practice using disposable single use instruments, supplied by a recognised manufacturer of sterile disposable instruments? If the answer is 'No' the practice should contact the local IPC advisors for advice.	 Decontamination involves a combination of processes and includes cleaning, disinfection and sterilisation, according to the intended use of the device. This aims to render a reusable item safe for further use on service users and for handling by staff; 	
3		 Effective decontamination of reusable medical devices is an essential part of infection risk control and is of special importance when the device comes into contact with service users or their body fluids. There should be a system to protect service users and staff that minimises the risk of transmission of infection from medical devices. This requires that the device or instrument set can be clearly linked in a traceable fashion to the individual process cycle that was used to decontaminate it, such that the success of that cycle in rendering the device safe for reuse can be verified; Reusable medical devices should be decontaminated in accordance with manufacturers' instructions and current partice users in the requirements. 	
		national or local best practice guidance. This must ensure that the device complies with the 'Essential Requirements' provided in the Medical Devices Regulations 2002 where applicable. This requires that the device should be clean and, where appropriate, sterilised at the end of the decontamination process and maintained in a clinically satisfactory condition up to the point of use;	
		Management systems should ensure adequate supplies of reusable medical devices, particularly where specific devices are essential to the continuity of care;	
		 Reusable medical devices employed in invasive procedures, for example, endoscopes and surgical instruments have to be either individually identifiable or identified to a set of which they are a consistent member, throughout the use and decontamination cycle in order to ensure subsequent traceability; 	
		 Systems should also be implemented to enable the identification of service users on whom the medical devices have been used; 	
		 Decontamination of single-patient use devices, i.e. that equipment designated for use only by one patient, should be subject to local policy and manufacturer's instructions 	
		M for Level 2 & 3 minor surgery	
		Criterion 9 of The Health & Social Care Act 2008 (Updated 2015) requires registered provider to:	
		Have and adhere to policies, designed for the individual's care and provider organisations that will help to prevent and control infections.	
	Where applicable, is an accredited external sterile supply service used for re-usable surgical instrument that need to be sterile at the point of use? If a GP contractor wishes to have advice about in-house decontamination arrangements, they should contact NEL to establish what this would entail, together with the associated	j. Decontamination of reusable medical devices	
		 Decontamination involves a combination of processes and includes cleaning, disinfection and sterilisation, according to the intended use of the device. This aims to render a reusable item safe for further use on service users and for handling by staff; 	
		• Effective decontamination of reusable medical devices is an essential part of infection risk control and is of special importance when the device comes into contact with service users or their body fluids. There should be a system to protect service users and staff that minimises the risk of transmission of infection from medical devices. This requires that the device or instrument set can be clearly linked in a traceable fashion to the individual process cycle that was used to decontaminate it, such that the success of that cycle in rendering the device safe for reuse can be verified;	
4		 Reusable medical devices should be decontaminated in accordance with manufacturers' instructions and current national or local best practice guidance. This must ensure that the device complies with the 'Essential Requirements' provided in the Medical Devices Regulations 2002 where applicable. This requires that the device should be clean and, where appropriate, sterilised at the end of the decontamination process and maintained in a clinically satisfactory condition up to the point of use; 	
	monitoring.	 Management systems should ensure adequate supplies of reusable medical devices, particularly where specific devices are essential to the continuity of care; 	
		 Reusable medical devices employed in invasive procedures, for example, endoscopes and surgical instruments have to be either individually identifiable or identified to a set of which they are a consistent member, throughout the use and decontamination cycle in order to ensure subsequent traceability; 	
		 Systems should also be implemented to enable the identification of service users on whom the medical devices have been used; 	
		Decontamination of single-patient use devices, i.e. that equipment designated for use only by one patient, should be subject to local policy and manufacturer's instructions	
		м	
_	Is the minor surgery room clean and	H&SC Act 2008 –	
5	free from extraneous items?	Criterion All parts of the premises from which it provides care are suitable for the purpose, kept clean and maintained in good physical repair and condition.	

			Ν
6	Are walls in good condition (no cracked, damaged or peeling paintwork or rough surface finishes), intact and have smooth easy-to-clean surfaces?	M for Level 2 & 3 minor surgery H&SC Act 2008 – All parts of the premises from which it provides care are suitable for the purpose, kept clean and maintained in good physical repair and condition.	The risk and cond Evidence be cons the ı
7	Is flooring impermeable, intact and have continuous edging coved up the walls?	M for Level 2 & 3 minor surgery H&SC Act 2008 – All parts of the premises from which it provides care are suitable for the purpose, kept clean and maintained in good physical repair and condition.	
8	Are ceilings intact and free from visible cracks or visible defects?	M for Level 2 & 3 minor surgery H&SC Act 2008 – All parts of the premises from which it provides care are suitable for the purpose, kept clean and maintained in good physical repair and condition.	
9	Are ceiling lights protected / enclosed from potential contamination?	M for Level 2 & 3 minor surgery H&SC Act 2008 – All parts of the premises from which it provides care are suitable for the purpose, kept clean and maintained in good physical repair and condition.	
10	Does the room have adequate ventilation - natural or mechanical (no desktop fans)? For level 3 minor surgery, the room should have mechanical ventilation with a minimum of 10 air exchanges as per current guidelines.	M for Level 2 & 3 minor surgery H&SC Act 2008 – All parts of the premises from which it provides care are suitable for the purpose, kept clean and maintained in good physical repair and condition.	
11	Is the heat source and pipe work in the room enclosed to prevent accumulation of dust and dirt?	M for Level 2 & 3 minor surgery H&SC Act 2008 – All parts of the premises from which it provides care are suitable for the purpose, kept clean and maintained in good physical repair and condition.	
12	Is the treatment couch intact and is protected with disposable paper towel that is changed after each patient?	M for Level 2 & 3 minor surgery H&SC Act 2008 – All parts of the premises from which it provides care are suitable for the purpose, kept clean and maintained in good physical repair and condition.	
13	Are all work surfaces intact, smooth, and impervious easy to clean and are able to withstand cleaning with chemical disinfectants?	M for Level 2 & 3 minor surgery H&SC Act 2008 – All parts of the premises from which it provides care are suitable for the purpose, kept clean and maintained in good physical repair and condition.	
14	Are all wall cabinets intact with doors? (open shelves are not recommended, wall cabinets should continue to ceiling level or have sloped tops).	EQR H&SC Act 2008 – All parts of the premises from which it provides care are suitable for the purpose, kept clean and maintained in good physical repair and condition.	
15	Does the clinical hand wash basin conforms to current recommended guidance (HTM 00-10; HBN 00-09) with the following available facilities: • Elbow, foot operated mixer taps • No swan neck fittings on taps • Thermostatically controlled hot / cold water • Free from sink plugs and overflows • Water from the tap does not drain directly into the drainage outlet • Hand hygiene facilities are intact.	M for Level 2 & 3 minor surgery H&SC Act 2008 – All parts of the premises from which it provides care are suitable for the purpose, kept clean and maintained in good physical repair and condition.	
16	Are there wall mounted dispensers for liquid soap and is the liquid soap dispensed via non-refillable cartridges? (Antiseptic hand wash must also be dispensed via wall mounted dispensers and dispensed from non-refillable cartridges)	M for Level 2 & 3 minor surgery H&SC Act 2008 – All parts of the premises from which it provides care are suitable for the purpose, kept clean and maintained in good physical repair and condition.	

		M for Level 2 & 3 minor surgery
17	Are there wall mounted dispensers with good quality disposable paper hand towels?	H&SC Act 2008 – All parts of the premises from which it provides care are suitable for the purpose, kept clean and maintained in good physical repair and condition.
		As detailed in HBN 00-09. The use of paper towels in rolls should be discouraged; they are difficult to tear off without contaminating the remaining roll.
18	Is the clinical hand wash basin free from re-usable nail brushes?	M for Level 2 & 3 minor surgery
19	Are single use sterile and non-sterile gloves available in latex and non-latex nitrile material?	M for Level 2 & 3 minor surgery
20	Is there is a designated stainless steel trolley available for use in this room only?	M for Level 2 & 3 minor surgery
21	Is there a clean clinical waste bin with a foot pedal with waste bag fully enclosed and is it in good operating condition?	M for Level 2 & 3 minor surgery
	(Waste bags must not be attached to cupboards / trolleys etc).	HTM 07-01: Management of Healthcare Wastes
		M for Level 2 & 3 minor surgery
		Criterion 9 of The Health & Social Care Act 2008 (Updated 2015) requires registered provider to:
		Have and adhere to policies, designed for the individual's care and provider organisations that will help to prevent and control infections.
		Section f. Prevention of occupational exposure to blood-borne viruses (BBVs) including prevention of sharps injuries
22	Do all staff use recommended PPE (including disposable goggles) when splashing of body fluids is anticipated?	M Measures to avoid exposure to BBV's (hepatitis B and C and HIV) should include:
		 immunisation against hepatitis B, as set out in <i>Immunisation against infectious disease</i>, better known as 'The Green Book' (published by Public Health England);
		 the wearing of gloves and other protective clothing;
		 the safe handling and disposal of sharps, including the provision of medical devices that incorporate sharps protection where there are clear indications that they will provide safe systems of working for staff; and
		measures to reduce risks during surgical procedures
23	Are disposable sterile drapes available and used for level 2 and level 3 minor surgeries?	M for Level 2 & 3 minor surgery
24	Does the GP Practice audit post- operative wound infections and are records kept?	M for Level 2 & 3 minor surgery

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