

# Infection Prevention & Control (IPC) Audit for Dental



## Introduction

Good infection prevention and control is essential to ensure that people who use Primary Care services receive safe and effective care. Effective prevention and control of infection must be part of everyday practice and be applied consistently by everyone. The Care Quality Commission (CQC) will assess a registered provider on how it complies with the infection prevention requirements, including environmental cleanliness.

## Purpose

The purpose of this audit is to:

- Ensure an infection prevention and control (IPC) audit is conducted on a six-monthly basis – as a minimum
- Ensure the audit accurately reflects findings, with reference to supportive information as examples of evidence of good practice
- Promote a standardised approach to evidence gathering and supportive documentation when preparing for and undertaking an Infection Prevention & Control Audit
- Ensure responsive action is taken and recorded to demonstrate delivery and progress for all findings recorded as 'No'

## Approach

A dedicated inspection date(s) will need to be identified, with named auditors identified to conduct the audit. On the agreed day(s), the allocated auditor(s) will:

- Ensure the auditor(s) answer the questions honestly and objectively
- In the event of answering 'Yes' to any question, the auditor(s) will need to document and collate examples of their findings in the 'Comments' column and/or refer to examples of supportive evidence where appropriate e.g. meeting minutes and date of meeting

This audit template covers the following areas:

1. Prevention of Bloodborne Virus Exposure
2. Decontamination
3. Environmental Design and Cleaning
4. Hand Hygiene
5. Management of Dental Medical Devices
6. Personal Protective Equipment
7. Waste Management
8. Infection Prevention & Control (IPC) Inspection – Action Plan

## Feedback

Ensure the completed audit tool is reviewed and the findings shared with the Practice Manager and/or Registered Manager.

## Actions

- Draft an action plan in response to all questions answered 'No', to demonstrate improvements, management of risk and delivery of progress
- The action plan must identify:
  - The issue
  - The action(s) taken/to be taken
  - Named individuals responsible for managing and delivering the actions
  - Anticipated delivery timeframes
- Give a copy of the audit findings plus action plan to the Practice Manager and/or Registered Manager
- Ensure the Practice Manager shares the audit findings and actions required with staff, celebrating success, and identifying areas of improvements required

Name of person carrying out the audit	Job role of the person doing the audit	Date Completed	Signature on completion

## Section 1: Prevention of Bloodborne Virus Exposure

1. Does the practice have a policy and procedure in place for the prevention of blood borne virus exposure? Does it include the management of spillages, sharps and inoculation incidents in accordance with national guidance?  
 YES  NO
2. Have all staff received training on the prevention and management of blood borne virus exposure?  
 YES  NO
3. Have all staff at risk from sharps injuries received an occupational health check in relation to risk reduction in blood borne virus transmission and general infection?  
 YES  NO
4. Can decontamination and clinical staff demonstrate current immunisation with the hepatitis B vaccine? Can the Practice Manager provide documented proof?  
 YES  NO
5. Are chlorine releasing agents available for blood/bodily fluid spillages and used as per the manufacturer's instructions? Is a spillage kit readily available?  
 YES  NO
6. Are sharps containers correctly assembled and labelled with the date, location and a signature?  
 YES  NO
7. Can you confirm that all sharps containers are not filled beyond the indicator mark? Are they locked shut when filled to the indicator mark?  
 YES  NO
8. Are full sharps containers stored in a secure place away from public access ready for collection?  
 YES  NO
9. Is there a readily available protocol and flowchart in place providing information on what to do in the event of a sharps injury  
 YES  NO
10. Are inoculation injuries recorded?  
 YES  NO

11. Can you confirm only the clinician using syringes is fully responsible for disposing of them?

YES  NO

12. Are safer sharps syringes used?

YES  NO

13. Are sharps containers available at the point of use and positioned safely? For example, wall mounted near the clinician

YES  NO

## Section 2: Decontamination

1. Does the practice have a policy and procedure in place that includes all aspects of the decontamination process?

YES  NO

2. Have all clinical and sterilisation team members received training on decontamination processes they are expected to perform?

YES  NO

3. Do you keep a record of instruments that are single use and therefore cannot be re-processed?

YES  NO

4. Are all wrapped, sterilised instruments date stamped with the date sterilised, use by date and name and signature of the decontaminator operator?

YES  NO

5. Does the practice have a nominated lead responsible for infection control and decontamination?

YES  NO

6. Can you provide evidence that all decontamination equipment (autoclaves, washer disinfectors etc.) are validated and serviced regularly as per the manufacturer's instructions?

YES  NO

7. Is there a procedure for the transportation of instruments to and from other locations that ensures the segregation of contaminated instruments from clean/sterilised instruments?

YES  NO

8. Is all documentation (including testing, servicing, maintenance and repair records) retained in the practice for at least two years?

YES  NO

9. Are reusable instrument trays decontaminated and sterilised after each use?

YES  NO

10. Are any instruments (used or unused) left on trays at the end of each session sterilised before further use?

YES  NO

11. Are instruments that are not decontaminated immediately kept moist until they are decontaminated?

YES  NO

12. Are instruments inspected under illuminated magnification for cleanliness and condition following cleaning?

YES  NO

13. Are handpieces decontaminated in between each patient in accordance with manufacturers' instructions?

YES  NO

14. Are separate canisters of lubricant used for unclean, cleaned and sterilised instruments?

YES  NO

15. Are those handpieces that are manually cleaned, wiped and lubricated with oil before steam sterilisation in accordance with manufacturers' instructions?

YES  NO

16. Are handpieces decontaminated by an automated washer disinfectant lubricated with oil before steam sterilisation in accordance with manufacturers' instructions?

YES  NO

17. Are handpieces decontaminated by an automated washer-disinfector (with a specific handpiece irrigation system) lubricated with oil before steam?

YES  NO

18. Are all instruments washed in a washer-disinfector before they are sterilised?

YES  NO

19. If the practice does not have a washer-disinfector are all instruments cleaned manually before they are sterilised?

YES  NO

20. When manual cleaning, are two sinks (or two bowls in a single unit) used? One for washing and the other for rinsing?

YES  NO

21. Are specially formulated detergents used for manually cleaning instruments and is it used at the specified concentration according to the manufacturer's instructions?

YES  NO

22. When manual cleaning is the water temperature checked and recorded to make sure it is 45°C or lower?

YES  NO

23. Are instruments fully submerged when manually cleaned? (unless the manufacturer's instructions suggest otherwise)

YES  NO

24. Are brushes used to manually clean instruments either single use or washed after each use and replaced at the manufacturer's recommended intervals or when damaged?

YES  NO

25. Are there contractual validation and testing arrangements in place to ensure all autoclaves are routinely maintained and validated in accordance with HTM-01-05 and manufacturers' instructions?

YES  NO

26. Are daily, weekly, quarterly and annual inspection testing and maintenance records available for autoclaves as described in section 12 of HTM-01-05?

YES  NO

27. Are fault logs kept for all autoclaves and are they removed from use until the fault has been rectified?

YES  NO

28. Are there arrangements to ensure all ultrasonic cleaners are maintained and validated in accordance with HTM-01-05 requirements and manufacturers' instructions?

YES  NO

29. Are daily, weekly, quarterly and annual inspections, testing and maintenance records available for ultrasonic cleaners?

YES  NO

30. Are there contractual arrangements in place to ensure all automated washer-disinfectors are routinely maintained and validated in line with manufacturers' instructions?

YES  NO

31. Are daily, weekly, quarterly and annual validation and testing results recorded for automated washer-disinfectors?

YES  NO

32. When using ultrasonic cleaners, are instruments placed in an instrument basket or cassette and fully immersed ensuring that all surfaces are in contact with the solution?

YES  NO

33. When using ultrasonic cleaners (and when not in use) is the lid closed to prevent contamination of the ultrasonic cleaning solution?

YES  NO

34. Is the ultrasonic cleaner solution emptied when visibly contaminated or otherwise at the end of every clinical session?

YES  NO

35. Where instruments are manually cleaned, are they rinsed after removing from the ultrasonic cleaner?

YES  NO



36. Are the relevant staff aware of the instrument loading procedure for the washer disinfector? (spray arms are free to rotate, cannulated instruments are loaded correctly)

YES  NO

37. Are washer-disinfector cycle parameters recorded?

YES  NO

38. For autoclaves, are there daily records available for the past two years of the temperature and pressure achieved on completion of each cycle

YES  NO

39. Can you confirm that autoclaves are never used if they display faults?

YES  NO

40. When using autoclaves, can you confirm that only pre-wrapped instruments are used in vacuum type sterilisers?

YES  NO

41. Is distilled water or RO (reverse osmosis) water used in the autoclave?

YES  NO

42. Are opened bottles of distilled water discarded at the end of each working day?

YES  NO

43. Are autoclave reservoirs drained and left clean and dry at the end of each day?

YES  NO

44. Is there a separate dedicated decontamination room which is restricted to those performing decontamination duties?

YES  NO

45. Is the decontamination room clearly zoned into clean and dirty areas?

YES  NO

46. Are decontamination areas and work surfaces clean and uncluttered?

YES  NO

47. Is there adequate ventilation in the decontamination room?

YES  NO

48. Where full mechanical ventilation is used in the decontamination room, is the direction of air flowing from the clean to the dirty area?

YES  NO

49. Where full mechanical ventilation is used in the decontamination room, is the direction of air flowing from the clean to the dirty area?

YES  NO

50. Are instruments maintained in a moist condition between use and decontamination?

YES  NO

51. Are instrument transport containers lidded, clean, leak proof and cleaned, disinfected and dried following each use?

YES  NO

52. Are instruments processed in a non-vacuum (type N) autoclave dried prior to packing using a disposable lint free cloth?

YES  NO

53. Does the practice have a system in place to ensure that the storage of non-wrapped instruments does not exceed 1 day if stored in a clinical area?

YES  NO

54. Does the practice have a system in place to ensure that the storage of wrapped instruments does not exceed 1 year?

YES  NO

55. Is there a system in place to identify instrument storage time, including the date by which they should be used or re-processed?

YES  NO

56. Are instruments stored in a dedicated secure, dry and cool environment?

YES  NO

### Section 3: Environmental Design and Cleaning

1. Does the practice have a policy and procedure for cleaning and maintaining the environment?

YES  NO

2. Are daily cleaning and decontamination checklists present and filled in for all clinical areas? Can you provide two years of evidence of this?

YES  NO

3. Does the practice have a daily checklist for the cleaner?

YES  NO

4. Does the practice have a COVID-19 Policy and Procedure in place?

YES  NO

5. Are two metre social distancing markers on the floor in all areas during the pandemic?

YES  NO

6. Is hand sanitising gel readily available in all rooms?

YES  NO

7. Is there a clear shield in front of the reception desk to protect staff and patients?

YES  NO

8. Have staff undertaking cleaning duties been fully trained to undertake such duties?undertake such duties?

YES  NO

9. Are surgery floors mopped after each session?

YES  NO

10. Is the overall appearance of the clinical and decontamination environment tidy and uncluttered?

YES  NO

11. Are all dental chairs free of rips or tears and cleaned in between each patient?

YES  NO

12. Are all walls, floors, ceilings, fixtures and fittings free from damage?

YES  NO

13. Are all work surface joints intact, seamless and with no visible damage?

YES  NO

14. Are all surfaces free from dust and visible dirt?

YES  NO

15. Are the surfaces of accessible ventilation fittings cleaned weekly?

YES  NO

16. Are easy clean keyboards used in all clinical areas?

YES  NO

17. Can you confirm that none of the clinical areas are carpeted?

YES  NO

18. Do all floor coverings in the clinical and decontamination areas have coved edges that are sealed?

YES  NO

19. Is cleaning equipment colour coded to determine what area it should be used in?

YES  NO

20. Is general cleaning equipment stored in a non-clinical area?

YES  NO

21. Where disposable single-use covers are used are they discarded after each patient contact?

YES  NO

22. Are surfaces and equipment cleaned in between each patient? (dental chairs, curing lamps, light handles, the dental cart, spittoon etc.)

YES  NO

23. Is fallow time used where needed following the latest guidelines during the pandemic?

YES  NO

24. Are AGP - Do not enter signs placed on doors when being undertaken?

YES  NO

25. Are all taps, drainage points, splashbacks, sinks, aspirators, drains and spittoons cleaned after every session with a surface disinfectant?

YES  NO

26. Are floors, cupboard doors and accessible high level surfaces cleaned daily?

YES  NO

27. Is there a designated area for the disposal of dirty water which is separate to the kitchen, clinical and decontamination areas?

YES  NO

28. Does the practice have a COSHH Policy and Procedure?

YES  NO

29. Are frequently touched sites/points, in particular waiting room areas cleaned regularly throughout the day?

YES  NO

## Section 4: Hand Hygiene

1. Does the practice have a policy and procedure for hand hygiene?

YES  NO

2. Is hand hygiene an integral part of staff induction?

YES  NO

3. Is hand hygiene training provided periodically throughout the year?

YES  NO

4. Are hand hygiene audits performed regularly for all staff?

YES  NO

5. Is hand hygiene carried out before donning gloves and after removing them?

YES  NO

6. Do all staff involved in any clinical and decontamination procedures have short nails that are clean and free from nail extensions and varnish?

YES  NO

7. Do all clinical and decontamination staff remove watches and rings with stones during clinical and decontamination procedures?

YES  NO

8. Are laminated or wipe clean hand hygiene posters on display by all sinks?

YES  NO

9. Is there a separate dedicated hand basin provided for hand hygiene in each surgery and is it clearly marked?

YES  NO

10. Are no contact liquid soap dispensers available in all surgeries and decontamination areas?

YES  NO

11. Are elbow or foot operated electronic mixers or thermostatically controlled taps available at all hand wash sinks in clinical and decontamination areas?

YES  NO

12. Can you confirm that nail brushes are never used?

YES  NO

13. Are good quality disposable and absorbent paper towels used when drying hands?

YES  NO

14. Is hand cream available for all clinical and decontamination staff?

YES  NO

## Section 5: Management of Dental Medical Devices

1. Does the practice's infection control policy include procedures for the use, maintenance, service and repair of all medical devices?

YES  NO

2. Does the practice have an individual responsible for ensuring that all staff comply with medical device procedures?

YES  NO

3. Has the practice had a legionella risk assessment of the building carried out?

YES  NO

4. Does the practice have a legionella policy and procedure?

YES  NO

5. Are regular temperature checks carried out and recorded according to the risk assessment?

YES  NO

6. Are all new reusable instruments sterilised prior to use?

YES  NO

7. Are instruments sent for repair labelled to identify they have been through the decontamination process?

YES  NO

8. Are contaminated patient devices (dentures, impressions etc.) decontaminated and labelled before being sent to the laboratory?

YES  NO

9. Are patient devices returned from the laboratory disinfected before being placed in the patient's mouth?

YES  NO

10. Are single use items always discarded after use?

YES  NO

11. Can you confirm that single use endodontic instruments are never re-used?

YES  NO

12. Are contaminated intra-oral x-ray digital sensor wrappers discarded after use, the sensor cleaned according to the manufacturer's instructions then re-wrapped?

YES  NO

13. Are x-ray film holders sterilised after every use according to the manufacturer's instructions?

YES  NO

14. Are difficult to clean instruments and devices identified as single use?

YES  NO

15. Are dental chair filters cleaned and replaced as per the manufacturer's instructions?

YES  NO

16. Does the dental chair have an independent bottled water system to dispense distilled or reverse osmosis water?

YES  NO

17. Is the bottle removed, cleaned and dried at the end of each day?

YES  NO

18. If a bottled water chair system is not used is there a physical air gap separating dental unit water lines from the mains water system?

YES  NO

19. For surgical procedures involving irrigation, is sterile saline available?

YES  NO

20. Are dental unit water lines flushed through for a minimum of two minutes at the start of the day and for a minimum of 20-30 seconds between every patient?

YES  NO

21. Are handpieces fitted with anti-retraction valves?

YES  NO

22. Are dental unit water lines purged using the manufacturer's recommended disinfectants?

YES  NO

23. Are dental unit water line filters changed according to the manufacturer's guidelines?

YES  NO

## Section 6: Personal Protective Equipment

1. Does the practice have a PPE policy and procedure that includes COVID government guidelines?

YES  NO

2. Are FFP3 masks/hoods, gloves, disposable aprons and visors worn for all AGPs

YES  NO

3. Are PPE donning and doffing areas clearly marked?

YES  NO

4. Are staff trained in the use of PPE as part of their practice induction?

YES  NO

5. Are powder free and latex free CE marked gloves used?

YES  NO

6. Are all single use PPE items disposed of after each episode of patient care?

YES  NO



7. Are heavy duty gloves available for decontamination processes where necessary, and washed with detergent and hot water and left to dry after each use?

YES  NO

8. Are visors, masks, gloves and disposable aprons worn during all decontamination procedures?

YES  NO

9. Are single use disposable aprons and masks disposed of as clinical waste after each procedure?

YES  NO

10. Are uniforms worn by staff changed at the end of each day and when visibly contaminated?

YES  NO

11. Can you confirm that staff uniforms are never worn outside of the practice?

YES  NO

12. Are staff uniforms washed separately and at the highest temperature possible for the fabric?

YES  NO

13. Are all PPE items stored in accordance with the manufacturer's instructions?

YES  NO

14. Are visors decontaminated after clinical sessions and decontamination procedures?

YES  NO

15. Is eye protection provided for the patient decontaminated after each use?

YES  NO

16. Are posters about the need to cover the nose and mouth during the pandemic present in the practice?

YES  NO

17. Do all patients wear a face mask on entering and exiting the building?

YES  NO

## Section 7: Waste Management

1. Does the practice have a policy and procedure for the management and disposal of waste?

YES  NO

2. Have all staff been trained at their induction on the process of waste disposal?

YES  NO

3. Is there evidence that your waste contractors are registered waste carriers?

YES  NO

4. Is there a contract in place for the removal of clinical waste, sharps, amalgam filled extracted teeth and out of date medicines?

YES  NO

5. Is the practice registered with the environmental agency if generating over 500KG of hazardous waste per year?

YES  NO

6. Are orange clinical waste bags used for infectious category B waste such as blood swabs, blood contaminated gloves and teeth without amalgam fillings?

YES  NO

7. Are yellow (with a black stripe) bags used for offensive/hygiene waste that is not contaminated with saliva, blood, medicines, chemicals or amalgam? for example, tissues and non-contaminated gloves

YES  NO

8. Are black or clear bags used for domestic waste including paper towels?

YES  NO

9. Are bins foot operated or sensor controlled, lidded and in good working order?

YES  NO

10. Are local anaesthetic cartridges disposed of in yellow sharps containers that conform to guidelines?

YES  NO

11. Are clinical waste bags tied securely and sharps containers clicked shut before they are collected by the waste company?

YES  NO

12. Are all clinical waste bags and sharps bins labelled with the practice name, address and date before they are collected?

YES
  NO

13. Is waste awaiting collection stored in a safe and secure location away from the public within the practice premises?

YES
  NO

14. Are waste consignment notes available for all hazardous waste for the past 3 years?

YES
  NO

15. Is there evidence that the practice is segregating waste in accordance with HTM-07-01?

YES
  NO

16. Has the practice been assured that a duty of care audit has been carried out and recorded from the producer to the final disposal?

YES
  NO

## Audit Results and Action Plan:

The purpose of this audit is to see what you are already doing well and identify areas for improvement. For all questions you should be aiming for a 'yes' answer.

The below table is broken down into sections for you to fill in so you can clearly see which areas need improvement and form an action plan to turn the 'no' answers into 'yes' answers.

Section	Number of questions	Number of questions answered yes	Number of questions answered no	Question numbers with 'no' answers requiring further action	Action plan	Date to be completed by
1	14					
2	56					

3	29					
4	14					
5	23					
6	17					
7	16					

Section	% of yes answers
1	
2	
3	
4	
5	
6	
7	

**Total overall % of yes answers**

Next audit review date: \_\_\_\_\_  
(6 months from now)