



The Regulation and
Quality Improvement
Authority

Announced Inspection

Name of Establishment: Quinndental
Establishment ID No: 11563
Date of Inspection: 26 August 2014
Inspector's Name: Emily Campbell
Inspection No: 18349

The Regulation and Quality Improvement Authority
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1.0 General Information

Name of establishment:	Quinndental
Address:	53 Main Street Randalstown BT41 3BB
Telephone number:	028 9447 2311
Registered organisation / registered provider:	Mr Liam Quinn
Registered manager:	Mr Liam Quinn
Person in charge of the establishment at the time of Inspection:	Mr Liam Quinn
Registration category:	IH-DT
Type of service provision:	Private dental treatment
Maximum number of places registered: (dental chairs)	2
Date and type of previous inspection:	Announced Inspection 16 December 2013
Date and time of inspection:	26 August 2014 10.00am – 1.35pm
Name of inspector:	Emily Campbell

2.0 Introduction

The Regulation and Quality Improvement Authority (RQIA) is empowered under The Health and Personal Social Services (Quality, Improvement and Regulation) (Northern Ireland) Order 2003 to inspect dental practices providing private dental care and treatment. A minimum of one inspection per year is required.

This is a report of the announced inspection to assess the quality of services being provided. The report details the extent to which the standards measured during inspection were met.

3.0 Purpose of the Inspection

The purpose of this inspection was to consider whether the service provided to patients was in accordance with their assessed needs and preferences and was in compliance with legislative requirements, minimum standards and other good practice indicators. This was achieved through a process of analysis and evaluation of available evidence.

RQIA not only seeks to ensure that compliance with regulations and standards is met but also aims to use inspection to support providers in improving the quality of services. For this reason, inspection involves in-depth examination of an identified number of aspects of service provision.

The aims of the inspection were to examine the policies, practices and monitoring arrangements for the provision of dental care, and to determine the provider's compliance with the following:

- The HPSS (Quality, Improvement and Regulation) (Northern Ireland) Order 2003;
- The Independent Health Care Regulations (Northern Ireland) 2005;
- The Regulation and Improvement Authority (Independent Health Care) (Fees and Frequency of Inspections) (Amendment) Regulations (Northern Ireland) 2011;
- The Minimum Standards for Dental Care and Treatment 2011; and
- Health Technical Memorandum HTM 01-05: Decontamination in Primary Care Dental Practices and Professional Estates Letter (PEL) (13) 13.

Other published standards which guide best practice may also be referenced during the inspection process.

4.0 Methods/Process

Committed to a culture of learning, the RQIA has developed an approach which uses self-assessment, a critical tool for learning, as a method for preliminary assessment of achievement of the Minimum Standards.

The inspection process has three key parts; self-assessment (including completion of self-declaration), pre-inspection analysis and the inspection visit by the inspector.

Specific methods/processes used in this inspection include the following:

- a self-assessment was submitted prior to the inspection and has been analysed;
- discussion with Mr Liam Quinn, registered provider;
- examination of relevant records;
- consultation with relevant staff;
- tour of the premises; and
- evaluation and feedback.

Any other information received by RQIA about this practice has also been considered by the inspector in preparing for this inspection.

5.0 Consultation Process

During the course of the inspection, the inspector spoke with staff on duty. Questionnaires were provided to staff prior to the inspection by the practice, on behalf of the RQIA to establish their views regarding the service. Matters raised by staff were addressed by the inspector during the course of this inspection:

	Number	
Discussion with staff	2	
Staff Questionnaires	4 issued	4 returned

Prior to the inspection the registered person/s were asked, in the form of a declaration, to confirm that they have a process in place for consulting with service users and that a summary of the findings has been made available. The consultation process may be reviewed during this inspection.

6.0 Inspection Focus

The inspection sought to establish the level of compliance achieved with respect to the selected DHSSPS Minimum Standards for Dental Care and Treatment and a thematic focus incorporating selected standards and good practice indicators. An assessment on the progress in relation to the issues raised during and since the previous inspection was also undertaken.

In 2012 the DHSSPS requested that RQIA make compliance with best practice in local decontamination, as outlined in HTM 01-05 Decontamination in Primary Care Dental Premises, a focus for the 2013/14 inspection year.

The DHSSPS and RQIA took the decision to review compliance with best practice over two years. The focus of the two years is as follows:

- Year 1 – Decontamination – 2013/14 inspection year
- Year 2 - Cross infection control – 2014/15 inspection year

Standard 13 – Prevention and Control of Infection [Safe and effective care]

The dental service takes every reasonable precaution to make sure you are not exposed to risk of infection.

The decontamination section of the Infection Prevention Society Audit tool, which has been endorsed by the Department of Health, was used as a framework for development of a self-assessment tool and for planned inspections during 2013/14.

The following sections of the 2013 edition of the Infection Prevention Society Audit tool, which has been endorsed by the Department of Health have been used as a framework for the development of a self-assessment tool and for planned inspections in 2014/15:

- prevention of blood-borne virus exposure;
- environmental design and cleaning;
- hand hygiene;
- management of dental medical devices;
- personal protective equipment; and
- waste.

A number of aspects of the Decontamination section of the Audit tool have also been revisited.

RQIA have highlighted good practice guidance sources to service providers, making them available on our website where possible. Where appropriate, requirements will be made against legislation and recommendations will be made against DHSSPS Minimum Standards for Dental Care and Treatment (2011) and other recognised good practice guidance documents.

The registered provider/manager and the inspector have each rated the practice's compliance level against each section of the self-assessment.

The table below sets out the definitions that RQIA has used to categorise the service's performance:

Guidance - Compliance statements		
Compliance statement	Definition	Resulting Action in Inspection Report
0 - Not applicable		A reason must be clearly stated in the assessment contained within the inspection report.
1 - Unlikely to become compliant		A reason must be clearly stated in the assessment contained within the inspection report.
2 - Not compliant	Compliance could not be demonstrated by the date of the inspection.	In most situations this will result in a requirement or recommendation being made within the inspection report.
3 - Moving towards compliance	Compliance could not be demonstrated by the date of the inspection. However, the service could demonstrate a convincing plan for full compliance by the end of the Inspection year.	In most situations this will result in a requirement or recommendation being made within the inspection report.
4 – Substantially Compliant	Arrangements for compliance were demonstrated during the inspection. However, appropriate systems for regular monitoring, review and revision are not yet in place.	In most situations this will result in a recommendation, or in some circumstances a requirement, being made within the inspection report.
5 – Compliant	Arrangements for compliance were demonstrated during the inspection. There are appropriate systems in place for regular monitoring, review and any necessary revisions to be undertaken.	In most situations this will result in an area of good practice being identified and comment being made within the inspection report.

7.0 Profile of Service

Quinndental is located within a terrace dwelling that has been converted to accommodate a dental practice. It is situated on a retail street in Randalstown. There is time limited on street car parking and a public car park is available within walking distance.

The practice is not accessible for patients with a disability. Arrangements are in place to accommodate patients with a disability who cannot access the surgery.

The practice provides two dental surgeries – one of which is on the ground floor and the other on the first floor of the building. In addition there is a reception area, a waiting area, a decontamination room, staff and storage facilities. Toilet facilities are available on the first floor for patient use.

Quinndental operates two dental chairs providing both private and NHS dental care.

General dentistry services are provided by Mr Liam Quinn with support from dental nurses and a hygienist. Mr Quinn had been the registered provider/manager since registration with RQIA in April 2012.

The establishment's statement of purpose outlines the range of services provided.

This practice is registered with RQIA as an independent hospital (IH) providing dental treatment (DT).

8.0 Summary of Inspection

This announced inspection of Quindental was undertaken by Emily Campbell on 26 August 2014 between the hours of 10.00am and 12.35pm. Mr Liam Quinn, registered provider, was available during the inspection and for verbal feedback at the conclusion of the inspection.

The requirements and recommendations made as a result of the previous inspection were also examined. Observations and discussion demonstrated that two of the three requirements and three of the four recommendations made have been addressed. One requirement regarding the refurbishment of the decontamination room has not been addressed and is stated for the second time. One recommendation has been partially addressed and the unaddressed aspect regarding the storage of items on the floor of the decontamination room is stated for the second time. The detail of the action taken by Mr Quinn can be viewed in the section following this summary.

Prior to the inspection, Mr Quinn completed a self-assessment using the standard criteria outlined in the theme inspected. The comments provided by Mr Quinn in the self-assessment were not altered in any way by RQIA. The self-assessment is included as appendix one in this report. Mr Quinn did not rate the practice compliance levels against each criterion; however, these were provided verbally to the inspector during the inspection.

During the course of the inspection the inspector met with staff, discussed operational issues, examined a selection of records and carried out a general inspection of the establishment.

Questionnaires were also issued to staff; four were returned to RQIA within the timescale required. Review of submitted questionnaires and discussion with staff evidenced that staff were knowledgeable regarding the inspection theme and that they have received training appropriate to their relevant roles. Staff confirmed that they are familiar with the practice policies and procedures and have received infection prevention and control training. Clinical staff also confirmed that they have been immunised against Hepatitis B.

Inspection Theme – Cross infection control

Dental practices in Northern Ireland have been directed by the DHSSPS, that best practice recommendations in the Health Technical Memorandum (HTM) 01-05, Decontamination in primary care dental practices, along with Northern Ireland amendments, should have been fully implemented by November 2012. HTM 01-05 was updated in 2013 and Primary Care Dental Practices were advised of this through the issue of Professional Estates Letter (PEL) (13) 13 on 1 October 2013. The PEL (13) 13 advised General Dental Practitioners of the publication of the 2013 version of HTM 01-05 and the specific policy amendments to the guidance that apply in Northern Ireland.

RQIA reviewed the compliance of the decontamination aspect of HTM 01-05 in the 2013/2014 inspection year. The focus of the inspection for the 2014/2015

inspection year is cross infection control. A number of aspects of the decontamination section of HTM 01-05 have also been revisited.

A copy of the 2013 edition of HTM 01-05 Decontamination in primary dental care practices is available at the practice for staff reference. Staff are familiar with best practice guidance outlined in the document and audit compliance on an ongoing basis.

The practice has a policy and procedure in place for the prevention and management of blood-borne virus exposure, including management of spillages, sharps and inoculation incidents in accordance with national guidance. A recommendation was made that the procedure for the management of blood/bodily spillages is further developed. Review of documentation and discussion with Mr Quinn and staff evidenced that appropriate arrangements are in place for the prevention and management of blood-borne virus exposure. Staff confirmed that they are aware of and are adhering to the practice policy in this regard. Sharps management at the practice was observed to be in line with best practice, with the exception of the provision of purple lidded sharps boxes for the disposal of pharmaceutical sharps waste. A recommendation was made in this regard.

The practice has a policy and procedure in place for cleaning and maintaining the environment. A recommendation was made that this is further developed to include the cleaning arrangements for the decontamination room and the flooring in the surgeries. Clinical and decontamination areas were tidy and uncluttered and work surfaces were intact and easy to clean. Floor coverings are impervious; however, they were not sealed at the edges or where cabinetry meets the flooring. A recommendation was made to address this. Fixtures, fittings, dental chairs and equipment were free from damage, dust and visible dirt, with the exception of a fan which had a build-up of dust on the blades and cover. A recommendation was made in this regard. Discussion with staff confirmed that although cupboard doors and accessible high level surfaces are cleaned on a daily basis, the floors of surgeries and the decontamination room are not cleaned daily. A recommendation was made in this regard.

The practice has a hand hygiene policy and procedure in place and staff demonstrated that good practice is adhered to in relation to hand hygiene. Dedicated hand washing basins are available in the appropriate locations. A recommendation was made that the overflow of the hand washing basin in the first floor surgery should be blanked off with a stainless steel plate sealed with antibacterial mastic. Information promoting hand hygiene is provided for staff.

A written scheme for the prevention of legionella is available. A recommendation was made that the legionella risk assessment is further developed to include the specific arrangements for the management of dental unit water lines (DUWLs). Procedures are in place for the use, maintenance, service and repair of all medical devices. Observations made and discussion with staff confirmed that DUWLs are appropriately managed.

The practice has a policy and procedure in place for the use of personal protective equipment (PPE) and staff spoken with demonstrated awareness of this. Observations made confirmed that PPE was readily available and used appropriately by staff.

Appropriate arrangements were in place for the management of general and clinical waste, including sharps, with the exception of the provision of purple lidded sharps boxes for pharmaceutical waste as discussed previously. Waste was appropriately segregated and suitable arrangements were in place for the storage and collection of waste by a registered waste carrier. Relevant consignment notes are retained in the practice for at least three years.

A decontamination room separate from patient treatment areas and dedicated to the decontamination process is available. During the previous inspection a requirement was made that the decontamination room should be refurbished. This requirement has not been addressed and is stated for the second time with a final three month timescale agreed. Further details can be seen in section 10.7 of the report. A recommendation was made for the second time that boxes, and used clinical waste bags should not be stored on the floor of the decontamination room.

Appropriate validated equipment, including a washer disinfecter, a DAC Universal and a steam steriliser have been provided to meet the practice requirements. Equipment logbooks evidenced that periodic tests are undertaken and recorded in keeping with HTM 01-05.

The evidence gathered through the inspection process concluded that Quinndental is moving towards compliance with this inspection theme.

Mr Quinn confirmed on the submitted self-assessment that arrangements are in place for consultation with patients, at appropriate intervals, that feedback provided by patients has been used by the service to improve and that results of the consultation have been made available to patients.

One requirement and eight recommendations were made as a result of the announced inspection; details can be found in the main body of the report and the attached Quality Improvement Plan (QIP).

The inspector wishes to thank Mr Quinn and staff for their helpful discussions, assistance and hospitality throughout the inspection process.

9.0 Follow-up on Previous Issues

No	Regulation Ref.	Requirements	Action taken - as confirmed during this inspection	Inspector's Validation of Compliance
1	25 (2) (b)	<p>The decontamination room should be refurbished.</p> <p>The layout of the room should be in keeping with HTM 01-05.</p>	<p>Observations made and discussion with Mr Quinn confirmed that this requirement has not been addressed. Further details in relation to this matter can be seen in section 10.7 of the report.</p> <p>The importance of ensuring that the decontamination room is suitably completed in keeping with HTM 01-05 was discussed with Mr Quinn and a final three month timescale was agreed. Mr Quinn should keep RQIA apprised regarding the refurbishment of the decontamination room. This requirement is stated for the second time and will be kept under review.</p>	Not compliant
2	15 (3)	Suction tips and any other lumen devices, dental burrs and all other dental instruments must be processed through the washer disinfectant unless the instrument is incompatible with the washer disinfectant.	Discussion with a dental nurse confirmed that this requirement has been addressed. Disposable suction tips are now in use in the practice.	Compliant
3	15 (2) (b)	The washer disinfectant and steriliser should be validated in accordance with manufacturer's instructions and best practice as outlined in HTM 01-05 and arrangements established for annual validation thereafter.	<p>Review of documentation confirmed that decontamination equipment has been validated. Mr Quinn confirmed that arrangements have been established for annual re-validation.</p> <p>Requirement addressed.</p>	Compliant

No	Minimum Standard Ref.	Recommendations	Action Taken – as confirmed during this inspection	Inspector's Validation of Compliance
1	13	<p>Boxes should not be stored on the floor of the decontamination room.</p> <p>The clinical waste bin in the decontamination room should be pedal operated.</p>	<p>A pedal operated clinical waste bin was available at the dirty side of the decontamination room. However, the inspector observed that a box, sharps boxes and used clinical waste bags were stored on the floor of the room.</p> <p>This recommendation has been partially addressed and a recommendation was made for the second time that boxes, and used clinical waste bags should not be stored on the floor of the decontamination room.</p>	Moving towards compliance.
2	13	Further develop the DAC Universal logbook to incorporate the periodic tests for a washer disinfectant including the daily cleaning efficacy test and the weekly protein residue test.	Review of the DAC Universal logbook evidenced that this recommendation has been addressed.	Compliant
3	13	A weekly protein residue test should be undertaken and recorded in the logbook for the washer disinfectant.	Discussion with a dental nurse and review of the washer disinfectant logbook confirmed that this recommendation has been addressed.	Compliant
4	13	Further develop the policies and procedures to reflect the arrangements for the decontamination of dental instruments.	Review of documentation confirmed that this recommendation has been addressed. The inspector suggested that some minor amendments are made regarding the transport of dental instruments and the use of the DAC Universal within the decontamination process.	Compliant

10.0 Inspection Findings

10.1 Prevention of blood-borne virus exposure

STANDARD 13 – Prevention and Control of Infection (Safe and effective care)

The dental service takes every reasonable precaution to make sure you are not exposed to risk of infection.

Criteria Assessed:

11.2 You receive care and treatment from a dental team (including temporary members) who have undergone appropriate checks before they start work in the service.

13.2 Your dental service adheres to the appropriate infection control policies and procedures in line with current best practice and legislation.

13.3 Your dental service has systems in place, including induction and ongoing training, to make sure these policies and procedures are known, and are being appropriately applied to the service at all times.

Inspection Findings:

Mr Quinn verbally rated the practice arrangements for the prevention of blood-borne virus exposure as compliant during the inspection.

The practice has a policy and procedure in place for the prevention and management of blood-borne virus exposure, including management of spillages, sharps and inoculation incidents in accordance with national guidance. Discussion with staff confirmed that staff are aware of the policies and procedures in in this regard. The procedure for the management of blood/bodily spillages indicates the use of a specified product, however, Mr Quinn and staff confirmed that household bleach is the product now available for this. Staff were not aware of the dilution strength to be used. A recommendation was made that the procedure is further developed to reflect the new arrangements in the practice including the relevant dilution rate. Staff should be made aware of these arrangements.

Review of documentation and discussion with staff evidenced that:

- the prevention and management of blood-borne virus exposure is included in the staff induction programme;
- staff training has been provided for clinical staff; and
- records are retained regarding the Hepatitis B immunisation status of clinical staff.

Mr Quinn confirmed that any new staff recruited will receive an occupational health check.

Discussion with staff confirmed that staff are aware of the policies and procedures in place for the prevention and management of blood-borne virus exposure.

Observations made and discussion with staff evidenced that sharps are appropriately handled. Sharp's boxes are wall mounted, appropriately used, signed and dated on assembly and final closure. A recommendation was made that purple lidded sharp's boxes for the disposal of pharmaceutical sharps waste are provided. Used sharp's boxes are locked with the integral lock and stored ready for collection away from public access.

Discussion with staff and review of documentation evidenced that arrangements are in place for the management of a sharps injury, including needle stick injury. Staff are aware of the actions to be taken in the event of a sharps injury.

Provider's overall assessment of the dental practice's compliance level against the standard assessed	Compliant
Inspector's overall assessment of the dental practice's compliance level against the standard assessed	Substantially compliant

10.2 Environmental design and cleaning

<p>STANDARD 13 – Prevention and Control of Infection (Safe and effective care) The dental service takes every reasonable precaution to make sure you are not exposed to risk of infection.</p>
<p>Criterion Assessed: 13.1 Your dental service's premises are clean.</p>
<p>Inspection Findings:</p> <p>Mr Quinn verbally rated the practice arrangements for environmental design and cleaning as substantially compliant during the inspection.</p> <p>The practice has a policy and procedure in place for cleaning and maintaining the environment. A recommendation was made that this is further developed to include the cleaning arrangements for the decontamination room and the flooring in the surgeries.</p> <p>The inspector undertook a tour of the premises which were found to be maintained to a good standard of cleanliness. Clinical and decontamination areas were tidy and uncluttered and work surfaces were intact and easy to clean. Floor coverings are impervious; however, they were not sealed at the edges or where cabinetry meets the flooring. A recommendation was made to address this. Mr Quinn confirmed that the new flooring in the decontamination room would be covered at the edges on refurbishment. Fixtures, fittings, dental chairs and equipment were free from damage, dust and visible dirt, with the exception of a fan which had a build-up of dust on the blades and cover. A recommendation was made that this should be addressed and included in the cleaning schedule.</p> <p>Discussion with staff confirmed that appropriate arrangements are in place for cleaning including:</p> <ul style="list-style-type: none"> • Equipment surfaces, including the dental chair, are cleaned between each patient; • Weekly/monthly cleaning schedule; • Cleaning equipment is colour coded; • Cleaning equipment is stored in a non-clinical area; and • Dirty water is disposed of at an appropriate location. <p>Discussion with staff confirmed that although cupboard doors and accessible high level surfaces are cleaned on a daily basis, the floors of surgeries and the decontamination room are not cleaned daily. A recommendation was made in this regard.</p> <p>Discussion with staff and review of submitted questionnaires confirmed that staff had received relevant training to undertake their duties.</p> <p>The practice has a local policy and procedure for spillage in accordance with the Control of Substances Hazardous to Health (COSHH) and staff spoken with demonstrated awareness of this.</p>

Provider's overall assessment of the dental practice's compliance level against the standard assessed	Substantially compliant
Inspector's overall assessment of the dental practice's compliance level against the standard assessed	Substantially compliant

10.3 Hand Hygiene

<p>STANDARD 13 – Prevention and Control of Infection (Safe and effective care) The dental service takes every reasonable precaution to make sure you are not exposed to risk of infection.</p>
<p>Criteria Assessed: 13.2 Your dental service adheres to the appropriate infection control policies and procedures in line with current best practice and legislation. 13.3 Your dental service has systems in place, including induction and ongoing training, to make sure these policies and procedures are known, and are being appropriately applied to the service at all times.</p>
<p>Inspection Findings: Mr Quinn verbally rated the practice arrangements for hand hygiene as compliant during the inspection.</p> <p>The practice has a hand hygiene policy and procedure in place.</p> <p>Staff confirmed that hand hygiene is included in the induction programme and that hand hygiene training is updated periodically.</p> <p>Discussion with staff confirmed that hand hygiene is performed before and after each patient contact and at appropriate intervals. Observations made evidenced that clinical staff had short clean nails and jewellery such as wrist watches and stoned rings were not worn in keeping with good practice.</p> <p>Dedicated hand washing basins are available in the dental surgeries and the decontamination room and adequate supplies of liquid soap, paper towels and disinfectant rub/gel were available. The hand washing basins in both surgeries have overflows. As the hand washing basin in the first floor surgery is made of stainless steel, the overflow can be blanked off with a stainless steel plate sealed with antibacterial mastic and a recommendation was made in this regard. The inspector suggested that the hand washing basin in the ground floor surgery is replaced with a clinical hand wash basin on the next refurbishment of the surgery. Staff confirmed that nail brushes and bar soap are not used in the hand hygiene process in keeping with good practice.</p> <p>The inspector observed that laminated posters promoting hand hygiene were on display in dental surgeries and the decontamination room. The inspector suggested that posters promoting hand hygiene are also displayed in toilet facilities.</p>

<p>Provider’s overall assessment of the dental practice’s compliance level against the standard assessed</p>	<p>Compliant</p>
<p>Inspector’s overall assessment of the dental practice’s compliance level against the standard assessed</p>	<p>Substantially compliant</p>

10.4 Management of Dental Medical Devices

<p>STANDARD 13 – Prevention and Control of Infection (Safe and effective care) The dental service takes every reasonable precaution to make sure you are not exposed to risk of infection.</p>
<p>Criterion Assessed: 13.4 Your dental service meets current best practice guidance on the decontamination of reusable dental and medical instruments.</p>
<p>Inspection Findings:</p> <p>Mr Quinn verbally rated the practice approach to the management of dental medical devices as compliant during the inspection.</p> <p>The practice has an infection control policy that includes procedures for the use, maintenance, service and repair of all medical devices.</p> <p>The inspector reviewed the written scheme for the prevention of legionella contamination in water pipes and other water lines and discussion with staff confirmed that this is adhered to. A recommendation was made that the legionella risk assessment is further developed to include the specific arrangements for the management of DUWLs. This should also be included in the practice’s infection control policy.</p> <p>Staff confirmed that impression materials, prosthetic and orthodontic appliances are decontaminated prior to despatch to laboratory and before being placed in the patient’s mouth.</p> <p>Observations made and discussion with staff confirmed that DUWLs are appropriately managed. This includes that:</p> <ul style="list-style-type: none"> • Filters are cleaned/replaced as per manufacturer’s instructions; • Water supply to the DUWLs is provided through the direct mains water supply. Mr Quinn confirmed on discussion that there is a physical air gap separating DUWLs from mains water systems; • DUWLs are flushed at the start of each working day and between every patient; • DUWLs and handpieces are fitted with anti-retraction valves; and • DUWLs are purged using disinfectant as per manufacturer’s recommendations.

<p>Provider’s overall assessment of the dental practice’s compliance level against the standard assessed</p>	<p>Compliant</p>
<p>Inspector’s overall assessment of the dental practice’s compliance level against the standard assessed</p>	<p>Substantially compliant</p>

10.5 Personal Protective Equipment

<p>STANDARD 13 – Prevention and Control of Infection (Safe and effective care) The dental service takes every reasonable precaution to make sure you are not exposed to risk of infection.</p>
<p>Criterion Assessed: 13.2 Your dental service adheres to the appropriate infection control policies and procedures in line with current best practice and legislation. 13.3 Your dental service has systems in place, including induction and ongoing training, to make sure these policies and procedures are known, and are being appropriately applied to the service at all times.</p>
<p>Inspection Findings: Mr Quinn verbally rated the practice approach to the management of personal protective equipment (PPE) as compliant during the inspection.</p> <p>The practice has a policy and procedure in place for the use of PPE and staff spoken with demonstrated awareness of this. Staff confirmed that the use of PPE is included in the induction programme.</p> <p>Observations made and discussion with staff evidenced that PPE was readily available and in use in the practice.</p> <p>Discussion with staff confirmed that:</p> <ul style="list-style-type: none"> • Hand hygiene is performed before donning and following the removal of disposable gloves; • Single use PPE is disposed of appropriately after each episode of patient care; • Heavy duty gloves are available for domestic cleaning and decontamination procedures where necessary; and • Eye protection for staff and patients is decontaminated after each episode. <p>Staff confirmed that they were aware of the practice uniform policy.</p>

<p>Provider’s overall assessment of the dental practice’s compliance level against the standard assessed</p>	<p>Compliant</p>
<p>Inspector’s overall assessment of the dental practice’s compliance level against the standard assessed</p>	<p>Compliant</p>

10.6 Waste

<p>STANDARD 13 – Prevention and Control of Infection (Safe and effective care) The dental service takes every reasonable precaution to make sure you are not exposed to risk of infection.</p>
<p>Criterion Assessed: 13.2 Your dental service adheres to the appropriate infection control policies and procedures in line with current best practice and legislation. 13.3 Your dental service has systems in place, including induction and ongoing training, to make sure these policies and procedures are known, and are being appropriately applied to the service at all times..</p>
<p>Inspection Findings: Mr Quinn verbally rated the practice approach to the management of waste as compliant during the inspection.</p> <p>The practice has a policy and procedure in place for the management and disposal of waste in keeping with HTM 07-01. Staff confirmed that the management of waste is included in the induction programme and that waste management training is updated periodically.</p> <p>Review of documentation confirmed that contracted arrangements are in place for the disposal of waste by a registered waste carrier and relevant consignment notes are retained in the practice for at least three years.</p> <p>Observations made and discussion with staff confirmed that staff are aware of the different types of waste and appropriate disposal streams.</p> <p>A pedal operated clinical waste bin is available in the decontamination room. Clinical waste is disposed of in waste bags within cupboards which are then transported to the clinical waste bin in the decontamination room. Staff demonstrated the non-touch technique in this regard.</p> <p>Appropriate arrangements are in place in the practice for the storage and collection of general and clinical waste, including sharps waste.</p> <p>The inspector observed adequate provision of sharps containers with the exception of the provision of purple lidded sharps boxes for pharmaceutical waste as discussed in section 10.1 of the report.</p>

<p>Provider’s overall assessment of the dental practice’s compliance level against the standard assessed</p>	<p>Compliant</p>
<p>Inspector’s overall assessment of the dental practice’s compliance level against the standard assessed</p>	<p>Compliant</p>

10.7 Decontamination

<p>STANDARD 13 – Prevention and Control of Infection (Safe and effective care) The dental service takes every reasonable precaution to make sure you are not exposed to risk of infection.</p>
<p>Criterion Assessed: 13.4 Your dental service meets current best practice guidance on the decontamination of reusable dental and medical instruments.</p>
<p>Inspection Findings: Mr Quinn verbally rated the decontamination arrangements of the practice as compliant during the inspection.</p> <p>A decontamination room separate from patient treatment areas and dedicated to the decontamination process is available. During the previous inspection a requirement was made that the decontamination room should be refurbished and a timescale for completion of six months was agreed. Observations made and discussion with Mr Quinn confirmed that this requirement has not been addressed. Mr Quinn advised that in order to refurbish the decontamination room, the electrical supply box has to be moved. This involves getting permission from NIE to carry out this work and the period for completion of this is time limited. Due to other circumstances, the work was unable to be completed within the timeframe stipulated previously and subsequently a new application for the works was submitted to NIE. Mr Quinn confirmed that he has recently been provided with new NIE approval and he hopes to get the works completed in the near future.</p> <p>The importance of ensuring that the decontamination room is suitably refurbished in keeping with HTM 01-05 was discussed with Mr Quinn and the requirement is stated for the second time with a final three month timescale agreed. Mr Quinn should keep RQIA apprised regarding the refurbishment of the decontamination room. This matter will be kept under review.</p> <p>As stated in section 9.0, a recommendation was made for the second time that boxes, and used clinical waste bags should not be stored on the floor of the decontamination room.</p> <p>Appropriate equipment, including a washer disinfector, a DAC Universal and a steam steriliser have been provided to meet the practice requirements.</p> <p>Review of documentation evidenced that equipment used in the decontamination process has been appropriately validated.</p> <p>Review of equipment logbooks evidenced that periodic tests are undertaken and recorded in keeping with HTM 01-05.</p>

<p>Provider’s overall assessment of the dental practice’s compliance level against the standard assessed</p>	<p>Compliant</p>
<p>Inspector’s overall assessment of the dental practice’s compliance level against the standard assessed</p>	<p>Moving towards compliance</p>

Inspector's overall assessment of the dental practice's compliance level against the standard assessed	Compliance Level
	Moving towards compliance

11.0 Additional Areas Examined

11.1 Staff Consultation/Questionnaires

During the course of the inspection, the inspector spoke with two dental nurses. Questionnaires were also provided to staff prior to the inspection by the practice on behalf of the RQIA. Four were returned to RQIA within the timescale required.

Review of submitted questionnaires and discussion with staff evidenced that staff were knowledgeable regarding the inspection theme and that they have received training appropriate to their relevant roles. Staff confirmed that they are familiar with the practice policies and procedures and have received infection prevention and control training. Clinical staff also confirmed that they have been immunised against Hepatitis B.

11.2 Patient Consultation

Mr Quinn confirmed on the submitted self-assessment that arrangements are in place for consultation with patients, at appropriate intervals, that feedback provided by patients has been used by the service to improve and that results of the consultation have been made available to patients. The inspector reviewed the summary of the most recent patient consultation completed in December 2013. Patient consultation is carried out through questionnaires being posted out to patients. As the most recent consultation had a low response rate, the inspector discussed other ways that patient consultation could be undertaken. The inspector also suggested further developing the questionnaires to provide a broader range of questions.

12.0 Quality Improvement Plan

The details of the Quality Improvement Plan appended to this report were discussed with Mr Liam Quinn as part of the inspection process.

The timescales for completion commence from the date of inspection.

The registered provider/manager is required to record comments on the Quality Improvement Plan.

Matters to be addressed as a result of this inspection are set in the context of the current registration of your premises. The registration is not transferable so that in the event of any future application to alter, extend or to sell the premises the RQIA would apply standards current at the time of that application.

Enquiries relating to this report should be addressed to:

Emily Campbell
The Regulation and Quality Improvement Authority
9th Floor
Riverside Tower
5 Lanyon Place
Belfast
BT1 3BT



Quality Improvement Plan

Announced Inspection

Quinndental

26 August 2014

The areas where the service needs to improve, as identified during this inspection visit, are detailed in the inspection report and Quality Improvement Plan.

The specific actions set out in the Quality Improvement Plan were discussed with Mr Liam Quinn either during or after the inspection visit.

Any matters that require completion within 28 days of the inspection visit have also been set out in separate correspondence to the registered persons.

Registered providers/managers should note that failure to comply with regulations may lead to further enforcement and/or prosecution action as set out in The HPSS (Quality, Improvement and Regulation) (Northern Ireland) Order 2003.

It is the responsibility of the registered provider/manager to ensure that all requirements and recommendations contained within the Quality Improvement Plan are addressed within the specified timescales.

Matters to be addressed as a result of this inspection are set in the context of the current registration of your premises. The registration is not transferable so that in the event of any future application to alter, extend or to sell the premises the RQIA would apply standards current at the time of that application.

STATUTORY REQUIREMENTS

This section outlines the actions which must be taken so that the registered person/s meets legislative requirements based on The HP (Quality, Improvement and Regulation) (Northern Ireland) Order 2003, and The Independent Health Care Regulations (NI) 2005 as amended.

NO.	REGULATION REFERENCE	REQUIREMENTS	NUMBER OF TIMES STATED	DETAILS OF ACTION TAKEN BY REGISTERED PERSON(S)	TIMESCALE
1	25 (2) (b)	<p>The decontamination room should be refurbished.</p> <p>The layout of the room should be in keeping with HTM 01-05.</p> <p>Mr Quinn should keep RQIA appraised regarding the refurbishment of the decontamination room.</p> <p>Ref 9.0 & 10.7</p>	Two	WORK TO BEGIN 25/10/14	Three months

RECOMMENDATIONS

These recommendations are based on The Minimum Standards for Dental Care and Treatment (2011), research or recognised sources. They promote current good practice and if adopted by the registered person may enhance service, quality and delivery.

NO.	MINIMUM STANDARD REFERENCE	RECOMMENDATIONS	NUMBER OF TIMES STATED	DETAILS OF ACTION TAKEN BY REGISTERED PERSON(S)	TIMESCALE
1	13	Boxes, and used clinical waste bags should not be stored on the floor of the decontamination room. Ref 9.0 & 10.7	Two	DONE	Immediate and ongoing
2	13	The procedure for the management of blood/bodily spillages should be further developed to reflect the current arrangements in the practice including the relevant dilution rate for the product used. Staff should be made aware of these arrangements. Ref 10.1	One	DONE	One month
3	13	Purple lidded sharps boxes for the disposal of pharmaceutical sharps waste should be provided. Ref 10.1 & 10.6	One	DONE	One month

4	13	<p>Floor coverings in surgeries should be sealed at the edges and where cabinetry meets the flooring in surgeries and the decontamination room.</p> <p>The new flooring in the decontamination room should be covered at the edges on refurbishment.</p> <p>Ref 10.2</p>	One	BEING DONE 28/10/14	Three months
5	13	<p>Fans should be retained dust free. This should be included in the cleaning schedule.</p> <p>Ref 10.2</p>	One	BEING DONE	Immediate and ongoing
6	13	<p>The floors of surgeries and the decontamination room should be cleaned daily.</p> <p>Ref 10.2</p>	One	BEING DONE	Immediate and ongoing
7	13	<p>The overflow of the hand washing basin in the first floor surgery should be blanked off with a stainless steel plate sealed with antibacterial mastic.</p> <p>Ref 10.3</p>	One	BEING DONE 28/10/14	Three months

8	13	<p>The legionella risk assessment is further developed to include the specific arrangements for the management of DUWLs. This should also be included in the practice's infection control policy.</p> <p>Ref 10.4</p>	One	BEING DONE	Three months
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Please complete the following table to demonstrate that this Quality Improvement Plan has been completed by the registered manager and approved by the responsible person / identified responsible person and return to independent.healthcare@rqia.org.uk

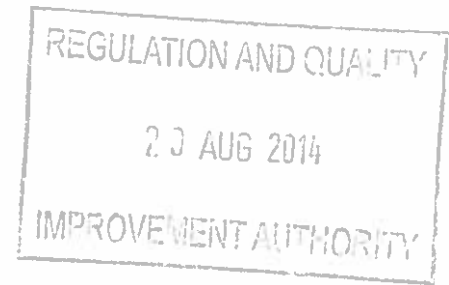
Name of Registered Manager Completing QIP	LIAM QUINN
Name of Responsible Person / Identified Responsible Person Approving QIP	LIAM QUINN

QIP Position Based on Comments from Registered Persons	Yes	Inspector	Date

Response assessed by inspector as acceptable	Yes	E Campbell	20.10.14
Further information requested from provider			



**The Regulation and
Quality Improvement
Authority**



**Self Assessment audit tool of compliance with
HTM01-05 - Decontamination - Cross Infection Control**

Name of practice: Quinndental
RQIA ID: 11563
Name of inspector: Emily Campbell

This self-assessment tool should be completed in reflection of the current decontamination and cross infection control arrangements in your practice.



THE REGULATION AND QUALITY IMPROVEMENT AUTHORITY
9th floor Riverside Tower, 5 Lanyon Place, Belfast, BT1 3BT
Tel: 028 9051 7500 Fax: 028 9051 7501

1 Prevention of bloodborne virus exposure			
Inspection criteria <i>(Numbers in brackets reflect HTM 01-05/policy reference)</i>	Yes	No	If NO provide rationale and actions to be taken with timescales to achieve compliance with HTM 01-05.
1.1 Does the practice have a policy and procedure/s in place for the prevention and management of blood borne virus exposure, including management of spillages, sharps and inoculation incidents in accordance with national guidance? (2.6)	✓		
1.2 Have all staff received training in relation to the prevention and management of blood-borne virus exposure? (1.22, 9.1, 9.5)	✓		
1.3 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? (2.6)	✓		
1.4 Can decontamination and clinical staff demonstrate current immunisation with the hepatitis B vaccine e.g. documentation? (2.4s, 8.8)	✓		
1.5 Are chlorine-releasing agents available for blood /bodily fluid spillages and used as per manufacturer's instructions? (6.74)	✓		
1.6 Management of sharps Any references to sharps management should be read in conjunction with The Health and Safety (Sharp Instruments in Healthcare) Regulations (Northern Ireland) 2013 Are sharps containers correctly assembled?	✓		

<p>1.7 Are in-use sharps containers labelled with date, locality and a signature?</p>	<p>✓</p>		
<p>1.8 Are sharps containers replaced when filled to the indicator mark?</p>	<p>✓</p>		
<p>1.9 Are sharps containers locked with the integral lock when filled to the indicator mark? Then dated and signed?</p>	<p>✓</p>		
<p>1.10 Are full sharps containers stored in a secure facility away from public access?</p>	<p>✓</p>		
<p>1.11 Are sharps containers available at the point of use and positioned safely (e.g. wall mounted)?</p>	<p>✓</p>		
<p>1.12 Is there a readily-accessible protocol in place that ensures staff are dealt with in accordance with national guidance in the event of blood-borne virus exposure? (2.6)</p>	<p>✓</p>		
<p>1.13 Are inoculation injuries recorded?</p>	<p>✓</p>		
<p>1.14 Are disposable needles and disposable syringes discarded as a single unit?</p>	<p>✓</p>		
<p>Provider's level of compliance</p>			<p>Provider to complete</p>


2 Environmental design and cleaning			
Inspection criteria	Yes	No	If NO provide rationale and actions to be taken with timescales to achieve compliance with HTM 01-05.
2.1 Does the practice have a policy and procedure for cleaning and maintaining the environment? (2.6, 6.54)	✓		
2.2 Have staff undertaking cleaning duties been fully trained to undertake such duties? (6.55)	✓		
2.3 Is the overall appearance of the clinical and decontamination environment tidy and uncluttered? (5.6)	✓		
2.4 Is the dental chair cleaned between each patient? (6.46, 6.62)	✓		
2.5 Is the dental chair free from rips or tears? (6.62)	✓		
2.6 Are all surfaces i.e. walls, floors, ceilings, fixtures and fittings and chairs free from damage and abrasion? (6.38)	✓		
2.7 Are all work-surface joints intact, seamless, with no visible damage? (6.46, 6.47)	✓		
2.8 Are all surfaces i.e. walls, floors, ceilings, fixtures and fittings and chairs free from dust and visible dirt? (6.38)	✓		
2.9 Are the surfaces of accessible ventilation fittings/grills cleaned at a minimum weekly? (6.64)	✓		
2.10 Are all surfaces including flooring in clinical and decontamination areas impervious and easy to clean? (6.46, 6.64)	✓		

<p>2.11 Do all floor coverings in clinical and decontamination areas have covered edges that are sealed and impervious to moisture? (6.47)</p>		<p>✓</p>	<p>LDU flooring to be redone.</p>
<p>2.12 Are keyboard covers or "easy-clean" waterproof keyboards used in clinical areas? (6.66)</p>	<p>✓</p>		
<p>2.13 Are toys provided easily cleaned? (6.73)</p>	<p>✓</p>		
<p>2.14 Confirm free standing or ceiling mounted fans are not used in clinical/ decontamination areas? (6.40)</p>	<p>✓</p>		
<p>2.15 Is cleaning equipment colour-coded, in accordance with the National Patient Safety Agency recommendations as detailed in HTM 01-05? (6.53)</p>	<p>✓</p>		
<p>2.16 Is cleaning equipment stored in a non-clinical area? (6.60)</p>	<p>✓</p>		
<p>2.17 Where disposable single-use covers are used, are they discarded after each patient contact? (6.65)</p>	<p>✓</p>		
<p>2.18 Are the surfaces of equipment cleaned between each patient (E.g. work surfaces, dental chairs, curing lamps, delivery units, inspection handles and lights, spittoons, external surface of aspirator and X-ray heads)? (6.62)</p>	<p>✓</p>		
<p>2.19 Are all taps, drainage points, splash backs, sinks, aspirators, drains, spittoons, cleaned after every session with a surfactant/detergent? (6.63)</p>	<p>✓</p>		
<p>2.20 Are floors, cupboard doors and accessible high level surfaces and floors cleaned daily? (6.63)</p>	<p>✓</p>		

<p>2.21 Is there a designated area for the disposal of dirty water, which is outside the kitchen, clinical and decontamination areas; for example toilet, drain or slop-hopper (slop hopper is a device used for the disposal of liquid or solid waste)?</p>			
<p>2.22 Does the practice have a local policy and procedure/s for spillage in accordance with COSHH? (2.4d, 2.6)</p>			
<p>Provider's level of compliance</p>			<p>Provider to complete</p>

3 Hand hygiene			
Inspection criteria	Yes	No	If NO provide rationale and actions to be taken with timescales to achieve compliance with HTM 01-05.
3.1 Does the practice have a local policy and procedure for hand hygiene? (2.6 Appendix 1)	✓		
3.2 Is hand hygiene an integral part of staff induction? (6.3)	✓		
3.3 Is hand hygiene training provided periodically throughout the year? (1.22, 6.3)	✓		
3.4 Is hand hygiene carried out before and after every new patient contact? (Appendix 1)	✓		
3.5 Is hand hygiene performed before donning and following the removal of gloves? (6.4, Appendix 1)	✓		
3.6 Do all staff involved in any clinical and decontamination procedures have short nails that are clean and free from nail extensions and varnish? (6.8, 6.23, Appendix 1)	✓		
3.7 Do all clinical and decontamination staff remove wrist watches, wrist jewellery, rings with stones during clinical and decontamination procedures? (6.9, 6.22)	✓		
3.8 Are there laminated or wipe-clean posters promoting hand hygiene on display? (6.12)	✓		
3.9 Is there a separate dedicated hand basin provided for hand hygiene in each surgery where clinical practice takes place? (2.4g, 6.10)	✓		

<p>3.10 Is there a separate dedicated hand basin available in each room where the decontamination of equipment takes place? (2.4u, 5.7, 6.10)</p>	<p>✓</p>		
<p>3.11 Are wash-hand basins free from equipment and other utility items? (2.4g, 5.7)</p>	<p>✓</p>		
<p>3.12 Are hand hygiene facilities clean and intact (check sinks taps, splash backs, soap and paper towel dispensers)? (6.11, 6.63)</p>	<p>✓</p>		
<p>3.13 Do the hand washing basins provided in clinical and decontamination areas have :</p> <ul style="list-style-type: none"> • no plug; and • no overflow. <p>Lever operated or sensor operated taps.(6.10)</p>	<p>✓</p>		
<p>3.14 Confirm nailbrushes are not used at wash-hand basins? (Appendix 1)</p>	<p>✓</p>		
<p>3.15 Is there good quality, mild liquid soap dispensed from single-use cartridge or containers available at each wash-hand basin?</p> <p>Bar soap should not be used. (6.5, Appendix 1)</p>	<p>✓</p>		
<p>3.16 Is skin disinfectant rub/gel available at the point of care? (Appendix 1)</p>	<p>✓</p>		
<p>3.17 Are good quality disposable absorbent paper towels used at all wash-hand basins? (6.6, Appendix 1)</p>	<p>✓</p>		

<p>3.18 Are hand-cream dispensers with disposable cartridges available for all clinical and decontamination staff? (6.7, Appendix 1)</p>			
<p>Provider's level of compliance</p>			<p>Provider to complete</p>

4 Management of dental medical devices			
Inspection criteria	Yes	No	If NO provide rationale and actions to be taken with timescales to achieve compliance with HTM 01-05.
4.1 Does the practice have an infection control policy that includes procedures for the use, maintenance, service and repair of all medical devices? (1.18, 2.4a, 2.6, 2.7, 3.54)	✓		
4.2 Has the practice carried out a risk assessment for legionella under the Health and Safety Commission's "Legionnaires' disease - the control of legionella bacteria in water systems Approved Code of Practice and Guidance" (also known as L8)? (6.75-6.90, 19.0)	✓		
4.3 Has the practice a written scheme for prevention of legionella contamination in water pipes and other water lines?(6.75, 19.2)	✓		
4.4 Impression material, prosthetic and orthodontic appliances: Are impression materials, prosthetic and orthodontic appliances decontaminated in the surgery prior to despatch to laboratory in accordance with manufacturer's instructions?(7.0)	✓		
4.5 Impression material, prosthetic and orthodontic appliances: Are prosthetic and orthodontic appliances decontaminated before being placed in the patient's mouth? (7.1b)	✓		By Lab
4.6 Dental Unit Water lines (DUWLs): Are in-line filters cleaned/replaced as per manufacturer's instructions?(6.89, 6.90)	✓		

<p>4.7 Dental Unit Water lines (DUWLs): Is there an independent bottled-water system used to dispense distilled, reverse osmosis (RO) or sterile water to supply the DUWL? (6.84)</p>		✓	
<p>4.8 Dental Unit Water lines (DUWLs): For dental surgical procedures involving irrigation; is a separate single-use sterile water source used for irrigation? (6.91)</p>			N/A
<p>4.9 Dental Unit Water lines (DUWLs): Are the DUWLs drained down at the end of every working day?(6.82)</p>	✓		
<p>4.10 Dental Unit Water lines (DUWLs): Are self-contained water bottles (bottled water system) removed, flushed with distilled or RO water and left open to the air for drying on a daily basis, and if necessary overnight, and in accordance with manufacturer's guidance? (6.83)</p>	✓		
<p>4.11 Dental Unit Water lines (DUWLs): Where bottled water systems are not used is there a physical air gap separating dental unit waterlines from mains water systems. (Type A)?(6.84)</p>	✓		
<p>4.12 Dental Unit Water lines (DUWLs): Are DUWLs flushed for a minimum of 2 minutes at start of each working day and for a minimum of 20-30 seconds between every patient? (6.85)</p>	✓		
<p>4.13 Dental Unit Water lines (DUWLs): Are all DUWL and hand pieces fitted with anti-retraction valves? (6.87)</p>	✓		
<p>4.14 Dental Unit Water lines (DUWLs): Are DUWLs either disposable or purged using manufacturer's recommended disinfectants? (6.84-6.86)</p>	✓		

4.15 Dental Unit Water lines (DUWLs): Are DUWL filters changed according to the manufacturer's guidelines? (6.89)	✓		
Provider's level of compliance			Provider to complete



5 Personal Protective Equipment			
Inspection criteria	Yes	No	If NO provide rationale and actions to be taken with timescales to achieve compliance with HTM 01-05.
5.1 Does the practice have a policy and procedures for the use of personal protective equipment? (2.6, 6.13)	✓		
5.2 Are staff trained in the use of personal protective equipment as part of the practice induction? (6.13)	✓		
5.3 Are powder-free CE marked gloves used in the practice? (6.20)	✓		
5.4 Are alternatives to latex gloves available? (6.19, 6.20)	✓		
5.5 Are all single-use PPE disposed of after each episode of patient care? (6.21, 6.25, 6.36c)	✓		
5.6 Is hand hygiene performed before donning and following the removal of gloves? (6.4 Appendix 1)	✓		
5.7 Are clean, heavy duty household gloves available for domestic cleaning and decontamination procedures where necessary? (6.23)	✓		
5.8 Are heavy-duty household gloves washed with detergent and hot water and left to dry after each use? (6.23)	✓		
5.9 Are heavy-duty household gloves replaced weekly or more frequently if worn or torn? (6.23)	✓		

<p>5.10 Are disposable plastic aprons worn during all decontamination processes or clinical procedures where there is a risk that clothing/uniform may become contaminated? (6.14, 6.24-6.25)</p>	<p>✓</p>		
<p>5.11 Are single-use plastic aprons disposed of as clinical waste after each procedure? (6.25)</p>	<p>✓</p>		
<p>5.12 Are plastic aprons, goggles, masks or face shields used for any clinical and decontamination procedures where there is a danger of splashes? (6.14, 6.26-6.29)</p>	<p>✓</p>		
<p>5.13 Are masks disposed of as clinical waste after each use? (6.27, 6.36)</p>	<p>✓</p>		
<p>5.14 Are all items of PPE stored in accordance with manufacturers' instructions? (6.14)</p>	<p>✓</p>		
<p>5.15 Are uniforms worn by all staff changed at the end of each day and when visibly contaminated? (6.34)</p>	<p>✓</p>		
<p>5.16 Is eye protection for staff used during decontamination procedures cleaned after each session or sooner if visibly contaminated? (6.29)</p>	<p>✓</p>		
<p>5.17 Is eye protection provided for the patient and staff decontaminated after each episode of patient care? (6.29)</p>	<p>✓</p>		
<p>Provider's level of compliance</p>			<p>Provider to complete</p>

6 Waste			
Inspection criteria	Yes	No	If NO provide rationale and actions to be taken with timescales to achieve compliance with HTM 07-01.
6.1 Does the practice have a policy and procedure/s for the management and disposal of waste? (2.6, 6.1 (07-01) 6.4 (07-01))	✓		
6.2 Have all staff attended induction and on-going training in the process of waste disposal? (1.22, 6.43 (07-01) 6.51 (07-01))	✓		
6.3 Is there evidence that the waste contractor is a registered waste carrier? (6.87 (07-01) 6.90 (07-01))	✓		
6.4 Are all disposable PPE disposed of as clinical waste? (6.26, 6.27, 6.36, HTM 07-01 PEL (13) 14)	✓		
6.5 Are orange bags used for infectious Category B waste such as blooded swabs and blood contaminated gloves? (HTM 07-01, PEL (13) 14, 5.39 (07-01) Chapter 10 - Dental 12 (07-01))	✓		
6.6 Are black/orange bags used for offensive/hygiene waste such as non-infectious recognisable healthcare waste e.g. gowns, tissues, non-contaminated gloves, X-ray film, etc, which are not contaminated with saliva, blood, medicines, chemicals or amalgam? (HTM 07-01, PEL (13) 14, 5.50 (07-01) Chapter 10-Dental 8 (07-01))	✓		
6.8 Are black/clear bags used for domestic waste including paper towels? (HTM 07-01, PEL (13) 14, 5.51 (07-01))	✓		

<p>6.9 Are bins foot operated or sensor controlled, lidded and in good working order? (5.90 (07-01))</p>	<p>✓</p>		
<p>6.10 Are local anaesthetic cartridges and other Prescription Only Medicines (POMs) disposed of in yellow containers with a purple lid that conforms to BS 7320 (1990)/UN 3291? (HTM 07-01 PEL (13) 14, Chapter 10 - Dental 11 (07-01))</p>	<p>✓</p>		
<p>6.11 Are clinical waste sacks securely tied and sharps containers locked before disposal? (5.87 (07-01))</p>	<p>✓</p>		
<p>6.12 Are all clinical waste bags and sharps containers labelled before disposal? (5.23 (07-01), 5.25 (07-01))</p>	<p>✓</p>		
<p>6.13 Is waste awaiting collection stored in a safe and secure location away from the public within the practice premises? (5.33 (07-01), 5.96 (07-01))</p>	<p>✓</p>		
<p>6.14 Are all clinical waste bags fully described using the appropriate European Waste Catalogue (EWC) Codes as listed in HTM 07-01 (Safe Management of Healthcare Waste)?(3.32 (07-01))</p>	<p>✓</p>		
<p>6.15 Are all consignment notes for all hazardous waste retained for at least 3 years?(6.105 (07-01))</p>	<p>✓</p>		
<p>6.16 Has the practice been assured that a "duty of care" audit has been undertaken and recorded from producer to final disposal? (6.1 (07-01), 6.9 (07-01))</p>	<p>✓</p>		
<p>6.17 Is there evidence the practice is segregating waste in accordance with HTM 07-01? (5.86 (07-01), 5.88 (07-01), 4.18 (07-01))</p>	<p>✓</p>		
<p>Provider's level of compliance</p>			<p>Provider to complete</p>

7 Decontamination			
Inspection criteria	Yes	No	If NO provide rationale and actions to be taken with timescales to achieve compliance with HTM 01-05.
7.1 Does the practice have a room separate from the patient treatment area, dedicated to decontamination meeting best practice standards? (5.3–5.8)	✓		
7.2 Does the practice have washer disinfector(s) in sufficient numbers to meet the practice requirements? (PEL(13)13)	✓		
7.3 Are all reusable instruments being disinfected using the washer disinfector? (PEL(13)13)	✓		
7.4 Does the practice have steam sterilisers in sufficient numbers to meet the practice requirements?	✓		
7.5 a Has all equipment used in the decontamination process been validated? 7.5 b Are arrangements in place to ensure that all equipment is validated annually? (1.9, 11.1, 11.6, 12,13, 14.1, 14.2, 15.6)	✓		
7.6 Have separate log books been established for each piece of equipment? Does the log book contain all relevant information as outlined in HTM01-05? (11.9)	✓		

<p>7.7 a Are daily, weekly, monthly periodic tests undertaken and recorded in the log books as outlined in HTM 01-05? (12, 13, 14)</p>			
<p>7.7 b Is there a system in place to record cycle parameters of equipment such as a data logger?</p>			
<p>Provider's level of compliance</p>			<p>Provider to complete</p>

<p>Please provide any comments you wish to add regarding good practice</p>
Empty space for comments

Appendix 1



Name of practice: **Quinndental**

Declaration on consultation with patients

The need for consultation with patients is outlined in The Independent Health Care Regulations (Northern Ireland) 2005, Regulation 17(3) and The Minimum Standards for Dental Care and Treatment 2011, Standard 9.

1 Do you have a system in place for consultation with patients, undertaken at appropriate intervals?

Yes No

If no or other please give details:

2 If appropriate has the feedback provided by patients been used by the service to improve?

Yes No

3 Are the results of the consultation made available to patients?

Yes No