



Nevada Board of Dental Examiners 2651 N Green Valley Parkway, Suite 104 Henderson, NV 89014 (702) 486-7044 • (800) DDS-EXAM • Fax (702) 486-7046

INFECTION CONTROL INSPECTION/SURVEY FORM						Rev 09/2023		
Dent	Dental Office Name: Date of Inspection:							
Licen	see/Owner Name:				1			
Addr	ess:				INSPECTOR(S)			
				(1)	(2)			_
City:		State:	Zip Code:		PURPOSE OF INSPEC	TION		
		Nevada		Initial Inspe	ction: Re-Inspection:	Random I	nspect	ion: 🔲
		СО	MPLIANCE LE	VEL CRITERIA	– LEVEL # 1-4			
Leve	l # 1 - CRITICAL: Failure to me	et these standard	ds may result in I	MMEDIATE term	nination of patient care at the tim	e of inspecti	on.	
Leve		JIRED: Failure to	meet these stan	dards will requir	e corrective action to become co	mpliant with	in 7 days	from the
Leve	date of inspection. I # 3 - ACTION REQUIRED: Faile	ure to meet these	standards will r	equire corrective	e action to become compliant wit	hin 30 davs f	rom the	date of
	inspection.							
Leve					required for compliance at this tiendations are updated.	me – complia	nce req	uirments
REC	are subject to change as the Centeres for Deisease Control (CDC) recommendations are updated. RECORD KEEPING — EACH PRACTICE MUST HAVE LEVEL 1.4 Y N							
1	Is there a written infection control program that is specific for the owner of this location and easily							N
EDU	CATION & TRAINING							
2	Is there documentation of r best practices?	eview of the infe	ection control pla	an at least annua	ally to ensure compliance with	3	Υ	N
3								N
4	4 Is there documentation of education and training that is appropriate to each dental personnel/staff member including hands-on training for personnel that process semi critical and critical instruments. 3 Y N							N
5	Are training records kept f			roceso genin erit	treat and enteed metronicites.	3	Υ	N
Is there written policies and procedures for corrective action for any deviation from the written policy including documentation of any corrective actions taken?						N		
CON	CONFIDENTIAL VACCINATION RECORDS, EXPOSURE AND POST EXPOSURE MANAGEMENT, MEDICAL CONDITIONS, WORK RELATED							_ATED
ILLN	ESS AND WORK RESTRICTION		11 /1	66 221	1 1 1 10		1	
7	that may expose others to infection?							N
8								N
	immunizations? Does the office maintain a confidental employee health record that includes any exposure and post							
9	exposure care recieved?							N
10	exposure medical plan (e.g. use CDC needle stick/sharps injury/exposure protocol?)							
11	Is a 24/7 contact telephone number for a qualified healthcare provider to handle occupational/post							N
12	Are there exposure and in			ıg a sharps injur	ry log?	3	Υ	N
13					e communicable disease upon	3	Υ	N
BLO	BLOODBORNE PATHOGEN ELEMENTS							

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14	Is there written policies and procedures for the prevention and transmission of bloodborne pathogens?	3	Υ	N
15	Is there written policies and procedures for hand hygiene, including documentation of training and appropriate selection of antiseptic agents?	3	Υ	N
16	Is there written policies and procedures for proper use of personal protective equipment?	3	Υ	N
17	Is there written policies and procedures for handling and management of sharps?	3	Υ	N

DISI	NFECTION AND STERILIZATION OF PATIENT CARE ITEMS						
18	Is there writen policies and procedures for managing semi-critical and critical items?		3	Υ		N	
19	Is there a written system outlining the entire sterilization process (written policies and procedures transporting and processing of all contaminated critical and semi-critical instruments, the instruments processing area, preparation and packaging of instruments, sterilization and storage of sterilized a clean dental instruments?)	ent	3	Y		N	
20	Is there written policy and procedures for sterilization biologic monitoring including how to handle failed biologic monitoring test?		3	Y		N	
21	Are weekly biological monitoring logs kept for each sterilizer that include the machine tested, date was sent, date test results were returned and the results of testing?	e test	1	Υ		N	
22	Are weekly biological monitoring logs kept for 2+ years or since opening date:?		3	Υ		N	
23	Are appropriate testing and maintenance logs kept for each piece of equipment such as sterilizers ultrasonic cleaners, etc?	,	3	Υ		N	
ENV	IRONMENTAL INFECTION CONTROL ELEMENTS						
24	Is there written policies and procedures for aseptic management during patient care?		3	Υ		N	
25	Is there written policies and procedures for surface disinfection and environmental barrier protect	ion?	3	Υ		N	
26	Is there written policies and procedures for medical waste management?		3	Υ		N	
27	Is the name/telephone number of licensed waste hauler for regulated waste available?		3	Υ		N	
28	Is there written Policies and procedures for decontaminating spills of blood or other body fluids w necessary supplies present for decontamination (i.e. Blood Spill Kit?)	ith	3	Υ		N	
29	Are there written policies and procedures for meeting EPA potable water standard and treating biofilm, including treating, testing and re-testing water lines?		2	Υ		N	
30	Are dental unit water lines flushed for 2 minutes each day prior to use and in between patients for minimum of 20 seconds?	ra	2	Υ		N	
31	Is documentation kept for dental unit water line testing to meet the potable water standard of EP (<500 CFU/ml?)		2	Υ		N	
32	Is there written policies and procedures to maintain aesepis and prevent cross contamination who taking and processing dental radiographs?	3	Υ	N	N/A		
33	Is there written policies and procedures to maintain asepsis and prevent cross contamination duri dental laboratory procedures?	3	Υ	N	N/A		
ОТН	ER						
34	Is a comprehensive and annually updated medical histroy form is used to evaluate patients?		3	Υ		N	
CON	MMUNICABLE DISEASE CONTROL PROCEDURES	LEVEL 1-4	Υ	N	١	I/A	
35	Are items used single use or sterilized for critical items?	1	Υ	N			
36	Are multi-dose vials used?		Υ	N			
37	a) If yes, are vials only entered with new sterile syringe with a new sterile needle?	1	Υ	N N/A			
38	b) Is the cap of multi-dose vial cleaned with alcohol based wipe before being accessed?	2	Υ	N	N	I/A	
39	c) Is the date of first access of a multi-use vials documented and discarded when expired or 28 days after initial access (as applicable?)	Υ	N	N	I/A		
40	Are fluid infusion and administration sets (IV bags, tubing and connectors) used?		Y	N			
41	a) If yes, are thery used only on one patient then discarded?	1	Υ	N N/A			
42	c) Is a single IV bag used to mix medications for ONE patient only?	1	Υ	N N/A			
43	d) Are single dose medication/infusions used for only ONE patient and discarded after use?	1	Υ	N N/A		/A	
44	Is there written policies and procedures and supplies available for personnel to wear utility gloves when processing contaminated instruments (not patient care type of gloves?)	2	Y N				
45	Are there supplies for hand hygiene accessible to employees at point of need (soap, water, alcohol rub if used?)	2	. Y N				

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46	Are team members adequately trained and able to demonstrate appropriate hand hygiene	2	V	N	
40	techniques?		I	IN	

APPRO	OPRIATE PPE SUPPLIES ACCESSIBLE & EMPLOYEES WITH EXPOSURE RISKS				
	Are gloves available in appropriate sizes (Latex and latex free or just latex free) including				
47	appropriately sized sterile surgical gloves if surgeries are performed in the office?	1	Υ	N	
48	Is the level of masks appropriate to the procedure type performed in the office?	1	Υ	N	
49	Are safety glasses with side shields and/or full-face shields used in conjunction with safety glasses available for use?	1	Υ	N	
50	Are disposable and/or laundered gowns available for use in the office?	1	Υ	N	
51	Do health care workers display appropriate use of PPE barriers?	2	Υ	N	
52	Is there a running water eyewash station accessible?	3	Υ	N	
53	Are appropriate barrier products available for patient use during procedures (dental dams, protective eyewear, etc?)	2	Υ	N	
54	Are basic first aid products and equipment available (Recommended to include: Nitrogylerin, Benadryl, Epinephrine Auto Injector for adult and child if applicaple, Oxygen, Aspirin, Albuterol, Glucose, etc?)	2	Υ	N	
CLEA	INING, DISINFECTION & STERILIZATION OF PATIENT CARE ITEMS	1		1	
55	Are biofilm and organic matter removed from critical and semi-critical instruments using detergents or enzymatic cleaners prior to sterilization following manufacture recommendations that may require temperature and time?	2	Υ	N	
56	Is sterilization equipment available and fully functional?	1	Υ	N	
57	What are the number of working autoclaves?	1	Υ	N	N/A
58	What are the number of working chemiclaves?	1	Υ	N	N/A
59	What are the number of working dry heat sterilizers?	1	Υ	N	N/A
60	What are the number of working Flash steam sterilizers (Statim)?	1	Υ	N	N/A
61	What are the number of working ultrasonic cleaners?	1	Υ	N	
62	Is biological testing of sterilizer(s) completed weekly on each cycle used (pouched, plastics, solids, etc) and with a full bio burden load under normal processiong parameters (full load of instruments, not overloaded, spore test strip or vial in a pouch according to manufacture recommendations?)	1	Υ	N	
63	Is a mail in biological testing service used? If yes, name:		Υ	N	N/A
64	Is in-office biological testing used and is control processed for each test?	2	Υ	N	N/A
65	Are sterilization cycles verified with chemical/heat indicators including a class V integrator for closed cassettes and containers?	2	Υ	N	
66	Are critical items (any instrument that penetrates soft tissue or bone) sterilized after each use?	1	Υ	N	
67	Is there written policies and procedures for proper sterilzer loading techniques and is demonstrated by the staff?	2	Y	N	
68	Are heat tolerant handpieces sterilized after each use (including high & low speed handpieces, prophylaxis angles and motors, ultrasonic and sonic handpiece and scaling tips, air abrasion devices, air and water syringe tips, and motorswith exception of electric type models?)	1	Υ	N	
69	Is event-related monitoring used to monitor package integrity, reprocessed when compromised and appropriately stored with a minimum of an initial date stamp and sterilizer used (if more than one sterilizer present?)	2	Υ	N	
70	Are single use items, supplies or devices and items labeled with $^{\textcircled{3}}$ discarded after use and not reprocessed?	1	Y	N	
71	Are semi-critical items sterilized after each use if not heat sensitive?	1	Υ	N	
72	Are heat sensitive semi-critical items processed at a minimum of high level disinfection or chemical sterilization after each use?	1	Υ	N	
73	Are semi-critical items that are not heat or chemical tolerant, such as digital sensors, intraoral cameras, intraoral scanners, curing lights, etc., utilizing FDA approved barriers and are cleaned then disinfected with an intermediate level disinfection agent between patients?	1	Υ	N	
74	Is the practice using an FDA approved chemical sterilant and has policies and procedures in place to ensure adequate exposure time is reached?	2	Υ	N	N/A

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75	Are all applicable label instructions followed on FDA approved chemical sterilant including mixing, dilution, expiration date, shelf life, storage, safe use, disposal and material compatibility?	Υ	N	N/A			
Asep	otic Techniques:						
76	Are splash shields and equipment guards used on dental laboratory lathes and grinders?	4	Υ	N	N/A		
77	Is fresh pumice and a sterilized or new rag wheel used for each patient?	2	Υ	N	N/A		
78	Are devices used to polish, trim or adjust contaminated intraoral devices disinfected and/or sterilized between patients?	2	Υ	N	N/A		
79	Are intraoral items such as impressions, bite registrations, prosthetics, crown and bridge, and orthodontic appliances cleaned and disinfected before lab procedures and before delivering to the patient?	2	Υ	N	N/A		
Envi	ronmental Infection Control		LEVEL 1-4	Υ	N		
80	Are clinical contact surfaces (frequently touched surface that could potentially allow secondary transmission to the DHCW or patient) that are not barrier protected cleaned then disinfected using an registered hospital disinfectant with low to intermediate claim after each patient following manufacturecommendations? Intermediate level disinfectant (TB claim) to be used if visibly contaminated with leading to the contaminated	re	2	Υ	N		
81	Are housekeeping surfaces (sinks, floors, walls, drawers, supply containers, etc.) cleaned on a routine basis?						
82	Are environmental surfaces cleaned then disinfected with an EPA registered low to intermediate level disinfectant at beginning and end of day?						
83	shelf life, storage, use of material compatibility?)						
84	Are all clinical contact surfaces protected with barriers (especially areas that are difficult to clean?)		2	Υ	N		
85	Are barriers removed then surfaces cleaned then disinfected prior to applying new barrier in between patients?	2	Υ	N			
86	Are the decontamination and clean areas adequately separated in the instrument processing area?	2	Υ	N			
87	Is biohazardous waste stored properly?		2	Υ	N		
Shar	ps						
88	Are approved sharps containers utilized, accessible and secured to counter/wall?		2	Υ	N		
89	Are sharps containers taken out of service when full and processed appropriately?	2	Υ	Ν			
90	Are safe recapping techniques/devices used and is the technique demonstrated by the staff?	2	Υ	N			
91	Single use sharps (blades, neeldes, sutures, etc) are disposed of after use?		1	Υ	N		
92	Do employees use engineering controls (e.g., forceps, hemostat, etc) to retrieve contaminated sharps from syringe, handles, trays or containers?		2	Υ	N		

ACKNOWLEDGEMENT AND RECEIPT OF COPY BY OWNER/AUTHORIZED AGENT

- 1. The owner of the dental practice hereby acknowledges that by executing this document below and initialing each page's lower right-hand corner on the line "Licensee Initials," receipt of a copy of this inspection/survey form is acknowledged.
- 2. In the event the dental practice has satisfactorily completed the inspection, as noted in this inspection/survey form, the owner/licensee will receive from the Board's Executive Director and/or representative, written notice of satisfactorily completing the inspection conducted.
- 3. If an owner/licensee has commenced the practice of dentistry prior to an Initial Inspection (NAC 631.1785) at any given location that inspection shall be deemed to be a Random Inspection pursuant to NAC 631.179.
- 4. If the inspection indicates level 1 deficiencies, the owner/licensee will receive written notice from the Board's Executive Director and/or representative of the deficiencies and a re-inspection will be conducted within seventy-two (72) hours of the written notice. In the event the deficiencies pose an immediate threat to the safety and/or welfare of the public, the President of the Board, may without any further action of the Board, issue an Order of Summary Suspension pursuant to NAC 631.179(4).
- 5. In the event the inspection indicates level 2 deficiencies, the owner/licensee will receive written notice from the Board's Executive Director and/or representative of the deficiencies and a re-inspection will be conducted within seven (7) days of the written notice.
- 6. In the event the inspection indicates "action required" deficiencies (items listed with a "#3"), the owner/licensee will receive written notice from the Board's Executive Director and/or representative of the "action required" deficiencies and that a re-inspection will be conducted within thirty (30) days of the written notice.

Receip	t of a copy of the fo	regoing is her	eby ack	nowled	lged;	
Ву				_		Print name:
this	day of	, 20_	at	:	m.	Title and/or position/capacity: