

OPR/Electrology

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Complete

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Disclaimer & Confidentiality

Disclaimer

The Inspector believes the information contained within this report to be correct at the time of the inspection. The Inspector does not accept responsibility for any consequences arising from the use of the information herein, outside of official OPR sanctioned actions. The report is based on matters which were observed at the time of the inspection and is not an exhaustive record of all possible risks or hazards that may exist, or potential improvements that can be made.

Information on the latest Rules and Regulations can be found at the Vermont Secretary of State Office of Professional Regulation home page.

All inspection related correspondence concerning this report shall be addressed to:

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Confidentiality Statement

In order to maintain the integrity and credibility of the risk assessment process, and to protect the parties involved, it is understood that the Inspector will not divulge to unauthorized persons any information obtained during this inspection unless legally obligated to do so.

GENERAL INSPECTION INFORMATION

Store Name, Address & Telephone Number.	
Unanswered	
Shop license number(s).	
Unanswered	
Owner	
Unanswered	
Is the owner the manager of the establishment?	Unanswered
Designated Registrant. / Title 26 V.S.A. § 4105(c)(1)	
Unanswered	
Shop Hours.	
Unanswered	
Hours:	Unanswered
Website:	
Unanswered	
Email Address:	
Unanswered	
Date of Inspection.	
Unanswered	
Reason for Inspection.	New Shop

PERSONNEL

Do any personnel maintain a special license endorsement for laser hair removal? / Title 26 V.S.A. § 4404(d)	Unanswered
Are licenses / registrations displayed in a conspicuous manner? / Rule 4.2	Unanswered
All licenses active?	Unanswered
Other personnel work in shop?	Unanswered

GENERAL FACILITY STANDARDS

Treatment rooms separate from waiting room(s). / Rule 4.3(A)(1)	Unanswered
Each treatment room floor plan contains 48 square feet. / Rule 4.3(A) (2)	Unanswered

Each treatment room contains a sink with hot/cold running water. / Rule 4.3(A)(3)	Unanswered
Treatment rooms do not incorporate any residential functions. / Rule 4.3(A)(8)	Unanswered
Adequate lighting. / Rule 4.3(A)(4)	Unanswered
Professional lamp focused on treatment area at all times. / Rule 4.3(C)	Unanswered
Counter tops are smooth, non-porous material. / Rule 5.16(B)	Unanswered
Adequate ventilation. / Rule 4.3(A)(5)	Unanswered
Is smoking prohibited in the treatment room? / Rule 4.3(E)	Unanswered
Sanitary conditions. / Rule 4.3(A)(6)	Unanswered
Toilet facilities available for consumers. / Rule 4.3(A)(7)	Unanswered

GENERAL EQUIPMENT STANDARDS

Professional type forceps. / Rule 4.3(D)	Unanswered
Forceps/instruments accumulated in a covered holding container. / Rule 5.5(B)(1)	Unanswered
Holding container equipped with protein-dissolving enzyme detergent. / Rule 5.5(B)(1)	Unanswered
Packaged instruments from autoclave or dry heat sterilizer show chemical indicator. / Rule 5.5(B)(6)	Unanswered
Sterilized, packaged instruments stored in a clean, dry, covered container. / Rule 5.5(B)(7)	Unanswered
Anaphoresis/cataphoresis rollers sterilized in the same manner as forceps. / Rule 5.5(F)	Unanswered
Tips for epilator needle holders are accumulated in a covered holding container. / Rule 5.5(E)(1)	Unanswered
Holding container equipped with protein-dissolving enzyme detergent. / Rule 5.5(E)(1)	Unanswered
Packaged tips are:	
Mechanically sterilized. / Rule 5.5(E)(5)	
Submerged in 1:99 bleach/water solution. / Rule 5.5(E)(5)	
Sterilized/bleached, packaged tips stored in a clean, dry, covered container. / Rule 5.5(E)(6)	Unanswered

Single-use, pre-sterilized, and disposable needles. / Rule 4.6(A)	Unanswered
Needles stored in a manner that maintains sterile condition. / Rule 5.5(A)(2)	Unanswered
Sharps container. / Rule 5.5(A)(4)	Unanswered
Sharps container securely sealed? / Rule 5.5(A)(5)	Unanswered
Method of sharps removal by medical waste removal companies: / Rule 5.5(A)(5)	Unanswered
Non-sterile, medical grade, disposable examination gloves. / Rule 4.6(C)	Unanswered
Disposable paper towels. / Rule 5.1(B)(4)-(5)	Unanswered
Soap for hand washing. / Rule 5.1(B)(1)	Unanswered
Alcohol based water-less handrub. / Rule 5.1(C)	Unanswered
FDA approved antiseptic. / Rules 4.5(A)(2), 5.18(A)-(B)	Unanswered
EPA approved Low-Level Disinfectant. / Rules 4.5(A)(27), 5.14(E)	Unanswered
EPA approved hospital-grade disinfectant/germicide. / Rules 4.5(A) (20), 5.15(A)-(B), 5.16(A)	Unanswered
Disposal protective barrier material. / Rule 5.14(F)	Unanswered
Fresh, clean drapes for treatment table/chair. / Rule 5.14(A)	Unanswered
Trash can with disposable plastic bag liner. / Rule 5.14(C)	Unanswered
Blood spill kit available? / Rule 5.5(K)	Unanswered

STERILIZATION EQUIPMENT

Is a dry heat sterilizer or autoclave in use? / Rule 5.5(I)	Unanswered
Is spore testing being conducted on a monthly basis? / Rule 5.5(J)	Unanswered
Date of last spore test? Unanswered	
Testing results filed in a permanent Sterility Assurance file? / Rule 5.5(J), 4.5(A)(38)	Unanswered

DOCUMENTATION

Disclosure forms on hand? / Rule 4.1(A)	Yes
Does the disclosure form include:	

The nature of the treatment or procedure to be performed. / Rule 4.1(A)(1)	
The potential benefits and risks of undergoing said treatment. / Rule 4.1(A)(2)	
The nature of any after-treatment care to be provided. / Rule 4.1(A)(3)	
The cost of said treatment. / Rule 4.1(A)(4)	
Other information reasonably necessary to allow the patient to make a decision intelligently about whether or not to undergo electrology treatment. / Rule 4.1(A)(5)	
Consent forms on hand? / Rule 4.1(B)	Unanswered
Consent forms signed by patient(s)? / Rule 4.1(B)	Unanswered
Health assessment forms on hand? / Rule 5.17(B)	Unanswered
Completed health assessment forms on hand?	Unanswered

PUNCTURE INJURY PROTOCOL

Has an employee of the facility been exposed to a puncture incident?	Unanswered
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LASER DEVICE

Is a Laser device currently in use? / Title 26 V.S.A. § 4404(d), Rule 2.3	Unanswered
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NOTES REGARDING INSPECTION / DEFICIENCIES FOUND

NOTES:	
Unanswered	
DEFICIENCIES FOUND:	
Unanswered	

SIGNATURES:

Inspector Signature:

Unanswered

Reviewing Designated Registrant Email Address

Unanswered

By signing below, the reviewing Designated Registrant acknowledges he/she has reviewed this inspection report with Inspector and any deficiencies contained herein.

Registrant Signature:

Unanswered

FINDING OF NO DEFICIENCIES: This report will serve as notice of satisfactorily closing the current inspection process.

DEFICIENCIES FOUND: Within the next ten days, please provide response correspondence outlining the corrective measures addressing the discrepancies outlined within the report. This will be added to the inspection record.