

Medical Commission

Licensing. Accountability. Leadership.

Rules Workshop

Chapter 246-919 WAC

March 6, 2019

Capital Event Center 6005 Tyee Drive SW Tumwater, WA 98512 (360) 464-6700



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Rule Workshop Notice



Public Notification and Meeting Announcement For the Washington Medical Commission *CR-101 for Chapter 246-919 WAC

Rulemaking

The Washington Medical Commission (Commission) has officially filed a <u>CR-101</u> with the Office of the Code Reviser on January 2, 2018. The Commission is considering amending the chapter to update and modernize the rules to align with current law and practice. The WSR# is 18-02-079.

Several sections within chapter 246-919 WAC have not been updated since 1996. Since then, several changes have occurred that may require rule revisions. The commission is considering updating rules to more closely align with current industry standards and provide clearer rules language for licensed allopathic physicians. In addition, RCW 43.70.041 requires the commission to review its administrative rules every five years to ensure that regulations are current and relevant. Rule amendments being considered will potentially benefit the public's health by ensuring participating providers are informed and regulated by current national industry and best practice standards.

Chapter 246-919 WAC Rules Committee Meeting

In response to the filing, the Commission will conduct their second open public rules workshop and committee meeting on Wednesday, January 16, 2019, beginning at 2:00 pm, at the:

Capital Event Center Chehalis B Room 6005 Tyee Drive SW Tumwater, WA 98512 (360) 464-6700

This meeting will be open to the public.

Approved 11/16/2018 Updated: February 25, 2019

The purpose of the rules workshop will be to discuss draft language and proposed changes to the current chapter.

Interested parties, stakeholders, and the general public are invited to participate in the rules workshops or provide comments on draft rules. For continued updates on rule development, interested parties are encouraged to join the Commission's rules GovDelivery.

For more information, please contact Amelia Boyd, Program Manager, Washington Medical Commission at (360) 236-2727 or by email at amelia.boyd@wmc.wa.gov.

Attachment: CR-101

Approved 11/16/2018 Updated: February 25, 2019

^{*}CR means Code Reviser



Rules Workshops Agenda March 6, 2019

> Capital Event Center 6005 Tyee Drive SW Tumwater, WA 98512

> > (360) 464-6700

Wednesday, March 6, 2019

Open Meeting

1:00 pm Clinical Support Program – WSR #18-06-007

Chehalis B

Discussion of draft and proposed rules

2:00 pm Chapter 246-919 WAC – WSR #18-02-079

Chehalis B

Discussion of draft and proposed rules

In accordance with the Open Public Meetings Act, this meeting notice was sent to individuals requesting notification of the Department of Health, Medical Quality Assurance Commission meetings.

Times and Order: The meeting will commence at 1:00 pm on Wednesday, March 6, 2019.



This agenda is subject to change.

Accessibility: These meetings are accessible to persons with disabilities. Special aids and services can be made available upon advance request. Advance request for special aids and services must be made no later than five days prior to the meeting. If you wish general information about this meeting, please call the program at (360) 236-2727. If you need assistance with special needs and services, you may leave a message with that request at 1-800-525-0127 or, if calling from outside Washington State, call (360) 236-4052. TTY users dial 711 for Washington State Relay Service. If you need assistance due to a speech disability, Speech-to-Speech provides human voices for people with difficulty being understood. The Washington State Speech-to-Speech toll free access number is 1-877-833-6341.

Smoking is prohibited at these meetings.



CR-101

WSR 18-02-079 PREPROPOSAL STATEMENT OF INQUIRY DEPARTMENT OF HEALTH

(Medical Quality Assurance Commission)
[Filed January 2, 2018, 12:29 p.m.]

Subject of Possible Rule Making: Chapter 246-919 WAC, Medical quality assurance commission (allopathic physicians), the medical quality assurance commission (commission) is considering amending the chapter to update and modernize the rules to align with current law and practice.

Statutes Authorizing the Agency to Adopt Rules on this Subject: RCW 18.71.017, 18.130.050.

Reasons Why Rules on this Subject may be Needed and What They Might Accomplish: Several sections within chapter 246-919 WAC have not been updated since 1996. Since then, several changes have occurred that may require rule revisions. The commission is considering updating rules to more closely align with current industry standards and provide clearer rules language for licensed allopathic physicians. In addition, RCW 43.70.041 requires the commission to review its administrative rules every five years to ensure that regulations are current and relevant. Rule amendments being considered will potentially benefit the public's health by ensuring participating providers are informed and regulated by current national industry and best practice standards.

Other Federal and State Agencies that Regulate this Subject and the Process Coordinating the Rule with These Agencies: None.

Process for Developing New Rule: Collaborative rule making.

Interested parties can participate in the decision to adopt the new rule and formulation of the proposed rule before publication by contacting Daidria Amelia Underwood, P.O. Box 47866, Olympia, WA 98504-7866, phone 360-236-2727, fax 360-236-2795, TTY 360-833-6388 or 711, email daidria.underwood@doh.wa.gov.

Additional comments: Interested persons may sign up for the commission's interested parties list (GovDelivery) at https://public.govdelivery.com/accounts/WADOH/subscriber/new to participate in the commission's rule drafting workshops. All commission rule-making notices will be emailed via GovDelivery and interested parties will be invited to participate in public rules meetings and submit written comments for consideration.

January 2, 2018 Melanie de Leon Executive Director



Proposed Changes

Chapter 246-919 WAC

MEDICAL QUALITY ASSURANCE COMMISSION

Last Update: 9/6/17

WAC 246-919-010 Definitions. The definitions in this section apply throughout this chapter unless the context clearly requires otherwise.

- (1) "Applicant" is an individual who has completed the application form and has paid the application fee.
- (2) "Commission" means the Washington state medical quality assurance commission.
 - (3) "Emergent" means a circumstance calling for immediate action.
- (4) "Hospital" means any health care institution licensed pursuant under to chapter 70.41 RCW.
- (5) "Intermittent" means providing services on a part-time or fulltime nonpermanent basis.
- (6) "Mentally or physically disabled physician" means a physician who has either been determined by a court to be mentally incompetent or mentally ill or who is unable to practice medicine with reasonable skill and safety by reason of any mental or physical condition.

- (7) "Nursing home" means any health care institution which comes chapter 18.51 RCW.
- (8) "Physician" means a physician person licensed pursuant tounder chapter 18.71 RCW.
- (9) "Unprofessional conduct" as used in these regulations shall means the conduct described in RCW 18.71.0193 for conduct occurring before June 11, 1986, and the conduct described in RCW 18.130.180 for conduct occurring on or after June 11, 1986.

[Statutory Authority: RCW 18.71.017, 18.130.250, 18.71.440. WSR 11-05-025, § 246-919-010, filed 2/7/11, effective 3/10/11. Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, \$ 246-919-010, filed 1/17/96, effective 2/17/96.]

WAC 246-919-020 Commission address. The commission's official mailing address is:

Medical Quality Assurance Commission

Department of Health

P.O. Box 47866

Olympia, WA 98504-7866

[Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, § 246-919-020, filed 1/17/96, effective 2/17/96.]

WAC (2/8/2018 11:51 AM) [2]

NOT FOR FILING

Commented [UD(1]: Approved.

WAC 246-919-110 Commission meetings. Regular commission meetings shall be held at least four times yearly. Additional regular or special meetings may be called at the discretion of the chair or by a quorum of the commission.

[Statutory Authority: RCW 18.71.017. WSR 04-04-067, § 246-919-110, filed 2/2/04, effective 3/4/04. Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, § 246-919-110, filed 1/17/96, effective 2/17/96.]

APPLICATIONS AND EXAMINATIONS

WAC 246-919-300 Application withdrawals. An application for a license may not be withdrawn after the commission or the reviewing commission member determines that grounds exist for denial of the license or for the issuance of a conditional license. Applications which that are subject to investigation for unprofessional conduct or impaired practice may not be withdrawn.

[Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, § 246-919-300, filed 1/17/96, effective 2/17/96.]

WAC 246-919-310 Credentialing of physicians and surgeons. All completed applications, for either limited or full licensure, must be reviewed by a member of the commission or a designee authorized in writing by the commission prior to examination and/or licensure.

[Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, \$246-919-310, filed 1/17/96, effective 2/17/96.]

Commented [UD(2]: Repeal. Not required to use this process.

wac 246-919-320 Approved United States and Canadian medical schools. For the purposes of the Medical Practice Act RCW 18.71.055, the commission approves those medical schools accredited by the Liaison Committee on Medical Education.

Commented [FM(3]: RCW 18.71 is not officially called the Medical Practice Act. This is the informal name we gave it.

[Statutory Authority: RCW 18.71.017. WSR 04-04-067, § 246-919-320, filed 2/2/04, effective 3/4/04. Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, § 246-919-320, filed 1/17/96, effective 2/17/96.]

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WAC 246-919-330 Postgraduate medical training defined. (1) For the purposes of this chapter, pPostgraduate medical training means clinical

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training approved by the commission in general medicine or surgery, or a specialty or subspecialty in the field of medicine or surgery as recognized by the American Board of Medical Specialties and listed in

the 2004 Official ABMS Annual Report and Reference Handbook, published

March 18, 2004. Listed in the 2017-2018 ABMS Board Certification Report

and new specialties or subspecialties approved by the commission.

- (2) The commission approves only the following postgraduate clinical training courses:
- (a) Programs accredited by the Accreditation Council for Graduate Medical Education (ACGME) which are listed in the 1984-85 directory of residency programs, or programs approved by the Accreditation Council at the time of residency.
- (b) Programs accredited by the Royal College of Physicians and Surgeons of Canada (RCPSC) or the College of Family Physicians of Canada (CFPC), or programs accredited by the RCPSC or CFPC at the time of residency.
- (3) Postgraduate medical training includes, but is not limited to, internships, residencies and medical or surgical fellowships.
- (4) A physician must complete two consecutive years of post—graduate medical training in no more than two programs. The physician must acquire this training after completion of a formal course of undergraduate medical instruction outlined in RCW 18.71.055. The commission will accept only satisfactory clinical performance evaluations.

Commented [UD(6]: Approved 11/14/18

Commented [UD(7]: Add the most current version. Then say: "and new programs approved by the commission." (would be approved at a Commission meeting.)

Commented [UD(8R7]: This is no longer a book, it is a website. As we cannot add websites to rules, I suggest we just state "Programs accredited by the Accreditation Council for Graduate Medical Education (ACGME)."

Commented [UD(9R7]: Approved 1.16.19

Commented [UD(10]: Approved.

[Statutory Authority: RCW 18.71.017 and 18.71.050. WSR 05-07-024, § 246-919-330, filed 3/7/05, effective 4/7/05. Statutory Authority: RCW 18.71.017, 18.71.050 and chapter 18.71 RCW. WSR 01-18-087, § 246-919-330, filed 9/5/01, effective 10/6/01. Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, § 246-919-330, filed 1/17/96, effective 2/17/96.]

WAC 246-919-340 Additional requirements for international medical school graduates. All graduates of medical schools outside the United States, Canada, or Puerto Rico must satisfy one of the following requriments have either:

- (1) Been licensed in another state prior to 1958 Held a full and unrestricted license to practice medicine in another state prior to 1958;
- (2) Obtained a certificate with an indefinite status granted by the Educational Commission for Foreign Medical Graduates (ECFMG); or
- (3) Successfully completed one year of supervised academic clinical training in the United States, commonly referred to as a Fifth Pathway program.

[Statutory Authority: RCW 18.71.017, 18.71.050 and chapter 18.71 RCW. WSR 01-18-086, \$246-919-340, filed 9/5/01, effective 10/6/01. Statutory

Commented [UD(11]: Approved 11/14/18

Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, § 246-919-340, filed 1/17/96, effective 2/17/96.]

WAC 246-919-355 Examinations accepted by the commission scores. mmission:

- (1) The commission adopts—accepts the United States Medical Licensing Examination (USMLE) as the examination for licensureaccepted by commission.
- (2) The minimal passing scores for each component of any approved examination combination shall be a score of seventy-five as defined by the examining authority.
- (3) Applicants must have passed all components of the USMLE within seven years after passing the first examination.
- (a) The commission recognizes that an applicant with a combined degree may require an exception to the seven-year requirement. The commission will review exception requests on a case by case basis.
- (3) Applicants who do not pass Step 3 of the USMLE examination nation, either Step 1 or Step 2, or acceptable combination, shall demon-

remedial or refresher medical course approved by the commission prior
to being permitted to sit for the examination again. Applicants who do
not pass after the fourth sitting may not sit for another examination
without completing an additional year of postgraduate training or satisfying any other conditions specified by the commission.

(4) To be eligible for USMLE Step 3, the applicant must:
(a) Have obtained the M.D. degree;
(b) Have successfully completed the Federation Licensure Examination (FLEX) Component 1 or both National Boards Examination (NBE) Parts
I and II or USMLE Steps 1 and 2 or NBE Part I and USMLE Step 2 or Step
1 and NBE Part II; and
(c) Be certified by the ECFMG if a graduate of an international
medical school, or have successfully completed a fifth pathway program;
and postgraduate training year in a program of graduate medical education
accredited by the Accreditation Council for Graduate Medical Education.
[Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, \$
246-919-355, filed 1/17/96, effective 2/17/96.]

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water. (1) The commission may accept certain examinations as a basis for licensure. These examinations include USMLE, FLEX, NBE, or those

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WAC (2/8/2018 11:51 AM)

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given by the other states, or territories of the United States. Those who have taken the Licentiate of the Medical Council of Canada (L.M.C.C.) and holds a valid LMCC certification obtained after 1969, may be granted a license without examination.

(2) Examination combination acceptable. Any applicant who has successfully completed Part I (NBE) or Step 1 (USMLE) plus Part II or Step 2 plus Part III or Step 3; or FLEX Component 1 plus Step 3; or Part I or Step 1, plus Part II or Step 2, plus FLEX Component 2 shall be deemed to have successfully completed a medical licensure examination as required by RCW 18.71.070. (For clarification, see Table 1.)

Accepted Examinations taken in Sequence	Other Acceptable Combinations
NBME Part I plus NBME Part II plus NBME Part III	NBME Part I or USMLE Step 1 plus NBME Part II or USMLE Step 2 plus NBME Part III or USMLE Step 3
FLEX Component 1 plus FLEX Component 2	FLEX Component 1 plus USMLE Step 3 or NBME Part I or USMLE Step 1 plus NBME Part II or USMLE Step 2 plus FLEX Component 2
USMLE Step 1 plus USMLE Step 2 plus USMLE Step 3	

[Statutory Authority: RCW 18.71.017, 18.130.050, 18.71.090, and [18.71.]095. WSR 06-18-042, § 246-919-360, filed 8/30/06, effective

9/30/06. Statutory Authority: RCW 18.71.017. WSR 04-04-067, § 246-919-360, filed 2/2/04, effective 3/4/04. Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, § 246-919-360, filed 1/17/96, effective 2/17/96.]

WAC 246-919-365 FLEX examination standards. Reciprocity applicants licensed in another state by passing the FLEX examination be eligible for a waiver of examination if the applicant received a FLEX weighted average score of at least 75. The score may be obtained in a single setting of the three day examination or by averaging the individual day scores from different examinations. The individual day scores will be averaged according to the following formula:

Day 1 equals 1/6. Day 3 equals 3/6.

(i.e., an average of 74.9 equals 74). Single subject averaging is not permitted. The commission will accept the federation licensing examination (FLEX) weighted average of 75 reported from the Federation of State Medical Boards. All FLEX scores must be submitted directly from the Federation of State Medical Boards. FLEX scores reported by other states will not be accepted.

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[Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, § 246-919-365, filed 1/17/96, effective 2/17/96.]

wac 246-919-370 Special purpose examination. (1) The commission may require an applicant or licensee to pass the Special Purpose Examination (SPEX) or any other examination deemed appropriate. An applicant or licensee may be required to take an examination when the commission has concerns with the applicant's or licensee's ability to practice competently for reasons which may include, but are not limited to, the following:

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- (a) Resolved or pending malpractice suits;
- (b) Pending action by another state licensing authority;
- (c) Actions pertaining to privileges at any institution; or
- (d) Not having practiced for the immediate two years prior to the

application an interval of time.

(2) The minimum passing score on the SPEX examination shall be seventy-five. The passing score for any other examination under this rule shall be determined by the commission.

Commented [UD(16]: Approved 11/14/18

[Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, § 246-919-370, filed 1/17/96, effective 2/17/96.]

WAC 246-919-380 AIDS/HIV prevention and information education requirements. Applicants must complete four clock hours of AIDS/HIV education as required in chapter 246-12 WAC, Part 8.

[Statutory Authority: RCW 43.70.280. WSR 98-05-060, § 246-919-380, filed 2/13/98, effective 3/16/98. Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, § 246-919-380, filed 1/17/96, effective 2/17/96.]

WAC 246-919-395 Substantially equivalent licensing standards—Temporary practice permit. (1) An applicant who holds an unrestricted, active license in another state with licensing standards substantially equivalent to those in Washington may apply for a temporary practice permit authorizing the applicant to practice as a physician in Washington.

- (2) The commission will issue the physician a temporary practice permit if the following requirements are met:
- (a) The applicant submits a completed application for a physician and surgeon license on a form provided by the commission on which the

Commented [UD(17]: Approved 11/14/18

applicant indicates that he or she wishes to receive a temporary practice permit;

- (b) The applicant submits payment of the application fee and temporary practice permit fee pursuant tounder WAC 246-919-990;
- (c) The commission receives the American Medical Association's physicians' data profile verifying states in which the applicant is or was licensed;
- (d) The commission receives the practitioner profile from the Federation of State Medical Boards;
- (e) The applicant requests and the commission receives written verification attesting that the applicant has a license in good standing and is not subject to charges or disciplinary action for unprofessional conduct or impairment from all states which the applicant is or was licensed;
- (f) The applicant is not subject to denial of a license or issuance of a conditional license under chapter 18.130 RCW; and
- (g) The applicant is licensed in a state that has licensing standards substantially equivalent to Washington.
- (3) The temporary practice permit allows the applicant to work in the state of Washington as a physician without restriction until the

permit expires. The temporary practice permit is a license to practice medicine.

- (4) The temporary permit shall expire upon the issuance of a license by the commission; initiation of an investigation by the commission of the applicant; or ninety days after the temporary practice permit is issued, whichever occurs first. The temporary permit will not be renewed, reissued, or extended.
- (5) An applicant who receives a temporary practice permit and who does not complete the application process may not receive additional temporary practice permits even upon submission of a new application in the future.

[Statutory Authority: RCW 18.71.017 and 18.130.075. WSR 17-18-098, § 246-919-395, filed 9/6/17, effective 10/7/17. Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, § 246-919-395, filed 1/17/96, effective 2/17/96.]

WAC 246-919-396 Background check—Temporary practice permit. The medical quality assurance commission (MQAC)—conducts background checks on applicants to assure safe patient care. Completion of a national criminal background check may require additional time. The MQAC—commission may issue a temporary practice permit when the applicant has met

all other licensure requirements, except the national criminal background check requirement. The applicant must not be subject to denial of a license or issuance of a conditional license under this chapter.

(1) If there are no violations identified in the Washington criminal background check and the applicant meets all other licensure conditions, including receipt by the department of health of a completed Federal Bureau of Investigation (FBI) fingerprint card, the MQAC—commission may issue a temporary practice permit allowing time to complete the national criminal background check requirements.

The MQACcommission will issue a temporary practice permit that is valid for six months. A one timeone-time extension of six months will be granted if the national background check report has not been received by the MQACcommission.

- (2) The temporary practice permit allows the applicant to work in the state of Washington as a physician during the time period specified on the permit. The temporary practice permit is a license to practice medicine.
- (3) The MQACcommission issues a license after it receives the national background check report if the report is negative and the applicant otherwise meets the requirements for a license.

(4) The temporary practice permit is no longer valid after the license is issued or action is taken on the application because of the background check.

[Statutory Authority: RCW 18.130.064 and 18.130.075. WSR 10-05-029, § 246-919-396, filed 2/9/10, effective 2/11/10.]

NEW SECTION

WAC 246-919-397 How to obtain a temporary practice permit—Military spouse. A military spouse or state registered domestic partner of a military person may receive a temporary practice permit while completing any specific additional requirements that are not related to training or practice standards for physicians.

- (1) A temporary practice permit may be issued to an applicant who is a military spouse or state registered domestic partner of a military person and:
- (a) Is moving to Washington as a result of the military person's transfer to Washington;
- (b) Left employment in another state to accompany the military person to Washington;

- (c) Holds an unrestricted, active license in another state that has substantially equivalent licensing standards for a physician to those in Washington; and
- (d) Is not subject to any pending investigation, charges, or disciplinary action by the regulatory body of the other state or states.
- (2) A temporary practice permit grants the individual the full scope of practice for the physician.
- (3) A temporary practice permit expires when any one of the following occurs:
 - (a) The license is granted;
- (b) A notice of decision on the application is mailed to the applicant, unless the notice of decision on the application specifically extends the duration of the temporary practice permit; or
- (c) One hundred eighty days after the temporary practice permit is issued.
 - (4) To receive a temporary practice permit, the applicant must:
 - (a) Submit to the commission the necessary application, fee(s),
- fingerprint card if required, and documentation for the license;
- (b) Attest on the application that the applicant left employment in another state to accompany the military person;

- (c) Meet all requirements and qualifications for the license that are specific to the training, education, and practice standards for physicians;
- (d) Provide verification of having an active unrestricted license in the same profession from another state that has substantially equivalent licensing standards for physicians in Washington;
 - (e) Submit a copy of the military person's orders and a copy of:
- (i) The military-issued identification card showing the military person's information and the applicant's relationship to the military

person;

- (ii) A marriage license; or
- (iii) A state registered domestic partnership; and
- (f) Submit a written request for a temporary practice permit.
- (5) For the purposes of this section:
- (a) "Military spouse" means the husband, wife, or registered do-

mestic partner of a military person.

- (b) "Military person" means a person serving in the United States
- armed forces, the United States public health service commissioned

corps, or the merchant marine of the United States.

Commented [UD(18]: 7/11/18 – approved to add.

RENEWAL AND CME REQUIREMENTS

WAC 246-919-421 Two year renewal cycle. A licensed physician shall renew his or her license every two years in compliance with WAC 246-12-030. A licensed physician must also submit information about his or her current professional practice as required by RCW 18.71.080 (1)(b). [Statutory Authority: RCW 18.71.002, 18.71.017, and 18.71.080. WSR 16-16-028, § 246-919-421, filed 7/22/16, effective 8/22/16. Statutory Authority: RCW 18.71.017, 18.130.250, 18.71.440. WSR 11-05-025, § 246-919-421, filed 2/7/11, effective 3/10/11. Statutory Authority: RCW 18.71.017, 18.130.050(1), 18.130.040(4), 18.130.050(12) and 18.130.340. WSR 99-23-090, § 246-919-421, filed 11/16/99, effective 1/1/00.]

WAC 246-919-422 Transition from post-graduate limited license to full license. In order to obtain full license status, individuals—a physician with a post-graduate limited Washington state license will pay the fee difference between the limited license application and the full license application. This license will expire on their second birth date after issuance and every two years thereafter. [Statutory Authority: RCW 18.71.002, 18.71.017, and 18.71.080. WSR 16-16-028, § 246-919-422, filed 7/22/16, effective 8/22/16.]

Commented [UD(19]: Approved 11/14/18

WAC 246-919-430 Requirements for maintenance of licensure. A licensed physician must complete one of the following to satisfy maintenance of licensure requirements during renewal:

- (1) Complete two hundred hours of continuing education every four years as required in chapter 246-12 WAC and as described in WAC 246-919-460. Participation in a residency program accredited by the Accreditation Council for Graduate Medical Education or in a fellowship program, accredited or not, may be credited fifty hours of Category I continuing medical education per year of training towards the two hundred hour requirement;
- (2) Obtain a current Physician's Recognition Award from the American Medical Association in at least two of the four years preceding the renewal due date;
- (3) Become certified by a member board of the American Board of Medical Specialties in the four years preceding the renewal due date;
- (4) Meet the requirements for participation in maintenance of certification of a member board of the American Board of Medical Specialties at the time of renewal.

[Statutory Authority: RCW 18.71.002, 18.71.017, and 18.71.080. WSR 16-16-028, § 246-919-430, filed 7/22/16, effective 8/22/16. Statutory Authority: RCW 18.71.017, 18.130.050(1), 18.130.040(4), 18.130.050(12) and

18.130.340. WSR 99-23-090, § 246-919-430, filed 11/16/99, effective 1/1/00. Statutory Authority: RCW 43.70.280. WSR 98-05-060, § 246-919-430, filed 2/13/98, effective 3/16/98. Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, § 246-919-430, filed 1/17/96, effective 2/17/96.]

WAC 246-919-435 Training in suicide assessment, treatment, and management. (1) A licensed physician, other than a resident holding a limited license issued under RCW 18.71.095(3), must complete a one-time training in suicide assessment, treatment, and management. The training must be at least six hours in length and may be completed in one or more sessions.

(2) The training must be completed by the end of the first full continuing education reporting period after January 1, 2016, or during the first full continuing education period after initial licensure, whichever occurs later, or during the first full continuing education reporting period after the exemption in subsection (6) of this section no longer applies. The commission accepts training completed between June 12, 2014, and January 1, 2016, that meets the requirements of RCW 43.70.442 as meeting the one-time training requirement.

__(3) Until July 1, 2017, the commission must approve the training.

The commission will approve an empirically supported training in suicide assessment, suicide treatment, and suicide management that meets the requirements of RCW 43.70.442.

- (4) Beginning July 1, 2017, Tthe training must be on the model list developed by the department of health under RCW 43.70.442. The establishment of the model list does not affect the validity of training completed prior to July 1, 2017.
- (5) The hours spent completing training in suicide assessment, treatment, and management count toward meeting applicable continuing education requirements in the same category specified in WAC 246-919-460.
- (6) The commission exempts any licensed physician from the training requirements of this section if the physician has only brief or limited patient contact, or no patient contact.

[Statutory Authority: RCW 18.71.017 and 43.70.442. WSR 17-07-043, § 246-919-435, filed 3/8/17, effective 4/8/17.]

WAC 246-919-460 Categories of creditable continuing medical education activities. (1) Category I: Continuing medical education activities with accredited sponsorship. The licensed physician may earn all two

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hundred credit hours in Category I. The commission will accept attendance at a continuing education program that is recognized as Category I credit and is offered by an organization or institution that meets the standards adopted by the Accreditation Council for Continuing Medical Education or its designated interstate accrediting agency, the Washington State Medical Association.

- (2) Category II: Continuing medical education activities with nonaccredited sponsorship. A licensed physician may earn a maximum of eighty credit hours by attendance at continuing medical education programs that are not approved but which are in accordance with the provisions of Category I.
- (3) Category III: Teaching of physicians or other allied health professionals. A licensed physician may earn a maximum of eighty credit hours for serving as an instructor of medical students, house staff, other physicians or allied health professionals from a hospital or institution with a formal training program if the hospital or institution has approved the instruction.
- (4) Category IV: Books, papers, publications, exhibits. A licensed physician may earn:

- (a) A maximum of eighty credit hours under Category IV, with specific subcategories listed below. Credit may be earned only during the forty-eight-month period following presentations or publications.
- (b) Ten credit hours for a paper, exhibit, publication, or for each chapter of a book that is authored by the licensed physician and published. A paper must be published in a recognized medical journal. A licensed physician who presents a paper at a meeting or an exhibit must present to physicians or allied health professionals. Credit may be claimed only once for the scientific materials presented. Credit should be claimed as of the date materials were presented or published.

Medical editing will not be accepted in this or any other category for credit.

- (5) Category V: Self-directed activities. A licensed physician may earn:
 - (a) A maximum of eighty credit hours under Category V.
- (b) Self-assessment: Credit hours for completion of a multimedia medical education program.
- (c) Self-instruction: Credit hours for the independent reading of scientific journals and books.

- (d) Specialty board examination preparation: Credit hours for preparation for specialty board certification or recertification examinations.
- (e) Quality care or utilization review: Credit hours for participation on a staff committee for quality of care or utilization review in a hospital or institution or government agency.

[Statutory Authority: RCW 18.71.002, 18.71.017, and 18.71.080. WSR 16-16-028, § 246-919-460, filed 7/22/16, effective 8/22/16. Statutory Authority: RCW 18.71.017, 18.130.050(1), 18.130.040(4), 18.130.050(12) and 18.130.340. WSR 99-23-090, § 246-919-460, filed 11/16/99, effective 1/1/00. Statutory Authority: RCW 43.70.280. WSR 98-05-060, § 246-919-460, filed 2/13/98, effective 3/16/98. Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, § 246-919-460, filed 1/17/96, effective 2/17/96.]

WAC 246-919-470 Approval not required. (1) Except as required by law, the commission will not give prior approval for any continuing medical education. The commission will accept any continuing medical education that reasonably falls within these rules and relies upon each individual physician's integrity to comply with this requirement.

(2) The continuing medical education category will depend solely upon the accredited status of the organization or institution. The number of creditable hours may be determined by counting the contact hours of instruction and rounding to the nearest quarter hour. The commission relies upon the integrity of program sponsors to present continuing medical education that constitutes a meritorious learning experience. [Statutory Authority: RCW 18.71.002, 18.71.017, and 18.71.080. WSR 16-16-028, § 246-919-470, filed 7/22/16, effective 8/22/16. Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, § 246-919-470, filed 1/17/96, effective 2/17/96.]

WAC 246-919-475 Expired license. (1) If the license has expired for three years or less the practitioner physician must meet the requirements of chapter 246-12 WAC, Part 2.

- (2) If the license has expired for over three years, the practitioner physician must:
- (a) Reapply for licensing under current requirements as stipulated in RCW 18.71.050 (1)(b) and WAC 246-919-330; and
- (b) Meet the requirements of chapter 246-12 WAC, Part 2. [Statutory Authority: RCW 18.71.017. WSR 01-03-115, § 246-919-475, filed 1/22/01, effective 2/22/01.]

WAC 246-919-480 Retired active license. (1) To obtain a retired active license a physician must comply with chapter 246-12 WAC, Part 5, excluding WAC 246-12-120 (2)(c) and (d).

- (2) A physician with a retired active license may not receive compensation for health care services;
- (3) A physician with a retired active license may practice only in emergent or intermittent circumstances; and
- (4) A Pphysicians with a retired active license must renew every two years and must report one hundred hours of continuing medical education at every renewal. The commission will accept a maximum of A physician can receive a maximum of 40 hours of continuing medical education in each of categories II through V during each renewal period. There is no limit to the number of hours that may be earnedaccepted in Category I.

[Statutory Authority: RCW 18.71.017, 18.130.250, 18.71.440. WSR 11-05-025, \$ 246-919-480, filed 2/7/11, effective 3/10/11. Statutory Authority: RCW 43.70.280. WSR 98-05-060, \$ 246-919-480, filed 2/13/98, effective 3/16/98. Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, \$ 246-919-480, filed 1/17/96, effective 2/17/96.]

ADJUDICATIVE PROCEDURES

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WAC 246-919-520 Revocation of a physician's license. This section sets forth the procedure by which a respondent physician may request a review by the medical quality assurance commission of its decision to revoke the respondent physician's license under RCW 18.71.019:

- (1) If the commission issues a final order revoking a respondent physician's license following an adjudicative proceeding, the respondent physician may request a review of the decision by a review panel of the commission.
- (2) The respondent physician shall file a written request with the commission within twenty days of effective date of the final order. The respondent physician may not request an extension of the twenty-day period to file a request for review.
- (3) The respondentphysician's request for review of the final order does not change the effective date of the final order.
- (4) A review panel shall review the final order. The review panel is composed of the members of the commission who did not:
- (a) Review the initial investigation and make the decision to issue a statement of charges against the respondentphysician in this matter;
 or
- (b) Hear the evidence at the adjudicative proceeding and issue the final order revoking the $\frac{1}{1}$

- (5) Within seven days of receipt of the request for review of the final order, a scheduling order is issued setting a date for the review hearing, and a date for the filing of written argument by the parties.

 The review hearing must take place within sixty days of the respondent physician's request for review of the final order.
- (6) The review panel shall convene in person for the review hearing on the date set in the scheduling order. If a commission member is unavailable to meet on the scheduled date, a pro tempore member shall take that person's place on the review panel. At the review hearing, the review panel:
 - (a) Shall review the final order;
 - (b) Shall review written argument presented by the parties; and
 - (c) May hear oral argument by the parties.
- (7) If the review panel determines that revocation of the respondentphysician's license is not the appropriate sanction, it shall issue an amended order setting the appropriate sanction(s) necessary to protect the public.
- (8) If the review panel determines that revocation of the respondent physician's license is appropriate, it shall issue an order confirming that decision.

[Statutory Authority: RCW 18.71.019. WSR 97-21-053, § 246-919-520, filed 10/13/97, effective 11/13/97.]

OFFICE-BASED SURGERY RULES

WAC 246-919-601 Safe and effective analgesia and anesthesia administration in office-based surgical settings. (1) Purpose. The purpose of this rule is to promote and establish consistent standards, continuing competency, and to promote patient safety. The medical quality assurance commission establishes the following rule for physicians licensed under this chapter who perform surgical procedures and use anesthesia, analgesia or sedation in office-based settings.

- (2) Definitions. The following terms used in this subsection apply throughout this rule unless the context clearly indicates otherwise:
 - (a) "Commission" means the medical quality assurance commission.

(b) "Deep sedation" or "analgesia" means a drug-induced depression of consciousness during which patients cannot be easily aroused but respond purposefully following repeated or painful stimulation. The ability to independently maintain ventilatory function may be impaired. Patients may require assistance in maintaining a patent airway, and spontaneous ventilation may be inadequate. Cardiovascular function is usually maintained.

- (eb) "General anesthesia" means a state of unconsciousness intentionally produced by anesthetic agents, with absence of pain sensation over the entire body, in which the patient is without protective reflexes and is unable to maintain an airway. Sedation that unintentionally progresses to the point at which the patient is without protective reflexes and is unable to maintain an airway is not considered general anesthesia.
- (4c) "Local infiltration" means the process of infusing a local anesthetic agent into the skin and other tissues to allow painless wound irrigation, exploration and repair, and other procedures, including procedures such as retrobulbar or periorbital ocular blocks only when performed by a board eligible or board certified ophthalmologist. It does not include procedures in which local anesthesia is injected into areas of the body other than skin or muscle where significant cardiovascular or respiratory complications may result.
- (ed) "Major conduction anesthesia" means the administration of a drug or combination of drugs to interrupt nerve impulses without loss of consciousness, such as epidural, caudal, or spinal anesthesia, lumbar or brachial plexus blocks, and intravenous regional anesthesia. Major conduction anesthesia does not include isolated blockade of small peripheral nerves, such as digital nerves.

- (fe) "Minimal sedation" means a drug-induced state during which patients respond normally to verbal commands. Although cognitive function and coordination may be impaired, ventilatory and cardiovascular functions are unaffected. Minimal sedation is limited to oral or intramuscular medications, or both.
- (gf) "Moderate sedation" or "analgesia" means a drug-induced depression of consciousness during which patients respond purposefully to verbal commands, either alone or accompanied by tactile stimulation. No interventions are required to maintain a patent airway, and spontaneous ventilation is adequate. Cardiovascular function is usually maintained.
- (hg) "Office-based surgery" means any surgery or invasive medical procedure requiring analgesia or sedation, including, but not limited to, local infiltration for tumescent liposuction, performed in a location other than a hospital or hospital-associated surgical center licensed under chapter 70.41 RCW, or an ambulatory surgical facility licensed under chapter 70.230 RCW.
- _(i) "Physician" means an individual licensed under chapter 18.71

 RCW.
 - (3) Exemptions. This rule does not apply to physicians when:
- (a) Performing surgery and medical procedures that require only minimal sedation (anxiolysis), or infiltration of local anesthetic

around peripheral nerves. Infiltration around peripheral nerves does not include infiltration of local anesthetic agents in an amount that exceeds the manufacturer's published recommendations.

- (b) Performing surgery in a hospital or hospital-associated surgical center licensed under chapter 70.41 RCW, or an ambulatory surgical facility licensed under chapter 70.230 RCW.
- (c) Performing surgery utilizing or administering general anesthesia. Facilities in which physicians administer general anesthesia orperform procedures in which general anesthesia is a planned event are regulated by rules related to hospital or hospital-associated surgical center licensed under chapter 70.41 RCW, or an ambulatory surgical facility licensed under chapter 70.230 RCW, or a dental office under WAC 246-919-602.
- (d) Administering deep sedation or general anesthesia to a patient in a dental office under WAC 246-919-602
 - (e) Performing oral and maxillofacial surgery, and the physician:
- (i) Is licensed both as a physician under chapter $18.71\ \text{RCW}$ and as a dentist under chapter $18.32\ \text{RCW};$
 - (ii) Complies with dental quality assurance commission regulations;
 - (iii) Holds a valid:
 - (A) Moderate sedation permit; or

- (B) Moderate sedation with parenteral agents permit; or
- (C) General anesthesia and deep sedation permit; and
- (iv) Practices within the scope of his or her specialty.
- (4) Application of rule.

This rule applies to physicians practicing independently or in a group setting who perform office-based surgery employing one or more of the following levels of sedation or anesthesia:

- (a) Moderate sedation or analgesia; or
- (b) Deep sedation or analgesia; or
- (c) Major conduction anesthesia.
- (5) Accreditation or certification.
- (a) A physician who performs a procedure under this rule must ensure that the procedure is performed in a facility that is appropriately equipped and maintained to ensure patient safety through accreditation or certification and in good standing from an accrediting entity approved by the commission.
- (b) The commission may approve an accrediting entity that demonstrates to the satisfaction of the commission that it has:
- (i) Standards pertaining to patient care, recordkeeping, equipment, personnel, facilities and other related matters that are in accordance

with acceptable and prevailing standards of care as determined by the commission;

- (ii) Processes that assure a fair and timely review and decision on any applications for accreditation or renewals thereof;
- (iii) Processes that assure a fair and timely review and resolution of any complaints received concerning accredited or certified facilities; and
- (iv) Resources sufficient to allow the accrediting entity to fulfill its duties in a timely manner.
- (c) A physician may perform procedures under this rule in a facility that is not accredited or certified, provided that the facility has submitted an application for accreditation by a commission-approved accrediting entity, and that the facility is appropriately equipped and maintained to ensure patient safety such that the facility meets the accreditation standards. If the facility is not accredited or certified within one year of the physician's performance of the first procedure under this rule, the physician must cease performing procedures under this rule until the facility is accredited or certified.
- (d) If a facility loses its accreditation or certification and is no longer accredited or certified by at least one commission-approved

entity, the physician shall immediately cease performing procedures under this rule in that facility.

- (6) Competency. When an anesthesiologist or certified registered nurse anesthetist is not present, the physician performing office-based surgery and using a form of sedation defined in subsection (4) of this section must be competent and qualified both to perform the operative procedure and to oversee the administration of intravenous sedation and analgesia.
- (7) Qualifications for administration of sedation and analgesia may include:
- (a) Completion of a continuing medical education course in conscious sedation;
 - (b) Relevant training in a residency training program; or
- (c) Having privileges for conscious sedation granted by a hospital $\mbox{medical staff.}$
- (8) At least one licensed health care practitioner currently certified in advanced resuscitative techniques appropriate for the patient age group (e.g., ACLS, PALS or APLS) must be present or immediately available with age-size-appropriate resuscitative equipment throughout the procedure and until the patient has met the criteria for discharge from the facility.

- (9) Sedation assessment and management.
- (a) Sedation is a continuum. Depending on the patient's response to drugs, the drugs administered, and the dose and timing of drug administration, it is possible that a deeper level of sedation will be produced than initially intended.
- (b) If an anesthesiologist or certified registered nurse anesthetist is not present, a physician intending to produce a given level of sedation should be able to "rescue" a patient who enters a deeper level of sedation than intended.
- (c) If a patient enters into a deeper level of sedation than planned, the physician must return the patient to the lighter level of sedation as quickly as possible, while closely monitoring the patient to ensure the airway is patent, the patient is breathing, and that oxygenation, heart rate and blood pressure are within acceptable values. A physician who returns a patient to a lighter level of sedation in accordance with this subsection (c) does not violate subsection (10) of this section.
 - (10) Separation of surgical and monitoring functions.
- (a) The physician performing the surgical procedure must not administer the intravenous sedation, or monitor the patient.

- (b) The licensed health care practitioner, designated by the physician to administer intravenous medications and monitor the patient who is under moderate sedation, may assist the operating physician with minor, interruptible tasks of short duration once the patient's level of sedation and vital signs have been stabilized, provided that adequate monitoring of the patient's condition is maintained. The licensed health care practitioner who administers intravenous medications and monitors a patient under deep sedation or analgesia must not perform or assist in the surgical procedure.
- (11) Emergency care and transfer protocols. A physician performing office-based surgery must ensure that in the event of a complication or emergency:
- (a) All office personnel are familiar with a written and documented plan to timely and safely transfer patients to an appropriate hospital.
- (b) The plan must include arrangements for emergency medical services and appropriate escort of the patient to the hospital.
- (12) Medical record. The physician performing office-based surgery must maintain a legible, complete, comprehensive, and accurate medical record for each patient.
 - (a) The medical record must include:
 - (i) Identity of the patient;

- (ii) History and physical, diagnosis and plan;
- (iii) Appropriate lab, X-ray or other diagnostic reports;
- (iv) Appropriate preanesthesia evaluation;
- (v) Narrative description of procedure;
- (vi) Pathology reports, if relevant;
- (vii) Documentation of which, if any, tissues and other specimens have been submitted for histopathologic diagnosis;
 - (viii) Provision for continuity of postoperative care; and
 - (ix) Documentation of the outcome and the follow-up plan.
- (b) When moderate or deep sedation, or major conduction anesthesia is used, the patient medical record must include a separate anesthesia record that documents:
 - (i) The type of sedation or anesthesia used;
 - (ii) Drugs (name and dose) and time of administration;
- (iii) Documentation at regular intervals of information obtained from the intraoperative and postoperative monitoring;
 - (iv) Fluids administered during the procedure;
 - (v) Patient weight;
 - (vi) Level of consciousness;
 - (vii) Estimated blood loss;
 - (viii) Duration of procedure; and

(ix) Any complication or unusual events related to the procedure or sedation/anesthesia.

[Statutory Authority: RCW 18.71.017. WSR 17-18-032, § 246-919-601, filed 8/28/17, effective 9/28/17. Statutory Authority: RCW 18.71.017 and 18.130.050(4). WSR 10-16-109, § 246-919-601, filed 8/2/10, effective 9/2/10.1

NEW SECTION

WAC 246-919-602 Administration of deep sedation and general anesthesia by physicians in dental offices.

- (1) Purpose. The purpose of this rule is to govern the administration of deep sedation and general anesthesia by physicians in dental offices. The commission establishes these standards to promote effective perioperative communication and appropriately timed interventions, and mitigate adverse events and outcomes.
- (2) Definitions. The definitions in this subsection apply throughout this section unless the context clearly requires otherwise.
- (a) "Administering physician" means an individual licensed under chapter 18.71 RCW who administers deep sedation or general anesthesia to a patient in a dental office.
 - (b) "Deep sedation" has the same meaning as in WAC 246-919-601.

- (c) "Dental office" means any facility where dentistry is practiced, as defined in chapter 18.32 RCW, except a hospital licensed under chapter 70.41 RCW or ambulatory surgical facility licensed under chapter 70.230 RCW.
- (d) "General anesthesia" has the same meaning as in WAC 246-919-601.
- (e) "Perioperative" includes the three phases of surgery: preoperative, intraoperative, and postoperative.
- (3) An administering physician is responsible for the perioperative anesthetic management and monitoring of a patient and must ensure patient care, recordkeeping, equipment, personnel, facilities, and other related matters are in accordance with acceptable and prevailing standards of care, including, but not limited to, the following:
- (a) Preoperative requirements. An administering physician must ensure the patient has undergone a preoperative health evaluation and document review of the evaluation. The physician must also conduct and document a risk assessment to determine whether a patient is an appropriate candidate for deep sedation or general anesthesia and discussion of the risks of deep sedation or general anesthesia with the patient. For a pediatric patient, this assessment must include:

- (i) Whether the patient has specific risk factors that may warrant additional consultation before administration of deep sedation or general anesthesia, and how each patient meets criteria for deep sedation or general anesthesia in an outpatient environment;
- (ii) A discussion with a parent or guardian of a pediatric patient of the particular risks of deep sedation or general anesthesia for a patient who (A) is younger than six years old; (B) has special needs; (C) has airway abnormalities; or (D) has a chronic condition. This discussion must include reasoning why the pediatric patient can safely receive deep sedation or general anesthesia in an outpatient environment and any alternatives.
- (b) Medical record. The recordkeeping requirements under WAC 246-919-601 and 246-817-770 apply to an administering physician, including the elements of a separate anesthesia record. The anesthesia record must be complete, comprehensive, and accurate for each patient, including documentation at regular intervals of information from intraoperative and postoperative monitoring. For a pediatric patient, the administering physician must ensure vital signs are postoperatively recorded at least at five-minute intervals until the patient begins to awaken, then recording intervals may be increased to ten-to-fifteen minutes.

- (c) Equipment. An administering physician must ensure the requirements for equipment and emergency medications under WAC 246-817-724 and 246-817-770 are met, regardless of any delineated responsibility for furnishing of the equipment or medications in a contract between the physician and dental office. The physician must also ensure equipment is available in the recovery area to meet the requirements in this section for monitoring during the recovery period. The physician must ensure all equipment and medications are checked and maintained on a scheduled basis.
- (d) Recovery and discharge requirements. An administering physician must ensure that:
- (i) A physician licensed under chapter 18.71 RCW capable of managing complications, providing cardiopulmonary resuscitation, and currently certified in advanced cardiac life support measures appropriate for the patient age group is immediately available for a patient recovering from anesthesia. For a pediatric patient, the physician must also be trained and experienced in pediatric perioperative care;
- (ii) At least one licensed health care practitioner experienced in postanesthetic recovery care and currently certified in advanced cardiac life support measures appropriate for the patient age group visually monitors the patient, at all times, until the patient has met

the criteria for discharge from the facility. A practitioner may not monitor more than two patients simultaneously, and any such simultaneous monitoring must take place in a single recovery room. The practitioner must provide: (A) continuous respiratory and cardiovascular monitoring during the recovery period; and (B) monitoring, at regular intervals, of the patient for color of mucosa, skin, or blood, oxygen saturation, blood pressure, and level of consciousness;

- (iii) Emergency equipment, supplies, and services are immediately available in all areas where anesthesia is used and for a patient recovering from anesthesia. Resuscitative equipment must be age and size-appropriate and support personnel must have knowledge of the emergency care inventory. All equipment and drugs must be checked and maintained on a scheduled basis;
- (iv) Before discharge, the patient is awake, alert, and behaving appropriately for age and developmental status, normal patient vital signs, and if applicable, a capable parent or guardian present to assume care of the patient.
- (e) Emergency care and transfer protocol. An administering physician must monitor for, and be prepared to treat, complications involving compromise of the airway and depressed respiration, particularly with a pediatric patient. The physician must ensure that in the event of a

complication or emergency, his or her assistive personnel and all dental office clinical staff are well-versed in emergency recognition, rescue, and emergency protocols, and familiar with a written and documented plan to timely and safely transfer a patient to an appropriate hospital.

(4) (a) An administering physician must submit to the commission a report of any patient death or serious perioperative complications, which is or may be the result of anesthesia administered by the physician. When a patient comes into an office with an existing condition, and hospital admission is the result of that condition and not the anesthesia, it is not reportable.

- (b) The physician must notify the commission or the department of health, by telephone, email, or fax within seventy-two hours of discovery and must submit a complete written report to the commission within thirty days of the incident. The written report must include the following:
- (i) Name, age, and address of the patient.
- (ii) Name of the dentist and other personnel present during the incident.
- (iii) Address of the facility or office where the incident took place.
- (iv) Description of the type of anesthetic being utilized at the time of the incident.
- (v) Dosages, if any, of any other drugs administered to the patient.

(vi) A narrative description of the incident including approximate times and evolution of symptoms.

(vii) Additional information which the commission may require or request.

STANDARDS FOR PROFESSIONAL CONDUCT

WAC 246-919-605 Use of laser, light, radiofrequency, and plasma devices as applied to the skin. (1) For the purposes of this rule, laser, light, radiofrequency, and plasma devices (hereafter LLRP devices) are medical devices that:

- (a) Use a laser, noncoherent light, intense pulsed light, radiofrequency, or plasma to topically penetrate skin and alter human tissue; and
- (b) Are classified by the federal Food and Drug Administration as prescription devices.
- (2) Because an LLRP device penetrates and alters human tissue, the use of an LLRP device is the practice of medicine under RCW 18.71.011. The use of an LLRP device can result in complications such as visual impairment, blindness, inflammation, burns, scarring, hypopigmentation and hyperpigmentation.

(3) Use of medical devices using any form of energy to penetrate or alter human tissue for a purpose other than the purpose set forth in subsection (1) of this section constitutes surgery and is outside the scope of this section.

PHYSICIAN RESPONSIBILITIES

- (4) A physician must be appropriately trained in the physics, safety and techniques of using LLRP devices prior to using such a device, and must remain competent for as long as the device is used.
- (5) A physician must use an LLRP device in accordance with standard medical practice.
- (6) Prior to authorizing treatment with an LLRP device, a physician must take a history, perform an appropriate physical examination, make an appropriate diagnosis, recommend appropriate treatment, obtain the patient's informed consent (including informing the patient that a non-physician may operate the device), provide instructions for emergency and follow-up care, and prepare an appropriate medical record.
- (7) Regardless of who performs LLRP device treatment, the physician is ultimately responsible for the safety of the patient.
- (8) Regardless of who performs LLRP device treatment, the physician is responsible for assuring that each treatment is documented in the patient's medical record.

- (9) The physician must ensure that there is a quality assurance program for the facility at which LLRP device procedures are performed regarding the selection and treatment of patients. An appropriate quality assurance program shall include the following:
- (a) A mechanism to identify complications and untoward effects of treatment and to determine their cause;
- (b) A mechanism to review the adherence of supervised professionals to written protocols;
 - (c) A mechanism to monitor the quality of treatments;
- (d) A mechanism by which the findings of the quality assurance program are reviewed and incorporated into future protocols required by subsection (10)(d) of this section and physician supervising practices; and
- (e) Ongoing training to maintain and improve the quality of treatment and performance of treating professionals.

PHYSICIAN DELEGATION OF LLRP TREATMENT

(10) A physician who meets the above requirements may delegate an LLRP device procedure to a properly trained and licensed professional, whose licensure and scope of practice allow the use of an LLRP device, provided all the following conditions are met:

- (a) The treatment in no way involves surgery as that term is understood in the practice of medicine;
- (b) Such delegated use falls within the supervised professional's lawful scope of practice;
 - (c) The LLRP device is not used on the globe of the eye;
- (d) A physician has a written office protocol for the supervised professional to follow in using the LLRP device. A written office protocol must include at a minimum the following:
- (i) The identity of the individual physician authorized to use the device and responsible for the delegation of the procedure;
- (ii) A statement of the activities, decision criteria, and plan the supervised professional must follow when performing procedures delegated pursuant tounder this rule;
- (iii) Selection criteria to screen patients for the appropriateness
 of treatments;
- (iv) Identification of devices and settings to be used for patients who meet selection criteria;
- (v) Methods by which the specified device is to be operated and maintained;
- (vi) A description of appropriate care and follow-up for common complications, serious injury, or emergencies; and

- (vii) A statement of the activities, decision criteria, and plan the supervised professional shall follow when performing delegated procedures, including the method for documenting decisions made and a plan for communication or feedback to the authorizing physician concerning specific decisions made;
- (e) The supervised professional has appropriate training in, at a minimum, application techniques of each LLRP device, cutaneous medicine, indications and contraindications for such procedures, preprocedural and postprocedural care, potential complications and infectious disease control involved with each treatment;
- (f) The delegating physician ensures that the supervised professional uses the LLRP device only in accordance with the written office protocol, and does not exercise independent medical judgment when using the device;
- (g) The delegating physician shall be on the immediate premises during the patient's initial treatment and be able to treat complications, provide consultation, or resolve problems, if indicated. The supervised professional may complete the initial treatment if the physician is called away to attend to an emergency;

(h) Existing patients with an established treatment plan may continue to receive care during temporary absences of the delegating physician provided that there is a local back-up physician who satisfies the requirements of subsection (4) of this section. The local back-up physician must agree in writing to treat complications, provide consultation or resolve problems if medically indicated. The local back-up physician shall be reachable by phone and able to see the patient within sixty minutes. The delegating physician's absence from the site where the treatment occurs must be for brief and, intermittent or limited periods of time. The delegating physician's absence from the site where the treatment occurs cannot be an ongoing arrangement.

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(11) The use of, or the delegation of the use of, an LLRP device by a physician assistant is covered by WAC 246-918-125.

[Statutory Authority: RCW 18.71.017, 18.71A.020 and 18.130.050(12). WSR

07-03-177, § 246-919-605, filed 1/24/07, effective 3/1/07.]

WAC 246-919-606 Nonsurgical medical cosmetic procedures. (1) The purpose of this rule is to establish the duties and responsibilities of a physician who delegates the injection of medication or substances for cosmetic purposes or the use of prescription devices for cosmetic pur-

poses. These procedures can result in complications such as visual impairment, blindness, inflammation, burns, scarring, disfiguration, hypopigmentation and hyperpigmentation. The performance of these procedures is the practice of medicine under RCW 18.71.011(3).

- (2) This rule does not apply to:
- (a) Surgery;
- (b) The use of prescription lasers, noncoherent light, intense pulsed light, radiofrequency, or plasma as applied to the skin; this is covered in WAC 246-919-605 and 246-918-125;
- (c) The practice of a profession by a licensed health care professional under methods or means within the scope of practice permitted by such license;
 - (d) The use of nonprescription devices; and
 - (e) Intravenous therapy.
- (3) Definitions. These definitions apply throughout this section unless the context clearly requires otherwise.
- (a) "Nonsurgical medical cosmetic procedure" means a procedure or treatment that involves the injection of a medication or substance for cosmetic purposes, or the use of a prescription device for cosmetic purposes. Laser, light, radiofrequency and plasma devices that are used to topically penetrate the skin are devices used for cosmetic purposes,

but are excluded under subsection (2)(b) of this section, and are covered by WAC 246-919-605 and 246-918-125.

(b) "Physician" means an individual licensed under chapter 18.71
RCW.

(eb)—"Prescription device" means a device that the federal Food and Drug Administration has designated as a prescription device, and can be sold only to persons with prescriptive authority in the state in which they reside.

PHYSICIAN RESPONSIBILITIES

- (4) A physician must be fully and appropriately trained in a non-surgical medical cosmetic procedure prior to performing the procedure or delegating the procedure. The physician must keep a record of his or her training in the office and available for review upon request by a patient or a representative of the commission.
- (5) Prior to authorizing a nonsurgical medical cosmetic procedure, a physician must:
 - (a) Take a history;
 - (b) Perform an appropriate physical examination;
 - (c) Make an appropriate diagnosis;
 - (d) Recommend appropriate treatment;
 - (e) Obtain the patient's informed consent;

- (f) Provide instructions for emergency and follow-up care; and
- (g) Prepare an appropriate medical record.
- (6) Regardless of who performs the nonsurgical medical cosmetic procedure, the physician is ultimately responsible for the safety of the patient.
- (7) Regardless of who performs the nonsurgical medical cosmetic procedure, the physician is responsible for ensuring that each treatment is documented in the patient's medical record.
- (8) The physician must ensure that there is a quality assurance program for the facility at which nonsurgical medical cosmetic procedures are performed regarding the selection and treatment of patients. An appropriate quality assurance program must include the following:
- (a) A mechanism to identify complications and untoward effects of treatment and to determine their cause;
- (b) A mechanism to review the adherence of supervised health care professionals to written protocols;
 - (c) A mechanism to monitor the quality of treatments;
- (d) A mechanism by which the findings of the quality assurance program are reviewed and incorporated into future protocols required by subsection (10)(d) of this section and physician supervising practices; and

- (e) Ongoing training to maintain and improve the quality of treatment and performance of supervised health care professionals.
- (9) A physician may not sell or give a prescription device to an individual who does not possess prescriptive authority in the state in which the individual resides or practices.
- (10) The physician must ensure that all equipment used for procedures covered by this section is inspected, calibrated, and certified as safe according to the manufacturer's specifications.

PHYSICIAN DELEGATION

- (11) A physician who meets the above requirements may delegate a nonsurgical medical cosmetic procedure to a properly trained physician assistant, registered nurse or licensed practical nurse, provided all the following conditions are met:
- (a) The treatment in no way involves surgery as that term is understood in the practice of medicine;
- (b) The physician delegates procedures that are within the delegate's lawful scope of practice;
 - (c) The delegate has appropriate training in, at a minimum:
 - (i) Techniques for each procedure;
 - (ii) Cutaneous medicine;
 - (iii) Indications and contraindications for each procedure;

- (iv) Preprocedural and postprocedural care;
- (v) Recognition and acute management of potential complications that may result from the procedure; and
 - (vi) Infectious disease control involved with each treatment.
- (d) The physician has a written office protocol for the delegate to follow in performing the nonsurgical medical cosmetic procedure. A written office protocol must include, at a minimum, the following:
- (i) The identity of the physician responsible for the delegation of the procedure;
- (ii) Selection criteria to screen patients for the appropriateness of treatment;
- (iii) A description of appropriate care and follow-up for common complications, serious injury, or emergencies; and
- (iv) A statement of the activities, decision criteria, and plan the delegate shall follow when performing delegated procedures, including the method for documenting decisions made and a plan for communication or feedback to the authorizing physician concerning specific decisions made.
- (e) The physician ensures that the delegate performs each procedure in accordance with the written office protocol;

- (f) Each patient signs a consent form prior to treatment that lists foreseeable side effects and complications, and the identity and license of the delegate or delegates who will perform the procedure; and
- (g) Each delegate performing a procedure covered by this section must be readily identified by a name tag or similar means so that the patient understands the identity and license of the treating delegate.
- (12) If a physician delegates the performance of a procedure that uses a medication or substance that the federal Food and Drug Administration has not approved, or that the federal Food and Drug Administration has not approved for the particular purpose for which it is used, the physician must be on-site during the entire duration of the procedure.
- (13) If a physician delegates the performance of a procedure that uses a medication or substance that is approved by the federal Food and Drug Administration for the particular purpose for which it is used, the physician need not be on-site during the procedure, but must be reachable by phone and able to respond within thirty minutes to treat complications.
- (14) If the physician is unavailable to supervise a delegate as required by this section, the physician must make arrangements for an alternate physician to provide the necessary supervision. The alternate

supervisor must be familiar with the protocols in use at the site, will be accountable for adequately supervising the treatment under the protocols, and must have comparable training as the primary supervising physician.

- (15) A physician performing or delegating nonsurgical cosmetic procedures may not sponsor more than three physician assistants at any one time.
- (16) A physician may not permit a delegate to further delegate the performance of a nonsurgical medical cosmetic procedure to another individual.

[Statutory Authority: RCW 18.71.017, 18.71A.020 and 18.130.050(4). WSR 10-11-001, § 246-919-606, filed 5/5/10, effective 6/5/10.]

WAC 246-919-610 Use of drugs or autotransfusion to enhance athletic ability. (1) A physician challmay not prescribe, administer or dispense anabolic steroids, growth hormones, testosterone or its analogs, human chorionic gonadotropin (HCG), other hormones, or any form of autotransfusion for the purpose of enhancing athletic ability.

(2) A physician shall complete and maintain patient medical records which accurately reflect the prescribing, administering or dispensing

of any substance or drug described in this rule or any form of autotransfusion. Patient medical records shall indicate the diagnosis and purpose for which the substance, drug or autotransfusion is prescribed, administered or dispensed and any additional information upon which the diagnosis is based.

(3) A violation of any provision of this rule shall constitutes grounds for disciplinary action under RCW 18.130.180(7). A violation of subsection (1) of this section shall also constitutes grounds for disciplinary action under RCW 18.130.180(6).

[Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, § 246-919-610, filed 1/17/96, effective 2/17/96.]

wac 246-919-620 Cooperation with investigation. (1) A licenseephysician must comply with a request, under RCW 70.02.050, for health care records or documents from an investigator who is acting on behalf of the disciplining authority pursuant tounder RCW 18.130.050(2). The physician must by submitting the requested items within fourteen_twenty-one_calendar days of receipt of the request by the licenseephysician or the licenseephysician's attorney, whichever is first. If the licenseephysician cian fails to comply with the request within fourteen_twenty-one_calendar

days, the investigator shall contact the licenseephysician or the licenseephysician attorney by letter as a reminder.

- (a) Investigators may extend the time for response if the licenseephysician requests an extension for good cause for a period not to
 exceed seven—thirty calendar days. Other requests for extension may be
 granted by the commission chair or the commission's designee.
- (b) If the licenseephysician fails to comply with the request within three business days after the receipt of the written reminder, a statement of charges shall be issued pursuant tounder RCW 18.130.180(8) and, if there is sufficient evidence to support additional charges, those charges may be included in the statement of charges.
- (2) A licenseephysician must comply with a request for nonhealth care records or documents from an investigator who is acting on behalf of the commission pursuant tounder RCW 18.130.050(2) by submitting the requested items within fourteen—twenty-one calendar days of receipt of the request by the licenseephysician or the licenseephysician's attorney, whichever is first. If the licenseephysician fails to comply with the request within fourteen—twenty-one calendar days, the investigator shall contact the licenseephysician or the licensee's attorney by letter as a reminder.

- (a) Investigators may extend the time for response if the licenseephysician requests an extension for good cause for a period not to exceed seven thirty calendar days. Other requests for extension may be granted by the commission chair or the commission's designee.
- (b) If the licenseephysician fails to comply with the request within three business days after the receipt of the written reminder, then a subpoena shall be served upon the licenseephysician to obtain the requested items.
- (c) If the <u>licenseephysician</u> fails to comply with the subpoena, a statement of charges shall be issued <u>pursuant tounder</u> RCW 18.130.180(8) and, if there is sufficient evidence to support additional charges, then those charges may be included in the statement of charges.
- (3) A licenseephysician must comply with a request for information from an investigator who is acting on behalf of the commission pursuant tounder RCW 18.130.050(2). This information may include, but is not limited to, an explanation of the matter under investigation, curriculum vitae, continuing medical education credits, malpractice action summaries, or hospital affiliations, or nonhealth care records or documents. The licenseephysician will submit the requested information within fourteen calendar days of receipt of the request by the licenseephysician

or the licenseephysician sattorney, whichever is first. If the licenseephysician fails to comply with the request within fourteen calendar days, the investigator shall contact the licenseephysician or the licenseephysician 's attorney by letter as a reminder.

- (a) Investigators may extend the time for response if the licenseephysician requests an extension for a period not to exceed seven
 calendar days. Other requests for extension may be granted by the commission chair or the commission's designee.
- (b) If the physician fails to comply with the request within three business days after the receipt of the written reminder, then a subpoena shall be served upon the physician to obtain the requested items.
- (c) If the physician fails to comply with the subpoena, a statement of charges shall be issued under RCW 18.130.180(8) and, if there is sufficient evidence to support additional charges, then those charges may be included in the statement of charges. (b) If the licensee fails to comply with the written reminder within three business days after the receipt of the reminder, a statement of charges shall be issued pursuant to RCW 18.130.180(8) and, if there is sufficient evidence to support additional charges, then those charges may be included in the statement of charges.

Commented [UD(25]: From Heather: Do you need three sections? Is there a reason this "information" is carved out? Is the difference because of the subpoena issued in section (2)? It seems to me you just need a section for health care records (1) and for non-health care records (2). Section (3) could be worked in to section (2) by adding the examples listed here. If you determine that you need three sections, I would suggest making the timelines the 21/30 days to match the others and 18.130.230.

Commented [UD(26]: Ask investigators what their pro-

(4) In negotiating a settlement on a statement of charges based on RCW 18.130.180(8), the reviewing commission member may take into consideration whether the licenseephysician has complied with the request after the statement of charges has been issued. Any settlement proposal shall be presented to the commission or a duly constituted panel of the commission for a decision on ratification and until ratified, the settlement is not final.

[Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, § 246-919-620, filed 1/17/96, effective 2/17/96.]

WAC 246-919-630 Sexual misconduct. (1) The following definitions apply throughout this section unless the context clearly requires otherwise.

(a) "Patient" means a person who is receiving health care or treatment, or has received health care or treatment without a termination of the physician-patient relationship. The determination of when a person is a patient is made on a case-by-case basis with consideration given to a number of factors, including the nature, extent and context of the professional relationship between the physician and the person. The fact that a person is not actively receiving treatment or professional services is not the sole determining factor.

(b) "Physician" means a person licensed to practice medicine and

surgery under chapter 18.71 RCW.

(eb) "Key third party" means a person in a close personal relationship with the patient and includes, but is not limited to, spouses, partners, parents, siblings, children, guardians and proxies.

(2) A physician shall not engage in sexual misconduct with a current patient or a key third party. A physician engages in sexual misconduct when he or she engages in the following behaviors with a patient or key third party:

- (a) Sexual intercourse or genital to genital contact;
- (b) Oral to genital contact;
- (c) Genital to anal contact or oral to anal contact;
- (d) Kissing in a romantic or sexual manner;
- (e) Touching breasts, genitals or any sexualized body part for any purpose other than appropriate examination or treatment;
- (f) Examination or touching of genitals without using gloves, except for examinations of an infant or prepubescent child when clinically appropriate;
 - (g) Not allowing a patient the privacy to dress or undress;
- (h) Encouraging the patient to masturbate in the presence of the physician or masturbation by the physician while the patient is present;

Commented [UD(27]: Already defined. Removal ap-

Commented [UD(28]: Approved 11/14/18

- (i) Offering to provide practice-related services, such as medications, in exchange for sexual favors;
 - (j) Soliciting a date;
- (k) Engaging in a conversation Communicating regarding the sexual history, preferences or fantasies of the physician.
- (3) A physician shall not engage in any of the conduct described in subsection (2) of this section with a former patient or key third party if the physician:
- (a) Uses or exploits the trust, knowledge, influence, or emotions derived from the professional relationship; or
- (b) Uses or exploits privileged information or access to privileged information to meet the physician's personal or sexual needs.
- (4) Sexual misconduct also includes sexual contact with any person involving force, intimidation, or lack of consent; or a conviction of a sex offense as defined in RCW 9.94A.030.
- (5) To determine whether a patient is a current patient or a former patient, the commission will analyze each case individually, and will consider a number of factors, including, but not limited to, the following:
 - (a) Documentation of formal termination;
 - (b) Transfer of the patient's care to another health care provider;

- (c) The length of time that has passed;
- (d) The length of time of the professional relationship;
- (e) The extent to which the patient has confided personal or private information to the physician;
 - (f) The nature of the patient's health problem;
 - (g) The degree of emotional dependence and vulnerability.
- (6) This section does not prohibit conduct that is required for medically recognized diagnostic or treatment purposes if the conduct meets the standard of care appropriate to the diagnostic or treatment situation.
- (7) It is not a defense that the patient, former patient, or key third party initiated or consented to the conduct, or that the conduct occurred outside the professional setting.
- (8) A violation of any provision of this rule shall constitute grounds for disciplinary action.

[Statutory Authority: RCW 18.71.017, 18.130.062, and Executive Order 06-03. WSR 16-06-010, § 246-919-630, filed 2/18/16, effective 3/20/16. Statutory Authority: RCW 18.130.180, 18.71.017, and 18.71A.020. WSR 06-03-028, § 246-919-630, filed 1/9/06, effective 2/9/06.]

WAC 246-919-640 Abuse. (1) A physician commits unprofessional conduct if the physician abuses a patient. A physician abuses a patient when he or she:

- (a) Makes statements regarding the patient's body, appearance, sexual history, or sexual orientation that have no legitimate medical or therapeutic purpose;
 - (b) Removes a patient's clothing or gown without consent;
- (c) Fails to treat an unconscious or deceased patient's body or property respectfully; or
- (d) Engages in any conduct, whether verbal or physical, which unreasonably demeans, humiliates, embarrasses, threatens, or harms a patient.
- (2) A violation of any provision of this rule shall constitute grounds for disciplinary action.

[Statutory Authority: RCW 18.130.180, 18.71.017, and 18.71A.020. WSR 06-03-028, § 246-919-640, filed 1/9/06, effective 2/9/06.]

MANDATORY REPORTING

WAC 246-919-700 Mandatory reporting. The commission adopts the rules for mandatory reporting in chapter 246-16 WAC. (1) All reports required by these regulations shall be submitted to the commission as

Commented [UD(29]: The Commission approved repeal of WAC 246-919-700 through -770 back in 2012. I believe this was not done because of the moratorium on rulemaking.

Commented [UD(30R29]: 7/11/18 – Approved repealing these sections and replacing with: Any person including, but not limited to, a physician and surgeon, health care facility, or governmental agency shall always report in compliance with the uniform mandatory reporting rules found in WAC 246-16-200 through 246-16-270.

Commented [FM(31R29]: How about stating: The commission adopts the rules for mandatory reporting in chapter 246-16 WAC.

Commented [UD(32R29]: Approved 11/14/18

soon as possible, but not later than sixty days after a determination is made.

- (2) A report should contain the following information if known:
- (a) The name, address and telephone number of the person making the
- (b) The name, address and telephone numbers of the physician being reported;
- (c) The case number of any patient whose treatment is a subject of the report;
- (d) A brief description or summary of the facts which gave rise to issuance of the report, including dates of occurrences;
- (e) If court action is involved, the name of the court in which the action is filed along with the date of filing and docket number; and
- (f) Any further information which would aid the evaluation of the report.
- (3) The mandatory reporting shall not act as a waiver of confidentiality of medical records and committee reports. The information reported or disclosed shall be kept for the confidential use of the commission as provided in the Uniform Disciplinary Act and shall not be subject to subpoena or discovery proceedings in any civil action as provided in RCW 4.24.250, and shall be exempt from public disclosure

pursuant tounder chapter 42.17 RCW except for review as provided in RCW 18.71.0195.

{Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073,
\$ 246-919-700, filed 1/17/96, effective 2/17/96.}

WAC 246-919-710 Mandatory reporting requirement satisfied. The requirement for a report to the commission under RCW 18.71.0193(1) may be satisfied by submitting the report to the impaired physician program approved by the commission under this chapter.

{Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, ... \$ 246-919-710, filed 1/17/96, effective 2/17/96.}

wac 246 919 730 Medical associations or societies. The presidents or chief executive officer of any medical association or society within this state shall report to the commission when a medical society hearing panel or committee determines that a physician has committed unprofessional conduct or that a physician may not be able to practice medicine with reasonable skill and safety to patients as the result of any mental or physical condition and constitutes an apparent risk to the public health, safety or welfare. The report required by this subsection shall be made without regard to whether the license holder appeals, accepts or acts upon the determination made by the association or society. Notification of appeal shall be included.

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Formatted: Indent: First line: 0.5" [Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, 919-730, filed 1/17/96, effective 2/17/96.] Formatted: Space Before: 0 pt WAC 246-919-740 Health care service contractors and disability insurance carriers. The executive officer of every health 48.21A and 48.44 RCW operating in the state of Washington, shall report services not actually provided. Formatted: Indent: First line: 0.5" 919-740, filed 1/17/96, effective 2/17/96.1 Formatted: Space Before: 0 pt WAC 246-919-750 Courts. The commission requests the assistance of -judgments and all convictions of licensed medical doctors, other than minor traffic violations. Formatted: Indent: First line: 0.5" {Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, 919-750, filed 1/17/96, effective 2/17/96.] Formatted: Space Before: 0 pt WAC 246-919-760 State and federal agencies. The commission assistance of executive officers of any state or federal program

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to provide patient care services, to report to the commission whenever such a physician has been judged to have demonstrated his/her incompetency or negligence in the practice of medicine, or has otherwise committed unprofessional conduct; or is a mentally or physically disabled physician unable to practice with reasonable skill and safety by reason of a mental or physical condition.

[Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, \$246-919-760, filed 1/17/96, effective 2/17/96.]

wac 246-919-770 Professional standards review organizations. When authorized by federal law, every professional standards review organization operating within the state of Washington shall report to the commission any determinations that a physician has engaged or is engaging in consistent, excessive utilization of any medical or surgical test, treatment or procedure when such procedures are clearly not called for under the circumstances in which such services were provided.

\frac{\{\text{15tatutory Authority: New 18.71.017 and 18.71A.020. WSR 96-03-073\}}{\frac{\}{246-919-770\}, \text{filed 1/17/96\}, \text{effective 2/17/96\}}

PAIN MANAGEMENT

WAC 246-919-850 Pain management—Intent. These rules govern the use of opioids in the treatment of patients for chronic noncancer pain. Formatted: Indent: First line: 0.5"

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The Washington state medical quality assurance commission (commission) recognizes that principles of quality medical practice dictate that the people of the state of Washington have access to appropriate and effective pain relief. The appropriate application of up-to-date knowledge and treatment modalities can serve to improve the quality of life for those patients who suffer from pain as well as reduce the morbidity and costs associated with untreated or inappropriately treated pain. For the purposes of this rule, the inappropriate treatment of pain includes nontreatment, undertreatment, overtreatment, and the continued use of ineffective treatments.

The diagnosis and treatment of pain is integral to the practice of medicine. The commission encourages physicians to view pain management as a part of quality medical practice for all patients with pain, acute or chronic, and it is especially urgent for patients who experience pain as a result of terminal illness. All physicians should become knowledgeable about assessing patients' pain and effective methods of pain treatment, as well as statutory requirements for prescribing controlled substances. Accordingly, this rule has been developed to clarify the commission's position on pain control, particularly as related to the use of controlled substances, to alleviate physician uncertainty and to encourage better pain management.

Inappropriate pain treatment may result from a physician's lack of knowledge about pain management. Fears of investigation or sanction by federal, state, and local agencies may also result in inappropriate treatment of pain. Appropriate pain management is the treating physician's responsibility. As such, the commission will consider the inappropriate treatment of pain to be a departure from standards of practice and will investigate such allegations, recognizing that some types of pain cannot be completely relieved, and taking into account whether the treatment is appropriate for the diagnosis.

The commission recognizes that controlled substances including opioid analgesics may be essential in the treatment of acute pain due to trauma or surgery and chronic pain, whether due to cancer or noncancer origins. The commission will refer to current clinical practice guidelines and expert review in approaching cases involving management of pain. The medical management of pain should consider current clinical knowledge and scientific research and the use of pharmacologic and nonpharmacologic modalities according to the judgment of the physician. Pain should be assessed and treated promptly, and the quantity and frequency of doses should be adjusted according to the intensity, duration of the pain, and treatment outcomes. Physicians should recognize that tolerance and physical dependence are normal consequences of sustained use of opioid analgesics and are not the same as addiction.

The commission is obligated under the laws of the state of Washington to protect the public health and safety. The commission recognizes that the use of opioid analgesics for other than legitimate medical purposes poses a threat to the individual and society and that the inappropriate prescribing of controlled substances, including opioid analgesics, may lead to drug diversion and abuse by individuals who seek them for other than legitimate medical use. Accordingly, the commission expects that physicians incorporate safeguards into their practices to minimize the potential for the abuse and diversion of controlled substances.

Physicians should not fear disciplinary action from the commission for ordering, prescribing, dispensing or administering controlled substances, including opioid analgesics, for a legitimate medical purpose and in the course of professional practice. The commission will consider prescribing, ordering, dispensing or administering controlled substances for pain to be for a legitimate medical purpose if based on sound clinical judgment. All such prescribing must be based on clear documentation of unrelieved pain. To be within the usual course of professional practice, a physician-patient relationship must exist and the prescribing should be based on a diagnosis and documentation of unrelieved pain.

Compliance with applicable state or federal law is required.

The commission will judge the validity of the physician's treatment of the patient based on available documentation, rather than solely on the quantity and duration of medication administration. The goal is to control the patient's pain while effectively addressing other aspects of the patient's functioning, including physical, psychological, social, and work-related factors.

These rules are designed to assist practitioners in providing appropriate medical care for patients. They are not inflexible rules or rigid practice requirements and are not intended, nor should they be used, to establish a legal standard of care outside the context of the medical quality assurance committee's jurisdiction.

The ultimate judgment regarding the propriety of any specific procedure or course of action must be made by the practitioner based on all the circumstances presented. Thus, an approach that differs from the rules, standing alone, does not necessarily imply that the approach was below the standard of care. To the contrary, a conscientious practitioner may responsibly adopt a course of action different from that set forth in the rules when, in the reasonable judgment of the practitioner, such

course of action is indicated by the condition of the patient, limitations of available resources, or advances in knowledge or technology subsequent to publication of these rules. However, a practitioner who employs an approach substantially different from these rules is advised to document in the patient record information sufficient to justify the approach taken.

The practice of medicine involves not only the science, but also the art of dealing with the prevention, diagnosis, alleviation, and treatment of disease. The variety and complexity of human conditions make it impossible to always reach the most appropriate diagnosis or to predict with certainty a particular response to treatment.

Therefore, it should be recognized that adherence to these rules will not assure an accurate diagnosis or a successful outcome. The sole purpose of these rules is to assist practitioners in following a reasonable course of action based on current knowledge, available resources, and the needs of the patient to deliver effective and safe medical care.

[Statutory Authority: RCW 18.71.450, 18.71A.100, 18.71.017, 18.71A.020. WSR 11-12-025, § 246-919-850, filed 5/24/11, effective 1/2/12.]

WAC 246-919-851 Exclusions. The rules adopted under WAC 246-919-850 through 246-919-863 do not apply:

- (1) To the provision of palliative, hospice, or other end-of-life care; or
- (2) To the management of acute pain caused by an injury or surgical procedure.

[Statutory Authority: RCW 18.71.450, 18.71A.100, 18.71.017, 18.71A.020. WSR 11-12-025, § 246-919-851, filed 5/24/11, effective 1/2/12.]

WAC 246-919-852 Definitions. The definitions in WAC 246-919-850 through 246-919-863 apply unless the context clearly requires otherwise.

- (1) "Acute pain" means the normal, predicted physiological response to a noxious chemical, thermal, or mechanical stimulus and typically is associated with invasive procedures, trauma, and disease. It is generally time-limited, often less than three months in duration, and usually less than six months.
- (2) "Addiction" means a primary, chronic, neurobiologic disease with genetic, psychosocial, and environmental factors influencing its development and manifestations. It is characterized by behaviors that include:

- (a) Impaired control over drug use;
- (b) Craving;
- (c) Compulsive use; or
- (d) Continued use despite harm.
- (3) "Chronic noncancer pain" means a state in which noncancer pain persists beyond the usual course of an acute disease or healing of an injury, or that may or may not be associated with an acute or chronic pathologic process that causes continuous or intermittent pain over months or years.
- (4) "Comorbidity" means a preexisting or coexisting physical or psychiatric disease or condition.
- (5) "Episodic care" means medical care provided by a practitioner other than the designated primary care practitioner in the acute care setting, for example, urgent care or emergency department.
- (6) "Hospice" means a model of care that focuses on relieving symptoms and supporting patients with a life expectancy of six months or less. Hospice involves an interdisciplinary approach to provide health care, pain management, and emotional and spiritual support. The emphasis is on comfort, quality of life and patient and family support. Hospice can be provided in the patient's home as well as freestanding

hospice facilities, hospitals, nursing homes, or other long-term care facilities.

- (7) "Morphine equivalent dose" means a conversion of various opioids to a morphine equivalent dose by the use of accepted conversion tables.
- (8) "Multidisciplinary pain clinic" means a clinic or office that provides comprehensive pain management and includes care provided by multiple available disciplines or treatment modalities, for example, medical care through physicians, physician assistants, osteopathic physicians, osteopathic physician assistants, advanced registered nurse practitioners, and physical therapy, occupational therapy, or other complementary therapies.
- (9) "Palliative" means care that improves the quality of life of patients and their families facing life-threatening illness. With palliative care particular attention is given to the prevention, assessment, and treatment of pain and other symptoms, and to the provision of psychological, spiritual, and emotional support.

[Statutory Authority: RCW 18.71.450, 18.71A.100, 18.71.017, 18.71A.020. WSR 11-12-025, § 246-919-852, filed 5/24/11, effective 1/2/12.]

WAC 246-919-853 Patient evaluation. The physician shall obtain, evaluate, and document the patient's health history and physical examination in the health record prior to treating for chronic noncancer pain.

- (1) The patient's health history shall include:
- (a) Current and past treatments for pain;
- (b) Comorbidities; and
- (c) Any substance abuse.
- (2) The patient's health history should include:
- (a) A review of any available prescription monitoring program or emergency department-based information exchange; and
- (b) Any relevant information from a pharmacist provided to a physician.
 - (3) The initial patient evaluation shall include:
 - (a) Physical examination;
 - (b) The nature and intensity of the pain;
 - (c) The effect of the pain on physical and psychological function;
 - (d) Medications including indication(s), date, type, dosage, and

quantity prescribed;

- (e) A risk screening of the patient for potential comorbidities and risk factors using an appropriate screening tool. The screening should address:
 - (i) History of addiction;
 - (ii) Abuse or aberrant behavior regarding opioid use;
 - (iii) Psychiatric conditions;
- (iv) Regular concomitant use of benzodiazepines, alcohol, or other central nervous system medications;
 - (v) Poorly controlled depression or anxiety;
- (vi) Evidence or risk of significant adverse events, including falls or fractures;
- (vii) Receipt of opioids from more than one prescribing practitioner or practitioner group;
 - (viii) Repeated visits to emergency departments seeking opioids;
 - (ix) History of sleep apnea or other respiratory risk factors;
 - (x) Possible or current pregnancy; and
 - (xi) History of allergies or intolerances.
 - (4) The initial patient evaluation should include:
 - (a) Any available diagnostic, the rapeutic, and laboratory results;

and

(b) Any available consultations.

- (5) The health record shall be maintained in an accessible manner, readily available for review, and should include:
 - (a) The diagnosis, treatment plan, and objectives;
- (b) Documentation of the presence of one or more recognized indications for the use of pain medication;
 - (c) Documentation of any medication prescribed;
 - (d) Results of periodic reviews;
- (e) Any written agreements for treatment between the patient and the physician; and
- (f) The physician's instructions to the patient. [Statutory Authority: RCW 18.71.450, 18.71A.100, 18.71.017, 18.71A.020. WSR 11-12-025, § 246-919-853, filed 5/24/11, effective 1/2/12.]

WAC 246-919-854 Treatment plan. (1) The written treatment plan shall state the objectives that will be used to determine treatment success and shall include, at a minimum:

- (a) Any change in pain relief;
- (b) Any change in physical and psychosocial function; and
- (c) Additional diagnostic evaluations or other planned treatments.

- (2) After treatment begins the physician should adjust drug therapy to the individual health needs of the patient. The physician shall include indications for medication use on the prescription and require photo identification of the person picking up the prescription in order to fill. The physician shall advise the patient that it is the patient's responsibility to safeguard all medications and keep them in a secure location.
- (3) Other treatment modalities or a rehabilitation program may be necessary depending on the etiology of the pain and the extent to which the pain is associated with physical and psychosocial impairment. [Statutory Authority: RCW 18.71.450, 18.71A.100, 18.71.017, 18.71A.020. WSR 11-12-025, § 246-919-854, filed 5/24/11, effective 1/2/12.]

WAC 246-919-855 Informed consent. The physician shall discuss the risks and benefits of treatment options with the patient, persons designated by the patient, or with the patient's surrogate or guardian if the patient is without health care decision-making capacity. [Statutory Authority: RCW 18.71.450, 18.71A.100, 18.71.017, and 18.71A.020. WSR 11-12-025, § 246-919-855, filed 5/24/11, effective 1/2/12.]

wac 246-919-856 Written agreement for treatment. Chronic noncancer pain patients should receive all chronic pain management prescriptions from one physician and one pharmacy whenever possible. If the patient is at high risk for medication abuse, or has a history of substance abuse, or psychiatric comorbidities, the prescribing physician shall use a written agreement for treatment with the patient outlining patient responsibilities. This written agreement for treatment shall include:

- (1) The patient's agreement to provide biological samples for urine/serum medical level screening when requested by the physician;
- (2) The patient's agreement to take medications at the dose and frequency prescribed with a specific protocol for lost prescriptions and early refills;
- (3) Reasons for which drug therapy may be discontinued (e.g., violation of agreement);
- (4) The requirement that all chronic pain management prescriptions are provided by a single prescriber or multidisciplinary pain clinic and dispensed by a single pharmacy or pharmacy system;
- (5) The patient's agreement to not abuse alcohol or use other medically unauthorized substances;
 - (6) A written authorization for:

- (a) The physician to release the agreement for treatment to local emergency departments, urgent care facilities, and pharmacies; and
- (b) Other practitioners to report violations of the agreement back to the physician;
- (7) A written authorization that the physician may notify the proper authorities if he or she has reason to believe the patient has engaged in illegal activity;
- (8) Acknowledgment that a violation of the agreement may result in a tapering or discontinuation of the prescription;
- (9) Acknowledgment that it is the patient's responsibility to safeguard all medications and keep them in a secure location; and
- (10) Acknowledgment that if the patient violates the terms of the agreement, the violation and the physician's response to the violation will be documented, as well as the rationale for changes in the treatment plan.

[Statutory Authority: RCW 18.71.450, 18.71A.100, 18.71.017, and 18.71A.020. WSR 11-12-025, § 246-919-856, filed 5/24/11, effective 1/2/12.]

WAC 246-919-857 Periodic review. The physician shall periodically review the course of treatment for chronic noncancer pain, the patient's

state of health, and any new information about the etiology of the pain. Generally, periodic reviews shall take place at least every six months. However, for treatment of stable patients with chronic noncancer pain involving nonescalating daily dosages of forty milligrams of a morphine equivalent dose (MED) or less, periodic reviews shall take place at least annually.

- (1) During the periodic review, the physician shall determine:
- (a) Patient's compliance with any medication treatment plan;
- (b) If pain, function, or quality of life have improved or diminished using objective evidence, considering any available information from family members or other caregivers; and
- (c) If continuation or modification of medications for pain management treatment is necessary based on the physician's evaluation of progress towards treatment objectives.
- (2) The physician shall assess the appropriateness of continued use of the current treatment plan if the patient's progress or compliance with current treatment plan is unsatisfactory. The physician shall consider tapering, changing, or discontinuing treatment when:
 - (a) Function or pain does not improve after a trial period;
 - (b) There is evidence of significant adverse effects;
 - (c) Other treatment modalities are indicated; or

- (d) There is evidence of misuse, addiction, or diversion.
- (3) The physician should periodically review information from any available prescription monitoring program or emergency department-based information exchange.
- (4) The physician should periodically review any relevant information from a pharmacist provided to the physician. [Statutory Authority: RCW 18.71.450, 18.71A.100, 18.71.017, 18.71A.020. WSR 11-12-025, § 246-919-857, filed 5/24/11, effective 1/2/12.]

WAC 246-919-858 Long-acting opioids, including methadone. Longacting opioids, including methadone, should only be prescribed by a physician who is familiar with its risks and use, and who is prepared to conduct the necessary careful monitoring. Special attention should be given to patients who are initiating such treatment. The physician prescribing long-acting opioids or methadone should have a one-time (lifetime) completion of at least four hours of continuing education relating to this topic.

[Statutory Authority: RCW 18.71.450, 18.71A.100, 18.71.017, and 18.71A.020. WSR 11-12-025, § 246-919-858, filed 5/24/11, effective 1/2/12.]

wac 246-919-859 Episodic care. (1) When evaluating patients for episodic care, such as emergency or urgent care, the physician should review any available prescription monitoring program, emergency department-based information exchange, or other tracking system.

- (2) Episodic care practitioners should avoid providing opioids for chronic pain management. However, if opioids are provided, the practitioner should limit the use of opioids for a chronic noncancer pain patient to the minimum amount necessary to control the pain until the patient can receive care from a primary care practitioner.
- (3) Prescriptions for opioids written by an episodic care practitioner shall include indications for use or the International Classification of Diseases (ICD) code and shall be written to require photo identification of the person picking up the prescription in order to fill.
- (4) If a patient has signed a written agreement for treatment and has provided a written authorization to release the agreement under WAC 246-919-856(6) to episodic care practitioners, then the episodic care practitioner should report known violations of the agreement back to the patient's treatment practitioner who provided the agreement for treatment.

[Statutory Authority: RCW 18.71.450, 18.71A.100, 18.71.017, 18.71A.020. WSR 11-12-025, § 246-919-859, filed 5/24/11, effective 1/2/12.]

WAC 246-919-860 Consultation—Recommendations and requirements. (1)

The physician shall consider, and document the consideration, referring the patient for additional evaluation and treatment as needed to achieve treatment objectives. Special attention should be given to those chronic noncancer pain patients who are under eighteen years of age, or who are at risk for medication misuse, abuse, or diversion. The management of pain in patients with a history of substance abuse or with comorbid psychiatric disorders may require extra care, monitoring, documentation, and consultation with, or referral to, an expert in the management of such patients.

(2) The mandatory consultation threshold for adults is one hundred twenty milligrams morphine equivalent dose (MED)(oral). In the event a physician prescribes a dosage amount that meets or exceeds the consultation threshold of one hundred twenty milligrams MED (orally) per day, a consultation with a pain management specialist as described in WAC 246-919-863 is required, unless the consultation is exempted under WAC 246-919-861 or 246-919-862. Great caution should be used when prescribing opioids to children with chronic noncancer pain and appropriate referrals to a specialist is encouraged.

- (a) The mandatory consultation shall consist of at least one of the following:
- (i) An office visit with the patient and the pain management specialist;
- (ii) A telephone consultation between the pain management specialist and the physician;
- (iii) An electronic consultation between the pain management specialist and the physician; or
- (iv) An audio-visual evaluation conducted by the pain management specialist remotely, where the patient is present with either the physician or a licensed health care practitioner designated by the physician or the pain management specialist.
- (b) A physician shall document each mandatory consultation with the pain management specialist. Any written record of the consultation by the pain management specialist shall be maintained as a patient record by the specialist. If the specialist provides a written record of the consultation to the physician, the physician shall maintain it as part of the patient record.

(3) Nothing in this chapter shall limit any person's ability to contractually require a consultation with a pain management specialist at any time. For the purposes of WAC 246-919-850 through 246-919-863, "person" means an individual, a trust or estate, a firm, a partnership, a corporation (including associations, joint stock companies, and insurance companies), the state, or a political subdivision or instrumentality of the state, including a municipal corporation or a hospital district.

[Statutory Authority: RCW 18.71.450, 18.71A.100, 18.71.017, 18.71A.020. WSR 11-12-025, § 246-919-860, filed 5/24/11, effective 1/2/12.]

WAC 246-919-861 Consultation-Exemptions for exigent and special circumstances. A physician is not required to consult with a pain management specialist as described in WAC 246-919-863 when he or she has documented adherence to all standards of practice as defined in WAC 246-919-850 through 246-919-863 and when any one or more of the following conditions apply:

- (1) The patient is following a tapering schedule;
- (2) The patient requires treatment for acute pain which may or may not include hospitalization, requiring a temporary escalation in opioid

dosage, with expected return to or below their baseline dosage level; or

- (3) The physician documents reasonable attempts to obtain a consultation with a pain management specialist and the circumstances justifying prescribing above one hundred twenty milligrams morphine equivalent dose (MED) per day without first obtaining a consultation; or
- (4) The physician documents the patient's pain and function is stable and the patient is on a nonescalating dosage of opioids.

 [Statutory Authority: RCW 18.71.450, 18.71A.100, 18.71.017, and 18.71A.020. WSR 11-12-025, § 246-919-861, filed 5/24/11, effective 1/2/12.]

WAC 246-919-862 Consultation—Exemptions for the physician. The physician is exempt from the consultation requirement in WAC 246-919-860 if one or more of the following qualifications are met:

- (1) The physician is a pain management specialist under WAC 246-919-863; or
- (2) The physician has successfully completed, within the last two years, a minimum of twelve (Category I) continuing education hours on chronic pain management with at least two of these hours dedicated to long-acting opioids; or

- (3) The physician is a pain management practitioner working in a multidisciplinary chronic pain treatment center, or a multidisciplinary academic research facility; or
- (4) The physician has a minimum three years of clinical experience in a chronic pain management setting, and at least thirty percent of his or her current practice is the direct provision of pain management care. [Statutory Authority: RCW 18.71.450, 18.71A.100, 18.71.017, and 18.71A.020. WSR 11-12-025, § 246-919-862, filed 5/24/11, effective 1/2/12.]

WAC 246-919-863 Pain management specialist. A pain management specialist shall meet one or more of the following qualifications:

- (1) If a physician or osteopathic physician:
- (a) Board certified or board eligible by an American Board of Medical Specialties-approved board (ABMS) or by the American Osteopathic Association (AOA) in physical medicine and rehabilitation, rehabilitation medicine, neurology, rheumatology, or anesthesiology; or
- (b) Has a subspecialty certificate in pain medicine by an ABMS-approved board; or
- (c) Has a certification of added qualification in pain management by the AOA; or

- (d) A minimum of three years of clinical experience in a chronic pain management care setting; and
- (i) Credentialed in pain management by an entity approved by the Washington state medical quality assurance commission for physicians or the Washington state board of osteopathic medicine and surgery for osteopathic physicians; and
- (ii) Successful completion of a minimum of at least eighteen continuing education hours in pain management during the past two years for physicians or three years for osteopathic physicians; and
- (iii) At least thirty percent of the physician's or osteopathic physician's current practice is the direct provision of pain management care or is in a multidisciplinary pain clinic.
- (2) If a dentist: Board certified or board eligible in oral medicine or orofacial pain by the American Board of Oral Medicine or the American Board of Orofacial Pain.
 - (3) If an advanced registered nurse practitioner (ARNP):
- (a) A minimum of three years of clinical experience in a chronic pain management care setting;
- (b) Credentialed in pain management by the Washington state nursing care quality assurance commission-approved national professional association, pain association, or other credentialing entity;

- (c) Successful completion of a minimum of at least eighteen continuing education hours in pain management during the past two years;
- (d) At least thirty percent of the ARNP's current practice is the direct provision of pain management care or is in a multidisciplinary pain clinic.
 - (4) If a podiatric physician:
- (a) Board certified or board eligible in a specialty that includes a focus on pain management by the American Board of Podiatric Surgery, the American Board of Podiatric Orthopedics and Primary Podiatric Medicine, or other accredited certifying board as approved by the Washington state podiatric medical board; or
- (b) A minimum of three years of clinical experience in a chronic pain management care setting; and
- (c) Credentialed in pain management by the Washington state podiatric medical board-approved national professional association, pain association, or other credentialing entity; and
- (d) Successful completion of a minimum of at least eighteen hours of continuing education in pain management during the past two years, and at least thirty percent of the podiatric physician's current practice is the direct provision of pain management care.

[Statutory Authority: RCW 18.71.450, 18.71A.100, 18.71.017, and 18.71A.020. WSR 11-12-025, § 246-919-863, filed 5/24/11, effective 1/2/12.]

Commented [UD(33]: All of the Pain Management sections are under review under a different CR-101.

PHYSICIAN AND SURGEON FEES

WAC 246-919-990 Physician and surgeon fees and renewal cycle. (1)

A physician Licenses must be renewed his or her license every two years on the practitioner'sphysician's birthday as provided in chapter 246-12 WAC, Part 2, except postgraduate training limited licenses.

- (2) Postgraduate training limited licenses must be renewed every year to correspond to the program's date.
- (3) A <u>physician with a retired active physicianlicense</u> who resides and practices in Washington and obtains or renews a retired active license is exempt from all licensing fees except for the impaired physician program surcharge authorized by RCW 18.71.310.
- (4) The aApplicants and licensedesphysicians must pay the following nonrefundable fees:

Title of Fee	Fee
Physicians and surgeons: Chapter 18.71 RCW	
Application (annual)*	\$491.00
Two-year renewal*	657.00
Late renewal penalty	262.50
Expired license reissuance	262.50
Certification of license	50.00
Duplicate license	15.00
Temporary permit	50.00
Application fee for transitioning from a postgraduate training limited license (annual)*	166.00

Title of Fee	Fee
Retired active physicians and surgeons: (Two-year cycle)	
Retired active physician who resides and practices in-state per RCW 18.71.080 and 18.130.250 (Washington physician health program surcharge)	100.00
Retired active physician license renewal *(does not meet in-state exemption)	332.00
Retired active late renewal penalty	50.00
Postgraduate limited license fees: RCW 18.71.095 (One-year cycle)	
Limited license application*	391.00
Limited license renewal*	391.00
Limited duplicate license	15.00

^{*} The application or renewal fee includes: The Washington physician health program surcharge (RCW 18.71.310(2)) assessed at \$50.00 per year, and the University of Washington (UW) HEAL-WA web portal access fee (RCW 43.70.110) assessed at \$16.00 per year.

[Statutory Authority: RCW 43.70.110 (3) (c) and 43.70.250. WSR 12-19-088, \$ 246-919-990, filed 9/18/12, effective 11/1/12. Statutory Authority: RCW 43.70.250, 43.70.280, 18.31.310, 18.71A.020, 18.71.080, and 43.70.110. WSR 09-16-120, \$ 246-919-990, filed 8/4/09, effective 8/15/09. Statutory Authority: RCW 43.70.110, 43.70.250, 2008 c 329. WSR 08-15-014, \$ 246-919-990, filed 7/7/08, effective 7/7/08. Statutory Authority: RCW 43.70.250. WSR 06-11-167, \$ 246-919-990, filed 5/24/06, effective 7/1/06. Statutory Authority: RCW 43.70.250, [43.70.]280 and 43.70.110. WSR 05-12-012, \$ 246-919-990, filed 5/20/05, effective 7/1/05. Statutory Authority: RCW 18.71.017, 18.71A.020 and 43.70.280. WSR 02-05-009, \$ 246-919-990, filed 2/8/02, effective 3/11/02. Statutory Authority: RCW 18.71.017, 18.130.040(4), 18.130.050(12) and 18.130.340. WSR 99-23-090, \$ 246-919-990, filed 11/16/99, effective

1/1/00. Statutory Authority: RCW 43.70.280. WSR 98-05-060, § 246-919-990, filed 2/13/98, effective 3/16/98. Statutory Authority: RCW 18.71.017 and 43.70.250. WSR 97-15-100, \$ 246-919-990, filed 7/21/97, effective 8/21/97. Statutory Authority: RCW 18.71.017 and 18.71A.020. WSR 96-03-073, § 246-919-990, filed 1/17/96, effective 2/17/96.]