

**BOARD OF DIRECT ENTRY MIDWIFERY**  
**PROPOSED ADMINISTRATIVE RULES – APRIL 1, 2026**

**332-015-0030**

**Application Requirements Direct Entry Midwifery License**

(1) An individual applying for licensure to practice direct entry midwifery must:

- (a) Meet the requirements of OAR 331 Division 30;
- (b) Submit a completed application on a form prescribed by the Office, which must contain the information listed in OAR 331-030-0000 and be accompanied by the required application and license fees;
- (c) Submit proof of being currently certified in cardiopulmonary resuscitation for infants and adults pursuant to ORS 687.420;
- (d) Submit proof of being currently certified in neonatal resuscitation.
- (e) Submit a written plan for emergency transport for birthing person or newborn pursuant to OAR 332-025-0020; and
- (f) Submit proof of having current Certified Professional Midwife credential from NARM;
- (g) Submit proof of having passed all four parts of the Legend Drugs and Devices examination, as described in OAR 332-015-0050(3)(a) and (b), within one year prior to the date of application on a form prescribed by the Office. The form must be submitted directly to the Office by the proctor who conducted and supervised the skills examination(s) and who has met all requirements listed in OAR 332-015-0075 and has been approved by the Office.**

(2) In addition to the requirements listed in subsection (1) of this rule and pursuant to ORS 687.420, an applicant must show proof of having participated in:

(a) 25 deliveries as an assistant.

(b) 25 deliveries as the primary or the primary under supervision.

(c) 100 prenatal care visits, 25 newborn examinations, and 40 postnatal examinations.

(d) 10 deliveries where continuity of care was provided as an assistant or primary care provider. The continuity of care must include four prenatal visits, one newborn examination, and one postpartum exam for each of the 10 deliveries.

(3) Of the 50 births listed in subsection (2) of this rule, at least 25 deliveries must have taken place as a community birth and 10 births must have occurred within the two years before the date of application.

(4) For the purpose of this rule, experience as an assistant, primary or primary under supervision, must have been obtained in one of the following ways:

(a) As an Oregon licensed health care practitioner while the services provided were within the scope of the practitioner's license;

(b) As a traditional midwife providing services in Oregon pursuant to ORS 687.415(2)(b);

(c) As an individual supervised by an LDM; ~~or~~ **Or**

(d) Under other lawful means.

**(5) If an applicant was issued a license without first taking the 40 hours Initial Legend Drugs and Devices Continuing Education course, required prior to July 1, 2026, they must pass the Legend Drugs and Devices examination as listed in OAR 332-015-0050 at the time of next**

**renewal and attest to having passed the examinations on a form prescribed by the Office.**

**(6) An applicant must maintain a record of documents required within this rule which may be subject to an audit or inspection by the Office. Documents may be stored electronically.**

**(7) The skills examination(s) listed under (1)(g) of this rule must be sent to the Office through mail, fax or email.**

~~(6) (8) If the applicant has not received the Initial Legend Drugs and Devices continuing education listed under OAR 332-020-0010(2) or (3) at the time of application this information must be disclosed to each patient on the patient disclosure form required under OAR 332-025-0020.~~

~~(5) If the applicant received the Initial Legend Drugs and Devices continuing education prior to applying for licensure, it must have been obtained within two years before the date of application. If applicant receives the continuing education within 12 months after applying for licensure, the applicant must attest to having received the continuing education at the time of next renewal on a form prescribed by the Office.~~

~~(6) If the applicant has not received the Initial Legend Drugs and Devices continuing education listed under OAR 332-020-0010(2) or (3) at the time of application this information must be disclosed to each patient on the patient disclosure form required under OAR 332-025-0020.~~

**332-015-0040**

**Education**

**(1) An Applicant's applicant's** education must incorporate the general educational requirements listed in the NARM CPM candidate information bulletin, including:

~~(1)~~ **(a)** Core competencies developed by MANA;

~~(2)~~ **(b)** NARM written test specifications;

~~(3)~~ **(c)** NARM skills assessment test specifications;

~~(4)~~ **(d)** NARM written examination primary reference list; and

~~(5)~~ **(e)** NARM skills assessment reference list.

**(2) The Board has adopted the July 1, 2026, Legend Drugs and Devices Standard Competencies as the theory and practical instructional outline for the legend drugs and devices.**

**(3) The July 1, 2026, Legend Drugs and Devices Standard Competencies can be obtained from the Office upon request.**

**332-015-0050**

**NARM Midwifery Examinations for Licensure and Retake Examination Requirements**

- (1) Pursuant to ORS 687.420 the Board may establish standard qualifications for licensure including but not limited to requiring passage of a Board approved examination.**
- (2) The Board has determined that an applicant for a license or renewal of a license must pass the Legend Drugs and Devices examination to purchase and administer authorized scheduled legend drugs and devices used in pregnancy, birth, postpartum care, newborn care, and resuscitation.**
- (3) The Legend Drugs and Devices examination consists of the following four parts:**
  - (a) The written examination which consists of multiple-choice questions related to pharmacology, administration of medications through injection, intravenous therapy, treatment of shock and suturing.**
  - (b) The skills examinations consist of three parts that assess the applicant's skills on how to safely administer authorized scheduled legend drugs. The three skills examinations are:**
    - (A) Administration of medication through injection.**
    - (B) Administration of intravenous fluids.**
    - (C) Suturing.**
- (4) The skills examinations listed in subsection (3)(b) of this rule must be supervised and conducted by a proctor who has met all requirements listed in OAR 332-015-0075 and has been approved by the Office.**

**(5) The skills examinations listed in subsection (3)(b) of this rule may be conducted and supervised by different proctors. Each proctor must meet all requirements listed in OAR 332-015-0075 and be approved by the Office prior to conducting a skills examination.**

**(6) The skills examinations listed in subsection (3)(b) of this rule are valid for three years and must be kept current.**

**(7) An applicant is responsible for payment of all examination fees including but not limited to fees paid directly to an approved proctor.**

**(8) An applicant who does not pass the written examination listed in subsection (3)(a) of this rule may retake the written examination the next business day.**

**(9) An applicant who does not pass a skills examination listed in subsection (3)(b) of this rule may retake that portion of the skills examination the next business day up to three times.**

**(10) An applicant who has failed any portion of the skills examination listed in subsection (3)(b) of this rule three times must wait six months before retaking that skills examination for the fourth time.**

~~The qualifying examination is the NARM examination. An applicant is responsible for payment of all fees for NARM applications, examinations, and any other fees paid directly to NARM.~~

**332-015-0075**

**Legend Drugs and Devices Skills Examination: Proctor Application and Requirements**

**(1) To be an approved proctor for a skills examination portion of the examination listed in OAR 332-015-0050 an individual must:**

**(a) Submit a completed application on a form prescribed by the Office, which must contain the information listed in OAR 331-030-0000.**

**(b) Submit proof of holding a qualifying credential or authorization listed in subsection (2), (3) or (4) of this rule that is active with no current disciplinary action imposed by the Board or other regulatory body.**

**(2) To be approved to proctor the administration of medication through injection skills examination the applicant must be one of the following:**

**(a) Oregon Licensed Direct Entry Midwife.**

**(b) NARM Certified Professional Midwife.**

**(c) Oregon Registered Nurse.**

**(d) Oregon Certified Nurse Midwife.**

**(e) Oregon Licensed Paramedic.**

**(f) Oregon Licensed Advanced Emergency Medical Technician.**

**(g) Oregon Licensed Medical Doctor.**

**(h) Oregon Licensed Doctor of Osteopathic Medicine. Or**

**(i) Oregon Licensed Doctor of Naturopathy.**

**(3) To be approved to proctor the intravenous therapy skills examination the applicant must be one of the following:**

**(a) Oregon Licensed Direct Entry Midwife.**

**(b) NARM Certified Professional Midwife.**

- (c) Oregon Registered Nurse.**
  - (d) Oregon Certified Nurse Midwife.**
  - (e) Oregon Licensed Paramedic.**
  - (f) Oregon Licensed Advanced Emergency Medical Technician.**
  - (g) Oregon Licensed Medical Doctor.**
  - (h) Oregon Licensed Doctor of Osteopathic Medicine. Or**
  - (i) Oregon Licensed Doctor of Naturopathy.**
- (4) To be approved to proctor the suturing skills examination the applicant must be one of the following:**
- (a) Oregon Licensed Direct Entry Midwife.**
  - (b) NARM Certified Professional Midwife.**
  - (c) Oregon Certified Nurse Midwife.**
  - (d) Oregon Licensed Medical Doctor.**
  - (e) Oregon Licensed Doctor of Osteopathic Medicine. Or**
  - (f) Oregon Licensed Doctor of Naturopathy.**
- (5) Prior to proctoring a skills examination listed in subsection (2), (3) or (4) of this rule, written approval must be received by the Office. Written approval may be obtained by the Office electronically.**
- (6) A person applying to be an approved proctor who does not meet the requisite requirements listed in subsection (2), (3) or (4) of this rule may submit satisfactory documentation to the office that demonstrates the applicant has substantially equivalent education and training in legend drugs and devices to those individuals listed in the requisite subsection (2), (3), or (4) of this rule, this may include training received outside of Oregon.**

**(7) A skills examination listed in subsection (2), (3) or (4) of this rule must be performed under direct supervision of an approved proctor. Supervision may be in person or virtual in real time.**

**(8) An approved proctor must:**

**(a) Supervise only one examinee, at a time.**

**(b) Exercise management, guidance, and control over the portion of the examination being administered and exercise professional judgment during the examination.**

**(c) Exercise professional judgment.**

**(d) Follow the skills examination(s) guidelines prescribed by the Office, verifying only the steps performed by the examinee.**

**(e) Document the skills performed under OAR 332-015-0050(3)(b) by the examinee, on a form prescribed by the Office.**

**(f) Maintain the skills examination documentation for a minimum of five (5) years following completion of the skills examination.**

**(g) Documentation listed under subsection (8)(f) of this rule may be stored electronically.**

**(h) Respond to all requests for information from the Office related to the skills examination(s).**

**(9) An approved proctor is responsible for all matters relative to the examinee during the skills examination.**

**(10) An approved proctor may also provide legend drugs and devices education listed under OAR 332-015-0040.**

**(11) An approved proctor must submit proof to the Office each time an applicant or licensee has passed a skills examination listed in OAR 332-015-0050. The proof must be submitted on a form prescribed by the Office.**

**332-015-0077**

**Qualification and Requirements for Legend Drugs and Devices**

**Written Examination**

**(1) Before an individual can take the written portion of the examination listed in OAR 332-015-0050 the individual must submit a form prescribed by the Office, which must contain the information listed in OAR 331-030-0000, and be accompanied by payment of all required fees.**

**(2) An individual must receive approval from the Office to take the written portion of the examination listed in OAR 332-015-0050. All documentation must be received and verified prior to taking the written portion of the examination.**

PROPOSED RULES

## 332-020-0000

### License Issuance and Renewal

(1) LICENSING: An LDM is subject to the provisions of OAR Chapter 331, Division 30 regarding the issuance and renewal of a license, and provisions regarding authorization to practice, identification, and requirements for issuance of a duplicate license.

(2) LICENSE RENEWAL: To avoid delinquency penalties, license renewal must be made prior to the license entering inactive status. To renew, an LDM must:

- (a) Submit a renewal application form.;
- (b) Submit payment of renewal fee.;
- (c) Attest to having obtained required continuing education under OAR 332-020-0010, on a form prescribed by the Office.;
- (d) Attest to being currently certified in cardiopulmonary resuscitation for infants and adults pursuant to ORS 687.425.;
- (e) Attest to being currently certified in neonatal resuscitation.;
- (f) Attest to having participated in peer review pursuant to 332-025-0015.
- (g) Attest to having passed the Legend Drugs and Devices examination as listed in OAR 332-015-0050 within the last three years.**

(3) INACTIVE LICENSE RENEWAL: A license may be inactive for up to three years. When renewing after entering inactive status the LDM must:

- (a) Submit a renewal application form.;
- (b) Submit payment of renewal and delinquency fees.;
- (c) Attest to having obtained required continuing education under OAR 332-020-0010, on a form prescribed by the Office, whether license is current or inactive.;

(d) Attest to being currently certified in cardiopulmonary resuscitation for infants and adults pursuant to ORS 687.425;

(e) Attest to being currently certified in neonatal resuscitation; and

(f) Attest to having participated in peer review pursuant to 332-025-0015.

**(g) Attest to having passed the Legend Drugs and Devices examination as listed in OAR 332-015-0050 within the last three years.**

(4) EXPIRED LICENSE: A license that has been inactive for more than three years is expired and the licensee must reapply and meet the requirements listed in OAR 332-015-0030.

**(5) An LDM must maintain documentation and records required for renewal of a license for at least five years. This information may be stored electronically and may be subject to an audit or inspection by the Office.**

PROPOSED RULES

## **332-020-0010**

### **Continuing Education**

(1) Standard Continuing Education Renewal Requirements: To maintain licensure, an LDM must complete at least ~~35~~ **18** continuing education hours every ~~year~~ ~~two years~~ from the date of initial licensure, ~~and every two years thereafter~~. Continuing education under this rule must be related to the services listed in ORS 687.405, cultural competency, patient charting, ethics, communication, or professional development.

(2) In addition to the requirements listed in section (1) of this rule, ~~five~~ **one** of the ~~35~~ **18** hours listed in subsection (1) of this rule must be related to legend drugs and devices and **pharmacology, covering drugs listed in ORS 687.493, OAR 332-026-0010 and OAR 332-026-0020**. ~~pursuant to OAR Chapter 332, Division 26, which must include at least one (1) of the following:~~

~~(a) Pharmacology covering drugs listed in ORS 687.493, OAR 332-026-0010 and 332-026-0020.~~

~~(b) Administration of medications through injection.~~

~~(c) Advanced treatment of shock.~~

~~(d) Intravenous therapy.~~

~~(e) Suturing.~~

~~(3) Initial Legend Drugs and Devices Continuing Education: An LDM must complete 40 hours of instruction in an approved curriculum prior to purchasing or administering legend drugs and devices listed in OAR Chapter 332, Division 26 of these rules or by the date of first renewal following initial licensing as an LDM. The initial continuing education is comprised of theory, hands-on practice, and skills testing for competency which must include:~~

~~(a) 10 hours in pharmacology, covering drugs listed in ORS 687.493, OAR Chapter 332, Division 26 including;~~

~~(b) Four hours of administration of medications through injection;~~

~~(c) Four hours in advanced treatment of shock;~~

~~(d) 12 hours in intravenous therapy; and~~

~~(e) 10 hours in suturing.~~

~~(4) In addition to the requirements listed in subsection (1) of this rule and in accordance with ORS 687.425, an LDM who has attended fewer than five births in the previous renewal year must obtain an additional 10 hours of continuing education separate from all other continuing education requirements. The additional 10 hours of continuing education must be obtained during the next license renewal cycle. The additional 10 hours of continuing education must be related to subjects listed in subsection (1) of this rule. An example pertaining to the timing of when the additional 10 hours of continuing education must be obtained is: An LDM who held a license from April 2018 to April 2019 who attended fewer than five births during that time period must obtain an additional 10 continuing education hours between April 2019 and April 2020.~~

**(3) In accordance with ORS 687.425, an LDM who attended fewer than five births during the previous renewal year must complete an additional 10 hours of continuing education during the next renewal cycle.**

**(a) The additional 10 hours must be separate from all other continuing education requirements.**

**(b) The additional hours must cover subjects listed in subsection (1) of this rule.**

**(c) An example pertaining to the timing of when the additional 10 hours of continuing education must be obtained is: An LDM who held a license from April 2024 to April 2025 who attended fewer than five births during that time period must obtain an additional 10 continuing education hours between April 2025 and April 2026.**

~~(5)~~ **(4)** Continuing education may be obtained through online courses, attendance at lectures, sessions, courses, workshops, symposiums seminars or other presentations offered by:

(a) Institutions or programs accredited by a federally recognized accrediting body;

(b) Institutions or programs approved by the Oregon Higher Education Coordinating Commission;

(c) An organization offering continuing medical education opportunities, including legend drugs and devices training, including but not limited to, Accreditation Council for Continuing Medical Education, MEAC accredited or pre-accredited schools and the Oregon Midwifery Council; ~~or~~ **Or**

(d) Any Board-approved professional organization, or association, hospital, or health care clinic offering continuing education related to the subject matter listed in subsection (1) ~~or (7)~~ of this rule.

~~(6)~~ **(5)** Continuing education relating to the subject matter listed in subsection (1) of this rule may also be obtained through research, authorship or teaching, provided that no more than half the required hours be in research, authorship or teaching.

~~(7)~~ **(6)** Up to ~~10~~ **(5)** hours of continuing education relating to subject matter listed in subsection (1) of this rule may be completed through self-study that may include clinical service learning. Documentation substantiating the

completion of continuing education through self-study must be **maintained by the licensee.** ~~submitted on forms provided by the Office.~~

~~(8)~~ **(7)** Notwithstanding any other continuing education requirements and of the ~~35~~ **18** continuing education hours listed under subsection (1) of this rule, in order to be eligible to renew a license, an authorization holder must obtain one hour of continuing education in cultural competency (CCCE) every other time the authorization is subject to renewal. The requirement of this paragraph is effective on and after July 1, 2021. The CCCE must be either:

(a) Approved by the Oregon Health Authority; or

(b) Meet the skills requirements established by the Oregon Health Authority by rule; or

(c) One of the educational institutions or organizations listed in subsection (4) of this rule which are in compliance with one of the requirements listed in subsection ~~(8)~~ **(7)** of this rule.

~~(9)~~ **(8)** The Board encourages completion of required CCCE from the approved list promulgated by the Oregon Health Authority, available on the Board's website. The Board shall also accept CCCE that is not obtained from the Oregon Health Authority's approved list, to the extent the Board determines the CCCE meets the skills requirements established by the Oregon Health authority in rules that are effective at the time the CCCE is obtained.

~~(10)~~ **(9)** ~~Section (8) does not apply to authorization holders who are:~~

**Requirements listed in subsection (8) of this rule do not apply to the authorization holders who are:**

(a) Retired and not practicing the profession in any state; or

(b) Not practicing the profession in this state; or

(c) Residing in this state but not practicing in the profession in any state.

~~(11)~~ **(10)** Obtaining and maintaining proof of participation in continuing education is the responsibility of the licensee. The licensee must ensure that adequate proof of attainment of required continuing education is available for audit or investigation or when otherwise requested by the Office. Adequate proof of participation is listed under OAR 332-020-0015~~(3)~~.

~~(12)~~ **(11)** Hours of continuing education that are obtained in excess of the minimum requirements listed in this rule will not be carried forward.

~~(13)~~ **(12)** Documentation of participation in continuing education requirements must be maintained for five years following renewal and must be available to the Office upon request.

~~(14)~~ **(13)** For the purpose of this rule, continuing education must include periods of continuous instruction and education. One hour of continuing education is one hour of actual class time, not to include breaks, rest periods, travel, registration or meals.

~~(15)~~ A copy of Board-approved curriculum objectives January 2021 Legend Drugs and Devices Program is available at the Health Licensing Office or on the Office website. Payment of administrative fees may be required. Refer to OAR 331-010-0030 for applicable public record request fees.

~~(16)~~ Requirements listed under OAR 332-020-0000 cannot be used towards the continuing education requirements listed in this rule.

### **332-020-0015**

#### **Continuing Education: Audit and Required Documentation**

- (1) The Office will audit a select percentage of licensees determined by the Board to verify compliance with continuing education requirements.
- (2) An LDM notified of selection for audit of continuing education attestation must submit to the Office, within 30 calendar days from the date of notification, satisfactory evidence of participation in required continuing education in accordance with OAR 332-020-0010.
- (3) If selected for audit, an LDM must provide documentation of the required continuing education. Documentation must include:
  - (a) Certificate of completion, official transcript, statement or affidavit from the sponsor attesting to attendance or other documentation approved by the Office;
  - (b) Name of sponsoring institution, association, or organization;
  - (c) Title of presentation and description of content;
  - (d) Name of instructor or presenter;
  - (e) Date of attendance and duration in hours; ~~and~~ **And**
  - (f) Course agenda.
- (4) If documentation of continuing education is incomplete, the licensee has 30 calendar days from the date of the notice sent to the licensee to submit further documentation to substantiate having completed the required continuing education.
- (5) An LDM notified of having been selected for an audit must show proof of having:
  - (a) Participated in peer review pursuant to OAR 332-025-0015;
  - (b) The required certification in cardiopulmonary resuscitation for infants and adults; ~~and~~ **And**

(c) The required certification in neonatal resuscitation-; **and**

**(d) Having passed the Legend Drugs and Devices examination as listed in OAR 332-015-0050 every three years.**

(6) Documentation listed in subsection (5) of this rule must be kept and maintained for five years following renewal and must be available to the Office upon request.

PROPOSED RULES

## **332-026-0000**

### **Access to and Administration of Legend Drugs and Devices**

~~(1) An LDM is prohibited from purchasing or administering legend drugs and devices, until the continuing education listed in OAR 332-020-0010 has been completed and attestation submitted to the Office upon renewal.~~

~~(2) Pursuant to ORS 687.493, an LDM who completes the continuing education listed in OAR 332-020-0010 is authorized to purchase and administer legend drugs and devices listed in OAR 332-026-0010, 332-026-0020, and 332-026-0030.~~

~~(3) (1) An LDM must comply with all local, state and federal laws and regulations regarding the administration, distribution, storage, transportation and disposal of legend drugs and devices listed in OAR 332-026-0010, 332-026-0020, 332-026-0030.~~

~~(4) (2) Approved legend drugs must be inventoried and securely stored when the product is not in use, including samples or any remaining portion of a drug.~~

~~(5) (3) Records regarding approved legend drugs and devices must be maintained for five years. Records must be kept on the business premises and available for inspection upon request by the Office. Upon request by the Board or Office, an LDM must provide a copy of records. Records must include, but are not limited, to the following:~~

~~(a) Name of drug, amount received, date of receipt, and drug expiration date;.~~

~~(b) Name of drug and to whom it was administered; date and amount of drug administered to client;.~~

~~(c) Name of drug, date and place or means of disposal.~~

~~(6)~~ **(4)** Expired, deteriorated or unused legend drugs must be disposed of in a manner that protects the LDM, client and others who may come into contact with the material during disposal.

~~(7)~~ An LDM is required to obtain the continuing education for intravenous antibiotics for Group B Streptococcal prophylaxis, however a LDM is not required to administer the antibiotic.

PROPOSED RULES

## **332-026-0010**

### **Approved Legend Drugs For Maternal Use**

An LDM may administer the following legend drugs as approved by the Board for maternal use:

(1) Anti-Hemorrhagics for use by intramuscular injection ~~includes~~ **is limited to:**

- (a) Synthetic Oxytocin (Pitocin, Syntocin and generic);
- (b) Methylergonovine (Methergine);
- (c) Ergonovine (Ergotrate); or

(2) Anti-Hemorrhagics by intravenous infusion is limited to:

- (a) Synthetic Oxytocin (Pitocin, Syntocin, and generic); or
- (b) Tranexamic acid.

(3) Anti-Hemorrhagics for oral administration is limited to:

- (a) Methylergonovine (Methergine);
- (b) Misoprostol (Cytotec).

(4) Anti-Hemorrhagics for rectal administration is limited to Misoprostol (Cytotec).

(5) Resuscitation is limited to medical oxygen and intravenous fluid replacement.

(6) Intravenous fluid replacement ~~includes~~ **is limited to:**

- (a) Lactated Ringers Solution;
- (b) 0.9% Saline Solution;
- (c) D5LR (5% Dextrose in Lactated Ringers); or
- (d) D5W (5% Dextrose in water).

(7) Anaphylactic treatment by subcutaneous injection is limited to Epinephrine.

(8) Local anesthetic includes:

(a) Lidocaine HCl (1% and 2%) (Xylocaine and generic);

(b) Topical anesthetic;

(c) Procaine HCl (Novocain, benzocaine, cetacane and generic);

(d) Sterile water papules.

(9) Rhesus Sensitivity Prophylaxis is limited to Rho(d) Immune Globulin (RhoGAM, Gamulin Rh, Bay Rho-D and others).

(10) Tissue adhesive (Dermabond or generic).

(11) Intravenous antibiotics for Group B Streptococcal prophylaxis is limited to the following and is only to be used solely for the purpose of Group B Streptococcal prophylaxis:

(a) Penicillin;

(b) Ampicillin;

(c) Cefazolin; or

(d) Clindamycin.

PROPOSED RULES

### **332-026-0020**

#### **Approved Legend Drugs For Neonatal Use**

An LDM may administer the following legend drugs as approved by the Board for neonatal use:

- (1) Eye Prophylaxis for disease of the newborn is limited to Erythromycin Ophthalmic (0.5%) Ointment (Ilotycin, AK-Mycin and generics).
- (2) Prophylaxis for hemorrhagic disease of the newborn for oral use is limited to Mephyton.
- (3) Prophylaxis for hemorrhagic disease of the newborn for intramuscular injection includes:
  - (a) AquaMephyton; and
  - (b) Konakion.
- (4) Resuscitation is limited to medical oxygen.

### **332-026-0030**

#### **Approved Devices**

An LDM may use the following Board approved devices:

- (1) Devices for injection of medications including:
  - (a) Needles.
  - (b) Syringes.
- (2) Devices for administration of intravenous fluids including:
  - (a) Drip sets.
  - (b) Catheters.
- (3) Devices for maternal and neonatal resuscitation including:
  - (a) Suction devices;
  - (b) Oxygen-delivery devices.
  - (c) Bag-Valve-Mask-Sets.
  - (d) Airway adjunct.**
- (4) Devices for rupturing the amniotic sac.
- (5) Devices for repairing the perineal area including:
  - (a) Sutures;
  - (b) Instruments for completing a repair.
  - (c) Local anesthetic administration devices.

## 332-040-0000

### Fees

(1) An applicant and licensee are subject to the provisions of OAR 331-010-0010 and 331-010-0020 regarding payment of fees, penalties and charges.

(2) Fees established by the Office pursuant to ORS 676.576 are as follows:

(a) Application:

(A) License: \$150.

(B) License by reciprocity: \$750.

(b) **Written portion of Examination — Oregon laws & rules Legend Drugs and Devices:** \$50.

(c) Original issuance of license: \$800 for one year.

(d) Renewal — License: \$800 for one year.

~~(e) Reactivation of license: \$150.~~

(f) Other administrative fees:

(A) Delinquency fee: \$50 for each year the license is inactive for up to three years.

(B) Replacement of license, including name change: \$25.

(C) Duplicate license document: \$25. ~~per copy, with a maximum of three.~~

(D) Affidavit of licensure **or credential** for reciprocity: \$50.

(E) An additional \$25 administrative processing fee will be assessed if a non-sufficient funds or non-negotiable instrument is received for payment of fees, penalties and charges. Refer to OAR 331-010-0010.

~~(3) As of April 1, 2020, an applicant applying for an original license totaling \$800 may be granted a \$350 license fee discount for a total cost for the license \$450 until July 1, 2024. An application fee of \$150 must be paid in order to grant the \$350 license fee discount. The license fee discount is~~

~~available to individuals who meet all application requirements for direct entry midwifery licensure under OAR 332-015-0030 and reside in Oregon.~~

~~(4) As of April 1, 2020, an applicant applying to renew a license totaling \$800 may be granted a \$200 discount for a total cost for the license \$600 until July 1, 2024. The license fee discount is available to individuals who meet all renewal requirements for direct entry midwifery licensure under OAR 332-020-0000 and reside in Oregon.~~

PROPOSED RULES