IV. Proposed Regulations to Add Title 16 California Code of Regulations (CCR) section 1746.5, Related to Travel Medications

At the June 2015 Board Meeting, the board approved proposed text to add Section 1746.5 of Title 16 CCR, related to Travel Medications. The 45 day comment period began on September 25, 2015 and ended November 9, 2015. At the January 19, 2016 Board meeting, in response to the comments received during the 45 day comment period, the board approved a modified text and initiated a 15 day comment period. The 15 day comment period began January 27, 2016 and ended February 11, 2016.

One comment was received during the 15 day comment period.

During the February 2016 Board meeting, the Board voted to adopt the language as noticed on January 27, 2016. The recommendation to amend the protocol will be discussed again at the Board meeting.

At this Meeting
The board will have the opportunity to discuss the regulation, the comment received and determine what course of action it wishes to pursue. Among its options:
   1. Adopt the regulation as approved at the February 2016 Board meeting
   2. Amend the regulation to address the concern expressed by the stakeholder and notice the modified text for a second 15 day comment period.

The Attachment immediately following this memo contains the proposed regulation text as adopted at the February 2016 Board meeting, a copy of the comment received during the 15 day comment period, and proposed modified text, as recommended by staff.
Travel Medications -
1746.5
Travel Medications – Adopted Text
February 2016
Add §1746.5 to Article 5 of Division 17 of Title 16 of the California Code of Regulations as follows:

§1746.5 Pharmacists Furnishing Travel Medications.

(a) For purposes of Business and Professions Code section 4052(a)(10)(A)(3), prescription medications “not requiring a diagnosis” means a prescription medication that is either:

    (1) For a condition that is both self-diagnosable and recognized as self-treatable by the federal Center for Disease Control and Prevention’s (CDC) Health Information for International Travel (commonly called the Yellow Book), or

    (2) A prophylactic.

(b) A pharmacist furnishing prescription medications not requiring a diagnosis that are recommended by the CDC for individuals traveling outside the 50 states and the District of Columbia pursuant to section 4052(a)(10) of the Business and Professions Code shall follow the requirements of this section.

(c) Training: A pharmacist who furnishes travel medications shall keep documentation of the following on site and available for inspection by the Board:

    (1) Completion of an immunization certification program that meets the requirements of Business and Professions Code section 4052.8(b)(1).

    (2) Completion of an approved travel medicine training program, which must consist of at least 10–20 hours of training and cover each medication related element of the International Society of Travel Medicine’s Body of Knowledge for the Practice of Travel Medicine (2012),

    (3) Completion of the CDC Yellow Fever Vaccine Course, and

    (4) Current basic life support certification.
(d) Continuing Education: Pharmacists must complete two hours of ongoing continuing education focused on travel medicine, separate from continuing education in immunizations and vaccines, from an approved provider once every two years.

(e) Prior to furnishing travel medication, a pharmacist shall perform a good faith evaluation of the patient, including evaluation of a patient travel history using destination-specific travel criteria. The travel history must include all the information necessary for a risk assessment during pre-travel consultation, as identified in the CDC Yellow Book. An example of an appropriate and comprehensive travel history is available on the Board’s website.

(f) Notifications: The pharmacist shall notify the patient’s primary care provider of any drugs and/or devices furnished to the patient within 30 days of the date of dispensing, or enter the appropriate information in a patient record system shared with the primary care provider, as permitted by the primary care provider. If the patient does not have a primary care provider, or is unable to provide contact information for his or her primary care provider, the pharmacist shall provide the patient with written record of the drugs and/or devices furnished and advise the patient to consult a physician of the patient’s choice.

(g) Documentation: For each travel medication furnished by a pharmacist, a patient medication record shall be maintained and securely stored in physical or electronic manner such that the information required by title 42, section 300aa-25 of the United States Code and title 16, sections 1707.1 and 1717 of the California Code of Regulations is readily retrievable during the pharmacy or facility’s normal operating hours. A pharmacist shall provide the patient with a progress note, which fully documents the clinical assessment and travel medication plan. An example of an appropriate and comprehensive progress note is available on the Board’s website.

Note: Authority cited: Section 4005, Business and Professions Code. Reference: Section 4052 and 4052.8, Business and Professions Code.
Travel Medications -
15-Day Comments
Comment Period
Closed February 11, 2016
Hi Lori,

Attached are some minor modifications. I made them directly on the draft, but they are also listed below:

(c)(1) "...an immunization certificate program..." – there are no certification programs for immunization, just certificate

(c)(2) "...of training and cover at least each medication related" – programs that only train on medications are not adequate.

Thanks!
Jeff

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Add §1746.5 to Article 5 of Division 17 of Title 16 of the California Code of Regulations as follows:

§1746.5 Pharmacists Furnishing Travel Medications.

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(1) For a condition that is both self-diagnosable and recognized as self-treatable by the federal Center for Disease Control and Prevention’s (CDC) Health Information for International Travel (commonly called the Yellow Book), or

(2) A prophylactic.

(b) A pharmacist furnishing prescription medications not requiring a diagnosis that are recommended by the CDC for individuals traveling outside the 50 states and the District of Columbia pursuant to section 4052(a)(10) of the Business and Professions Code shall follow the requirements of this section.

(c) Training: A pharmacist who furnishes travel medications shall keep documentation of the following on site and available for inspection by the Board:

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(2) Completion of an approved travel medicine training program, which must consist of at least 10 20-hours of training and cover each medication related element of the International Society of Travel Medicine’s Body of Knowledge for the Practice of Travel Medicine (2012),

(3) Completion of the CDC Yellow Fever Vaccine Course, and

(4) Current basic life support certification.
(d) Continuing Education: Pharmacists must complete two hours of ongoing continuing education focused on travel medicine, separate from continuing education in immunizations and vaccines, from an approved provider once every two years.

(e) Prior to furnishing travel medication, a pharmacist shall perform a good faith evaluation of the patient, including evaluation of a patient travel history using destination-specific travel criteria. The travel history must include all the information necessary for a risk assessment during pre-travel consultation, as identified in the CDC Yellow Book. An example of an appropriate and comprehensive travel history is available on the Board’s website.

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(g) Documentation: For each travel medication furnished by a pharmacist, a patient medication record shall be maintained and securely stored in physical or electronic manner such that the information required by title 42, section 300aa-25 of the United States Code and title 16, sections 1707.1 and 1717 of the California Code of Regulations is readily retrievable during the pharmacy or facility’s normal operating hours. A pharmacist shall provide the patient with a progress note, which fully documents the clinical assessment and travel medication plan. An example of an appropriate and comprehensive progress note is available on the Board’s website.

Note: Authority cited: Section 4005, Business and Professions Code. Reference: Section 4052 and 4052.8, Business and Professions Code.
Travel Medications
Staff Recommended
Modified Text
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