DIVISION OF HEALTH CARE FINANCING AND POLICY NEVADA MEDICAID DRUG USE REVIEW (DUR) BOARD PROPOSED PRIOR AUTHORIZATION CRITERIA

Eliquis[®] (apixaban), Pradaxa[®] (dabigatran) and Xarelto[®] (rivaroxaban) are a covered benefit of Nevada Medicaid for recipients who meet the criteria for coverage.

1. Coverage and Limitations:

Authorization will be given if the following criteria are met and documented:

Requests for Eliquis[®] (apixaban)

- 1. Must have the following:
 - a. The recipient has a diagnosis of nonvalvular atrial fibrillation.

AND

The recipient does not have an active pathological bleed.

The recipient has failed to consistently achieve and maintain an INR of 2.0 to 3.0 on warfarin therapy despite an adequate trial that included patient education and dose adjustments.

OR

The recipient has experienced an adverse event with warfarin or has a contraindication to treatment with warfarin.

Requests for Pradaxa® (dabigatran)

- 1. Must have the following:
 - a. The recipient has a diagnosis of nonvalvular atrial fibrillation.
 - AND

The recipient does not have an active pathological bleed.

AND

The recipient does not have mechanical prosthetic heart valves.

AND

The recipient has failed to consistently achieve and maintain an INR of 2.0 to 3.0 on warfarin therapy despite an adequate trial that included patient education and dose adjustments.

OR

The recipient has experienced an adverse event with warfarin or has a contraindication to treatment with warfarin.

Requests for Xarelto® (rivaroxaban)

- 1. Must have ONE of the following:
 - a. The recipient has a diagnosis of nonvalvular atrial fibrillation.

AND

The recipient does not have an active pathological bleed. **AND**

The recipient has failed to consistently achieve and maintain an INR of 2.0 to 3.0 on warfarin therapy despite an adequate trial that included patient education and dose adjustments.





OR

The recipient has experienced an adverse event with warfarin or has a contraindication to treatment with warfarin.

b. The requested medication will be used for the treatment of deep vein thrombosis (DVT), pulmonary embolism (PE), or for the reduction in the risk of recurrence of DVT and of PE.

AND

The recipient does not have an active pathological bleed.

AND

The recipient has failed to consistently achieve and maintain an INR of 2.0 to 3.0 on warfarin therapy despite an adequate trial that included patient education and dose adjustments.

OR

The recipient has experienced an adverse event with warfarin, or has a contraindication to treatment with warfarin.

2. PA Guidelines:

Prior Authorization approval will be 1 year.

3. Quantity Limitations:

Eliquis[®] (apixaban): 60 tablets per 30 days Pradaxa[®] (dabigatran): 60 capsules per 30 days Xarelto[®] (rivaroxaban): 30 tablets per 30 days



