

Clinical Information - Adstiladrin (nadofaragene firadenovec-vncg)

Is your patient is 18 years or older?

- Yes
 No

Does your patient have a documented diagnosis of Non-Muscle Invasive Bladder Cancer?

- Yes
 No

Does your patient have documentation of Bacillus Calmette-Guerin (BCG) – unresponsive disease?

- Yes
 No

Does your patient have documentation of **ONE** of the following:

- Yes, carcinoma in situ (CIS) with or without high-grade papillary Ta/T1 tumors
 Yes, high-grade papillary Ta/T1 tumors without CIS

Does your patient have documentation Adstiladrin is used for **ONE** of the following:

- Initial treatment
 Recurrent or persistent disease detected by positive cytology or on bladder biopsy despite negative imaging and no visible lesions identified on cytology

Adstiladrin is prescribed by, or in consultation with, a urologist or oncologist?

- Yes
 No

Is this request for new or continued therapy?

- New start
 Continuation of therapy (must provide documentation of beneficial response)

If any of the requirements listed above are not met and provider feels administration of Adstiladrin is medically necessary please provide clinical support and rationale for the use of Adstiladrin.

Additional pertinent information: (including recent history and physical, recent lab work, disease stage, prior therapy, performance status, and names/doses/admin schedule of any agents to be used concurrently).

Additional CPT and Administration Codes for Consideration Following Medical Necessity Determination

Please indicate any other CPT codes that will be billed for administration.

- J9029 Injection, nadofaragene firadenovec-vncg, per therapeutic dose
 Other

Agreement and Attestation

Do you and your patient agree to share any required plan specific outcome measures?

- Yes
 No

Attestation: I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan or insurer its designees may perform a routine audit and request the medical information necessary to verify the accuracy of the

information reported on this form.

Prescriber Signature: _____ **Date:** _____

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