

# Bladder Cancer Adstiladrin (Nadofaragene Firadenovec-vncg) J9029 Prior Authorization Request Medicare Part B Form

Instructions: \* Indicates required information – Form may be returned if required information is not provided. Please fax this request to the appropriate fax number listed at the bottom of the page.

	Date Req	uested	<del></del>					
	Requesto	r	Clinic name: _		Phone		/ Fax	
MEMBER INFORMATION								
*Name:			*I	*ID#:		*DOB:		
PRESCRIBER INFORMATION								
*Name:						*Phone:		
*Address:				*Fax:				· · · · · · · · · · · · · · · · · · ·
DISPENSING PROVIDER / ADMINISTRATION INFORMATION								
*Name: Phone:								
*Add	dress:			Fax:				
PROCEDURE / PRODUCT INFORMATION								
нс	PC Code	Name of Drug	☐ Self-administered	Dose (Wt: _	kg Ht:	)	Frequency	End Date if known
□ Chart notes attached. Other important information:								
Diagnosis: ICD10: Description:								
☐ Provider attests the diagnosis provided is an FDA-Approved indication for this drug								
CLINICAL INFORMATION								
<ul> <li>□ New Start or Initial Request: (Clinical documentation required for all requests)</li> <li>□ Provider has reviewed the attached "Criteria for Approval" and attests the member meets         ALL required PA criteria.     </li> <li>If not, please provide clinical rationale for formulary exception:</li> </ul>								
<ul> <li>□ Continuation Requests: (Clinical documentation required for all requests)</li> <li>□ Provider has reviewed the attached "Criteria for Continuation" and attests the member meets         ALL required PA Continuation criteria.</li> <li>□ Patient had an adequate response or significant improvement while on this medication.         If not, please provide clinical rationale for continuing this medication:</li> </ul>								
ACKNOWLEDGEMENT								
Request By (Signature Required):								



# **Prior Authorization Group - Bladder Cancer PA**

# Drug Name(s):

# **ADSTILADRIN**

### NADOFARAGENE FIRADENOVEC-VNCG

# Criteria for approval of Non-Formulary/Preferred Drug:

- 1. Prescribed for an approved FDA diagnosis (as listed below):
- 2. Drug meets the following utilization management criteria:
  - a. The member has non-muscle invasive bladder cancer; and
  - b. The disease is high-risk; and
  - c. The disease is Bacillus Calmette-Guerin (BCG)-unresponsive.
- 3. Member does not have any clinically relevant contraindications, or CMS/Plan exclusions, to the requested drug.
  - If the member meets all these criteria, they may be approved by the Plan for the requested drug.
  - Quantity limits and Tiering will be determined by the Plan.

### **Exclusion Criteria:**

N/A

## **Prescriber Restrictions:**

N/A

# **Coverage Duration:**

# Approval will be for 6 months

### **FDA Indications:**

• Treatment of adult patients with high-risk <u>Bacillus Calmette-Guerin (BCG)-unresponsive</u> non-muscle invasive bladder cancer (NMIBC) with carcinoma in situ (CIS) with or without papillary tumors

### Off-Label Uses:

N/A

### Age Restrictions:

Safety and effectiveness have not been established in pediatric patients

# Other Clinical Consideration:

N/A

### Resouces:

https://www.micromedexsolutions.com/micromedex2/librarian/CS/30DD1F/ND PR/evidencexpert/ND P/evidencexpert/DUPLICATI ONSHIELDSYNC/6B979B/ND PG/evidencexpert/ND B/evidencexpert/ND AppProduct/evidencexpert/ND T/evidencexpert/PFActionld/evidencexpert.IntermediateToDocumentLink?docId=933877&contentSetId=100&title=Nadofaragene+Firadenovecvncg&servicesTitle=Nadofaragene+Firadenovecvncg#