

Imfinzi[®] (durvalumab) Prior Authorization Form

Member Name: _____ Date of Birth: _____ Member ID#: _____

Drug Information

Physician billing (HCPCS code: _____) Start Date (or date of next dose): _____

Dose: _____ Dosing Regimen: _____

Billing Provider Information

Provider NPI: _____ Provider Name: _____

Provider Phone: _____ Provider Fax: _____

Prescriber Information

Prescriber NPI: _____ Prescriber Name: _____

Prescriber Phone: _____ Prescriber Fax: _____ Specialty: _____

Criteria

For Initial Authorization (Initial approval will be for the duration of 6 months):

1. Please indicate the diagnosis and information:

Non-Small Cell Lung Cancer (NSCLC)

A. Does member have resectable (tumors ≥ 4 cm and/or node positive) NSCLC? Yes No

i. If yes, will durvalumab be used in combination with platinum-containing chemotherapy as neoadjuvant treatment before surgery, followed by single agent durvalumab as adjuvant treatment after surgery? Yes No

ii. Are there any epidermal growth factor receptor (EGFR) mutations or anaplastic lymphoma kinase (ALK) rearrangements? Yes No

A. Does member have unresectable stage II or III NSCLC? Yes No

i. If yes, has member's disease progressed following concurrent platinum-based chemotherapy and radiation therapy? Yes No

B. Does member have metastatic NSCLC? Yes No

i. If yes, does member have an epidermal growth factor (EGFR) mutation or anaplastic lymphoma kinase (ALK) genomic tumor aberrations? Yes No

ii. Will durvalumab be used in conjunction with Imjudo[®] (tremelimumab-actl) and platinum-based chemotherapy? Yes No

Biliary Tract Cancer

A. Does member have locally advanced or metastatic biliary tract cancer? Yes No

B. Will durvalumab be used in combination with gemcitabine and cisplatin? Yes No

Extensive-Stage Small Cell Lung Cancer (ES-SCLC)

A. Will durvalumab be used in combination with etoposide and either cisplatin or carboplatin followed by single-agent maintenance? Yes No

(Page 1 of 2)

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Fax completed prior authorization request form to 888-601-8461 or submit Electronic Prior Authorization through CoverMyMeds[®] or SureScripts.
All requested data must be provided. Incomplete forms or forms without the chart notes will be returned. Pharmacy Coverage Guidelines are available at AetnaBetterHealth.com/Oklahoma.

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Criteria**For Initial Authorization:**1. Please indicate the diagnosis and information: *(continued)* **Bladder Cancer**A. Is diagnosis muscle invasive bladder cancer? Yes No B. Will durvalumab be used in combination with gemcitabine and cisplatin as neoadjuvant treatment for 4 cycles? Yes No C. Will durvalumab be followed by single-agent adjuvant treatment following radical cystectomy for up to 8 additional cycles? Yes No **Hepatocellular Carcinoma (HCC)**A. Does member have a diagnosis of unresectable HCC? Yes No B. Will durvalumab be used in combination with Imjudo® (tremelimumab-actl)? Yes No C. Will durvalumab be used as a single agent? Yes No **Endometrial Cancer**A. Is diagnosis primary advanced (FIGO measurable stage III/newly diagnosed stage IV) or recurrent endometrial cancer? Yes No B. Mismatch repair deficient (dMMR)? Yes No C. Will durvalumab be used in combination with carboplatin and paclitaxel followed by single-agent maintenance? Yes No **Limited-Stage Small Cell Lung Cancer (LS-SCLC)**A. Has disease progressed following concurrent platinum-based chemotherapy and radiation therapy? Yes No B. Will durvalumab be used as a single agent? Yes No

Additional Information: _____

For Continued Authorization:

1. Date of last dose: _____

2. Does member have any evidence of progressive disease while on durvalumab? Yes No 3. Has the member experienced adverse drug reactions related to durvalumab therapy? Yes No

If yes, please specify adverse reactions: _____

(Page 2 of 2)**Prescriber Signature:** _____ **Date:** _____**I certify that the indicated treatment is medically necessary and all information is true and correct to the best of my knowledge. Failure to complete this form in full will result in processing delays.****CONFIDENTIALITY NOTICE**

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