#### **Regimen Monograph**

 Regimen Name
 Drug Regimen
 Cycle Frequency
 Premedication and Supportive Measures
 Administrative Information

 References
 Other Notes
 Disclaimer

A - Regimen Name

# ABIRNIRPPRED Regimen

Abiraterone-Niraparib-Prednisone

Disease Site Genitourinary

Prostate

Intent Palliative

## Regimen Evidence-informed :

Category

Regimen is considered appropriate as part of the standard care of patients; meaningfully improves outcomes (survival, quality of life), tolerability or costs compared to alternatives (recommended by the Disease Site Team and national consensus body e.g. pan-Canadian Oncology Drug Review, pCODR). Recommendation is based on an appropriately conducted phase III clinical trial relevant to the Canadian context OR (where phase III trials are not feasible) an appropriately sized phase II trial. Regimens where one or more drugs are not approved by Health Canada for any indication will be identified under Rationale and Use.

This **Regimen Abstract** is an **abbreviated** version of a Regimen Monograph and contains only top level information on usage, dosing, schedule, cycle length and special notes (if available). Information in regimen abstracts is accurate to the extent of the ST-QBP regimen master listings, and has not undergone the full review process of a regimen monograph. Full regimen monographs will be published for each ST-QBP regimen as they are developed.

Rationale andFor the treatment of metastatic castration-resistant prostate cancer (mCRPC)Usesin patients with a BRCA mutation

Refer to EAP criteria for funding details.

Any use of the information is subject, at all times, to CCO's Terms and Conditions.

Supplementary Public Funding	niraparib / abiraterone Exceptional Access Program (niraparib / abiraterone - For the treatment of metastatic castration resistant prostate cancer in patients with a BRCA mutation) (EAP Website)			
	<b>prednisone</b> ODB - General Benefit (prednisone) ( <u>ODB Formulary</u> )			
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B - Drug Regimen				
Patients should continue to receive a GnRH agonist unless they have had prior bilateral orchiectomy.				
<u>niraparib / abirate</u>	erone*	200 mg / 1000 mg	PO	Daily
*Available as a combination product with 100 mg niraparib / 500 mg abiraterone, or 50 mg niraparib / 500 mg abiraterone per tablet				
prednisone		10 mg	PO	Daily
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C - Cycle Frequency				
CONTINUOUS TREATMENT				
Until disease progression or unacceptable toxicity				
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D - Premedication and Supportive Measures				
Antiemetic Regim	<b>ien:</b> Modera	te – Consider prophy	/laxis daily	
<ul> <li>Also refer to <u>CCO Antiemetic Recommendations</u>.</li> </ul>				

Screen for hepatitis B virus in all cancer patients starting systemic treatment. Refer to the <u>hepatitis B virus screening and management</u> guideline.

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## J - Administrative Information

Outpatient prescription for home administration

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## K - References

CADTH reimbursement recommendation. Niraparib-abiraterone (Akeega). February 2024.

Chi KN, Sandhu S, Smith MR, et al. Niraparib plus abiraterone acetate with prednisone in patients with metastatic castration-resistant prostate cancer and homologous recombination repair gene alterations: second interim analysis of the randomized phase III MAGNITUDE trial. Ann Oncol 2023 Sep;34(9):772-82.

March 2025 Updated EAP funding (niraparib/abiraterone)

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### M - Disclaimer

#### **Regimen Abstracts**

A Regimen Abstract is an abbreviated version of a Regimen Monograph and contains only top level information on usage, dosing, schedule, cycle length and special notes (if available). It is intended for healthcare providers and is to be used for informational purposes only. It is not intended to constitute or be a substitute for medical advice, and all uses of the Regimen Abstract are subject to clinical judgment. Such information is provided on an "as-is" basis, without any representation, warranty, or condition, whether express, or implied, statutory or otherwise, as to the information's quality, accuracy, currency, completeness, or reliability, and Cancer Care Ontario disclaims all liability for the use of this information, and for any claims, actions, demands or suits that arise from such use.

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#### **Regimen Monographs**

Refer to the <u>New Drug Funding Program</u> or <u>Ontario Public Drug Programs</u> websites for the most up-to-date public funding information.

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The information set out in the drug monographs, regimen monographs, appendices and symptom management information (for health professionals) contained in the Drug Formulary (the "Formulary") is intended for healthcare providers and is to be used for informational purposes only. The information is not intended to cover all possible uses, directions, precautions, drug interactions or adverse effects of a particular drug, nor should it be construed to indicate that use of a particular drug is safe, appropriate or effective for a given condition. The information in the Formulary is not intended to constitute or be a substitute for medical advice and should not be relied upon in any such regard. All uses of the Formulary are subject to clinical judgment and actual prescribing patterns may not follow the information provided in the Formulary.

The format and content of the drug monographs, regimen monographs, appendices and symptom management information contained in the Formulary will change as they are reviewed and revised on a periodic basis. The date of last revision will be visible on each page of the monograph and regimen. Since standards of usage are constantly evolving, it is advised that the Formulary not be used as the sole source of information. It is strongly recommended that original references or product monograph be consulted prior to using a chemotherapy regimen for the first time.

Some Formulary documents, such as the medication information sheets, regimen information sheets and symptom management information (for patients), are intended for patients. Patients should always consult with their healthcare provider if they have questions regarding any information set out in the Formulary documents.

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