

NUBEQA- darolutamide tablet, film coated
Bayer HealthCare Pharmaceuticals Inc.

HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use NUBEQA safely and effectively. See full prescribing information for NUBEQA.

NUBEQA® (darolutamide) tablets, for oral use
Initial U.S. Approval: 2019

-----**RECENT MAJOR CHANGES**-----

Indications and Usage (1)	6/2025
Dosage and Administration (2.1)	6/2025
Dosage and Administration (2.1)	X/XXXX
Warnings and Precautions (5.1, 5.2)	6/2025

-----**INDICATIONS AND USAGE**-----

NUBEQA is an androgen receptor inhibitor indicated for the treatment of adult patients with:

- non-metastatic castration-resistant prostate cancer (nmCRPC). (1)
- metastatic castration-sensitive prostate cancer (mCSPC). (1)
- metastatic castration-sensitive prostate cancer (mCSPC) in combination with docetaxel. (1)

-----**DOSAGE AND ADMINISTRATION**-----

Recommended Dosage: NUBEQA 600 mg (two 300 mg tablets) administered orally twice daily. Swallow tablets whole. Take NUBEQA with food. (2.1)

For patients with mCSPC treated with NUBEQA in combination with docetaxel, administer the first cycle of docetaxel within 6 weeks after the start of NUBEQA treatment. (2.1)

Patients should also receive a gonadotropin-releasing hormone (GnRH) agonist or antagonist concurrently or have had bilateral orchiectomy. (2.1)

-----**DOSAGE FORMS AND STRENGTHS**-----

Tablets: 300 mg (3)

-----**CONTRAINDICATIONS**-----

None. (4)

-----**WARNINGS AND PRECAUTIONS**-----

- Ischemic Heart Disease: Optimize management of cardiovascular risk factors. Monitor for signs and symptoms of coronary artery disease. Discontinue NUBEQA for Grade 3-4 events. (5.1)
- Seizure: Consider discontinuation of NUBEQA in patients who develop a seizure during treatment. (5.2)
- Embryo-Fetal Toxicity: NUBEQA can cause fetal harm and loss of pregnancy. Advise males with female partners of reproductive potential to use effective contraception. (5.3, 8.1, 8.3)

-----**ADVERSE REACTIONS**-----

In nmCRPC and mCSPC: The most common adverse reactions (>10% with a \geq 2% increase over placebo), including laboratory test abnormalities, are increased AST, decreased neutrophil count, increased bilirubin, fatigue, and increased ALT. (6.1)

In mCSPC in combination with docetaxel: The most common adverse reactions (\geq 10% with a \geq 2% increase over placebo) are constipation, rash, decreased appetite, hemorrhage, increased weight, and hypertension.

The most common laboratory test abnormalities (\geq 30%) are anemia, hyperglycemia, decreased lymphocyte count, decreased neutrophil count, increased AST, increased ALT, and hypocalcemia. (6.1)

To report SUSPECTED ADVERSE REACTIONS, contact Bayer HealthCare Pharmaceuticals Inc. at 1-888-842-2937 or FDA at 1-800-FDA-1088 or WWW.FDA.GOV/MEDWATCH.

-----**DRUG INTERACTIONS**-----

- Combined P-gp and Strong or Moderate CYP3A Inducers: Avoid concomitant use. (7.1)
- Combined P-gp and Strong CYP3A Inhibitors: Monitor patients more frequently for NUBEQA adverse reactions. (7.1)

- BCRP Substrates: Avoid concomitant use with drugs that are BCRP substrates where possible. If used together, monitor patients more frequently for adverse reactions and consider dose reduction of the BCRP substrate drug. (7.2)
- OATP1B1 and OATP1B3 Substrates: Concomitant use of NUBEQA may increase the plasma concentrations of OATP1B1 or OATP1B3 substrates. If used together, monitor patients more frequently for adverse reactions and consider dose reduction of these drugs. (7.2)

----- **USE IN SPECIFIC POPULATIONS** -----

- Severe Renal Impairment (not on hemodialysis): Recommended dose is 300 mg twice daily. (8.6)
- Moderate Hepatic Impairment: Recommended dose is 300 mg twice daily. (8.7)

See 17 for PATIENT COUNSELING INFORMATION and FDA-approved patient labeling.

Revised: 2/2026

FULL PRESCRIBING INFORMATION: CONTENTS*

1 INDICATIONS AND USAGE

2 DOSAGE AND ADMINISTRATION

2.1 Recommended Dosage

2.2 Dosage Modification

2.3 Recommended Dosage in Patients with Severe Renal Impairment

2.4 Recommended Dosage in Patients with Moderate Hepatic Impairment

3 DOSAGE FORMS AND STRENGTHS

4 CONTRAINDICATIONS

5 WARNINGS AND PRECAUTIONS

5.1 Ischemic Heart Disease

5.2 Seizure

5.3 Embryo-Fetal Toxicity

6 ADVERSE REACTIONS

6.1 Clinical Trials Experience

7 DRUG INTERACTIONS

7.1 Effects of Other Drugs on NUBEQA

7.2 Effects of NUBEQA on Other Drugs

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

8.2 Lactation

8.3 Females and Males of Reproductive Potential

8.4 Pediatric Use

8.5 Geriatric Use

8.6 Renal Impairment

8.7 Hepatic Impairment

10 OVERDOSAGE

11 DESCRIPTION

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

12.2 Pharmacodynamics

12.3 Pharmacokinetics

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

14 CLINICAL STUDIES

16 HOW SUPPLIED/STORAGE AND HANDLING

17 PATIENT COUNSELING INFORMATION

* Sections or subsections omitted from the full prescribing information are not listed.

FULL PRESCRIBING INFORMATION

1 INDICATIONS AND USAGE

NUBEQA is indicated for the treatment of adult patients with:

- non-metastatic castration resistant prostate cancer (nmCRPC)
- metastatic castration-sensitive prostate cancer (mCSPC)
- metastatic castration-sensitive prostate cancer (mCSPC) in combination with docetaxel.

2 DOSAGE AND ADMINISTRATION

2.1 Recommended Dosage

The recommended dose of NUBEQA is 600 mg (two 300 mg tablets) taken orally, twice daily, with food [see *Clinical Pharmacology (12.3)*].

Continue treatment until disease progression or unacceptable toxicity occurs.

Patients receiving NUBEQA should also receive a gonadotropin-releasing hormone (GnRH) agonist or antagonist concurrently or have had a bilateral orchiectomy.

When used in combination with docetaxel for mCSPC, administer the first of 6 cycles of docetaxel within 6 weeks after the start of NUBEQA treatment. Refer to docetaxel prescribing information for additional dosing information, including dosage modifications. Treatment with NUBEQA may be continued until disease progression or unacceptable toxicity, even if a cycle of docetaxel is delayed, interrupted, or discontinued [see *Dosage and Administration (2.2)*].

Advise patients to swallow tablets whole with food, to take any missed dose as soon as they remember. A missed dose may be taken together with the next scheduled dose. Thereafter, resume the regularly scheduled twice daily dosing.

2.2 Dosage Modification

If a patient experiences a greater than or equal to Grade 3 or an intolerable adverse reaction, withhold NUBEQA or reduce dosage to 300 mg twice daily until symptoms improve. NUBEQA may be resumed at a dose of 600 mg twice daily, when adverse reaction returns to baseline [see *Adverse Reactions (6.1)*]. Dosage reduction below 300 mg twice daily is not recommended.

For patients who experience ischemic heart disease or seizure, additional dose modifications may be required [see *Warnings and Precautions (5.1 and 5.2)*].

2.3 Recommended Dosage in Patients with Severe Renal Impairment

For patients with severe renal impairment (eGFR 15–29 mL/min/1.73 m²) not receiving hemodialysis, the recommended dose of NUBEQA is 300 mg twice daily [see *Use in Specific Populations (8.6) and Clinical Pharmacology (12.3)*].

2.4 Recommended Dosage in Patients with Moderate Hepatic Impairment

For patients with moderate hepatic impairment (Child-Pugh Class B), the recommended dose of NUBEQA is 300 mg twice daily [see *Use in Specific Populations (8.7) and Clinical Pharmacology (12.3)*].

3 DOSAGE FORMS AND STRENGTHS

Tablets (300 mg): white to off-white oval film-coated tablets marked with "300" on one side and "Bayer" on the other.

4 CONTRAINDICATIONS

None.

5 WARNINGS AND PRECAUTIONS

5.1 Ischemic Heart Disease

Ischemic heart disease, including fatal cases, occurred in patients receiving NUBEQA.

In a pooled analysis of ARAMIS and ARANOTE, ischemic heart disease occurred in 3.4% of patients receiving NUBEQA and 2.2% receiving placebo, including Grade 3-4 events in 1.4% and 0.3%, respectively. Ischemic events led to death in 0.4% of patients receiving NUBEQA and 0.4% receiving placebo.

In ARASENS, ischemic heart disease occurred in 3.2% of patients receiving NUBEQA with docetaxel and 2% receiving placebo with docetaxel, including Grade 3-4 events in 1.3% and 1.1%, respectively. Ischemic events led to death in 0.3% of patients receiving NUBEQA with docetaxel and 0% receiving placebo with docetaxel.

Monitor for signs and symptoms of ischemic heart disease. Optimize management of cardiovascular risk factors, such as hypertension, diabetes, or dyslipidemia. Discontinue NUBEQA for Grade 3-4 ischemic heart disease.

5.2 Seizure

Seizure occurred in patients receiving NUBEQA.

In a pooled analysis of ARAMIS and ARANOTE, Grade 1-3 seizure occurred in 0.2% of patients receiving NUBEQA. Seizure occurred from 261 to 665 days after initiation of NUBEQA.

In ARASENS, seizure occurred in 0.8% of patients receiving NUBEQA with docetaxel, including two Grade 3 events. Seizure occurred from 38 to 1754 days after initiation of NUBEQA.

It is unknown whether anti-epileptic medications will prevent seizures with NUBEQA. Advise patients of the risk of developing a seizure while receiving NUBEQA and of engaging in any activity where sudden loss of consciousness could cause harm to themselves or others. Consider discontinuation of NUBEQA in patients who develop a seizure during treatment.

5.3 Embryo-Fetal Toxicity

The safety and efficacy of NUBEQA have not been established in females. Based on its mechanism of action, NUBEQA can cause fetal harm and loss of pregnancy when administered to a pregnant female [see *Clinical Pharmacology (12.1)*].

Advise males with female partners of reproductive potential to use effective contraception during treatment and for 1 week after the last dose of NUBEQA [see *Use in Specific Populations (8.1, 8.3)*].

6 ADVERSE REACTIONS

6.1 Clinical Trials Experience

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in practice.

The pooled data in WARNINGS and PRECAUTIONS reflect two randomized clinical trials [ARAMIS, ARANOTE] in patients with nmCRPC (N = 954) and mCSPC (N = 445) treated with NUBEQA. In these trials, the median duration of treatment was 18.2 months (range 0.03 to 44.3) for patients who received NUBEQA [see *Clinical Studies (14)*]. In this pooled safety population, the most common adverse reactions (>10% with a $\geq 2\%$ increase over placebo), including laboratory test abnormalities were increased AST, decreased neutrophil count, increased bilirubin, fatigue and increased ALT.

The safety of NUBEQA in combination with docetaxel in mCSPC is based on data from 1302 patients of whom 652 received at least one dose of NUBEQA in the ARASENS study [see *Clinical Studies (14)*].

Non-Metastatic Castration Resistant Prostate Cancer

ARAMIS

The safety of NUBEQA in nmCRPC patients was evaluated in ARAMIS, a randomized (2:1), double-blind, placebo-controlled, multi-center clinical study [see *Clinical Studies (14)*]. Patients received either NUBEQA at a dose of 600 mg, or a placebo, twice a day. All patients in the ARAMIS study received a concomitant gonadotropin-releasing hormone (GnRH) analog or had a bilateral orchiectomy. Among patients who received NUBEQA, the median duration of exposure was 14.8 months (range: 0 to 44.3 months).

Serious adverse reactions occurred in 25% of patients receiving NUBEQA and in 20% of patients receiving placebo. Serious adverse reactions in $\geq 1\%$ of patients who received NUBEQA included urinary retention, pneumonia and hematuria. Fatal adverse reactions occurred in 3.9% of patients receiving NUBEQA and 3.2% of patients receiving placebo. Fatal adverse reactions that occurred in ≥ 2 patients who received NUBEQA included death (0.4%), cardiac failure (0.3%), cardiac arrest (0.2%), general physical health

deterioration (0.2%), and pulmonary embolism (0.2%).

Permanent discontinuation of NUBEQA due to adverse reactions occurred in 9% of patients receiving NUBEQA. The most common adverse reactions requiring permanent discontinuation in patients who received NUBEQA included cardiac failure (0.4%), and death (0.4%).

Dosage interruptions due to adverse reactions occurred in 13% of patients treated with NUBEQA. The most common adverse reactions requiring dosage interruption in patients who received NUBEQA included hypertension (0.6%), diarrhea (0.5%), and pneumonia (0.5%).

Dosage reductions due to adverse reactions occurred in 6% of patients treated with NUBEQA. The most common adverse reactions requiring dosage reduction in patients treated with NUBEQA included fatigue (0.7%), hypertension (0.3%), and nausea (0.3%).

The most common (>2% with a $\geq 2\%$ increase compared to placebo) adverse reactions, including laboratory test abnormalities, were increased AST, decreased neutrophil count, fatigue, increased bilirubin, pain in extremity, and rash.

Table 1 summarizes the adverse reactions in ARAMIS.

Table 1: Adverse Reactions (>2% with a $\geq 2\%$ increase compared to placebo) in Patients with nmCRPC in ARAMIS

Adverse Reaction	NUBEQA (N=954)		Placebo (N=554)	
	All Grades %	Grades 3 or 4 %	All Grades %	Grade 3 or 4 %
Fatigue*	16	0.6	11	1.1
Pain in extremity	6	0	3	0.2
Rash†	4	0.1	1.4	0

* Includes fatigue and asthenia

† Includes rash, eczema, rash maculo-papular, dermatitis, erythema multiforme, rash macular, rash papular, rash pustular, skin exfoliation

Clinically relevant adverse reactions occurring in 2% or more of patients treated with NUBEQA included ischemic heart disease (4%) and heart failure (2.1%).

Table 2 summarizes the laboratory test abnormalities in ARAMIS.

Table 2: Laboratory Test Abnormalities in ARAMIS

Laboratory Abnormality	NUBEQA (N=954)*		Placebo (N=554)*	
	All Grades %	Grade 3 or 4 %	All Grades %	Grade 3 or 4 %
AST increased	23	0.5	14	0.2
Neutrophil count decreased	20	4	9	0.6

Bilirubin increased	16	0.1	7	0
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* The denominator used to calculate the rate varied based on the number of patients with a baseline value and at least one post-treatment value.

Metastatic Castration-Sensitive Prostate Cancer

ARANOTE

The safety of NUBEQA in mCSPC patients was evaluated in ARANOTE, a randomized (2:1), double-blind, placebo-controlled, multi-center clinical study [see *Clinical Studies (14)*]. Patients received either NUBEQA at a dose of 600 mg, or a placebo, twice a day. All patients in the ARANOTE study received a concomitant gonadotropin-releasing hormone (GnRH) agonist or antagonist or had a bilateral orchiectomy. Among patients who received NUBEQA, the median duration of exposure was 24 months (range: 0.03 to 39 months).

Serious adverse reactions occurred in 24% of patients receiving NUBEQA. Serious adverse reactions in $\geq 1\%$ of patients who received NUBEQA included pneumonia (2%), urinary tract infection (1.8%), musculoskeletal pain (1.6%), hemorrhage (1.6%), arrhythmias (1.3%), and spinal cord compression (1.1%). Fatal adverse reactions occurred in 4.7% of patients receiving NUBEQA and those that occurred in ≥ 2 patients included sepsis (1.1%), craniocerebral injury (0.4%), and myocardial infarction (0.4%).

Permanent discontinuation of NUBEQA due to adverse reactions occurred in 6% of patients treated in the NUBEQA arm. The most common adverse reactions which resulted in permanent discontinuation of NUBEQA were increased ALT, increased AST, craniocerebral injury, myocardial infarction, and rash (all with 0.4%).

Dosage interruptions of NUBEQA due to adverse reactions occurred in 14% of patients treated in the NUBEQA arm. The most common adverse reactions requiring dosage interruption of NUBEQA were increased AST (1.6%), increased ALT (1.3%), and rash (1.3%).

Dosage reductions of NUBEQA due to adverse reactions occurred in 3.6% of patients treated in the NUBEQA arm. The most common adverse reactions requiring dosage reduction of NUBEQA were increased AST (0.7%), rash (0.7%), increased ALT (0.4%), and hypertension (0.4%).

Table 3 summarizes the adverse reactions in ARANOTE.

Table 3: Adverse Reactions ($\geq 10\%$ with a $\geq 2\%$ increase compared to placebo) in Patients with mCSPC in ARANOTE

Adverse Reaction	NUBEQA (N=445)		Placebo (N=221)	
	All Grades (%)	Grade 3 or 4 (%)	All Grades (%)	Grade 3 or 4 (%)
Urinary Tract Infection	12	1.8	8	0.5

Clinically relevant adverse reactions in $< 10\%$ of patients who received NUBEQA included arrhythmia (8.8%), pneumonia (3.6%), and myocardial infarction (0.7%).

Table 4 summarizes the laboratory test abnormalities in ARANOTE.

Table 4: Laboratory Test Abnormalities ($\geq 15\%$ with a $\geq 5\%$ increase over placebo) in ARANOTE

Laboratory Abnormality	NUBEQA (N=445)*		Placebo (N=221)*	
	All Grades† %	Grade 3 or 4† %	All Grades† %	Grade 3 or 4† %
AST increased	32	2.8	25	0.5
ALT increased	28	2.1	23	0.5
Bilirubin increased	17	0.5	7	0
Neutrophil count decreased	16	1.2	9	0.5

* The number of patients tested for a specific laboratory test parameter may be different. The incidence of each laboratory test abnormality was calculated accordingly.

† Common Terminology Criteria for Adverse Events (CTCAE) version 5.0. Grade 4 laboratory test values were limited to neutrophil count decreased.

ARASENS

The safety of NUBEQA, in combination with docetaxel, in mCSPC patients was evaluated in ARASENS, a randomized (1:1), double-blind, placebo-controlled, multi-center clinical study [see *Clinical Studies (14)*]. Patients were to receive either NUBEQA at a dose of 600 mg, or a placebo, twice a day in combination with docetaxel at a dose of 75 mg/m² every 21 days for 6 cycles. All patients in the ARASENS study received a concomitant gonadotropin-releasing hormone (GnRH) agonist or antagonist or had a bilateral orchiectomy. Among patients who received NUBEQA, the median duration of exposure was 41 months (range: 0.1 to 56.5 months) vs. 16.7 months (range 0.3 to 55.8) with placebo. Eighty-eight percent and 86% of patients received the 6 planned cycles of docetaxel, in the NUBEQA with docetaxel arm and placebo with docetaxel arm, respectively.

Serious adverse reactions occurred in 45% of patients receiving NUBEQA with docetaxel. Serious adverse reactions in $\geq 2\%$ of patients who received NUBEQA with docetaxel included febrile neutropenia (6%), neutrophil count decreased (2.8%), musculoskeletal pain (2.6%) and pneumonia (2.6%). Fatal adverse reactions occurred in 4% of patients receiving NUBEQA with docetaxel. Fatal adverse reactions in ≥ 2 patients who received NUBEQA included COVID-19/COVID-19 pneumonia (0.8%), myocardial infarction (0.3%), and sudden death (0.3%).

Permanent discontinuation of NUBEQA due to adverse reactions occurred in 14% of patients treated in the NUBEQA with docetaxel arm. The most common adverse reactions which resulted in permanent discontinuation of NUBEQA were rash (1.1%), musculoskeletal pain (0.9%), and increased aspartate aminotransferase (AST) (0.9%).

Dosage interruptions of NUBEQA due to adverse reactions occurred in 23% of patients treated in the NUBEQA with docetaxel arm. The most common ($>2\%$) adverse reactions requiring dosage interruption of NUBEQA were increased alanine aminotransferase (ALT) (3.2%), increased AST (3.1%) and febrile neutropenia (2.1%).

Dosage reductions of NUBEQA due to adverse reactions occurred in 9% of patients treated in the NUBEQA with docetaxel arm. The most common (>2%) adverse reactions requiring dosage reduction of NUBEQA were increased ALT (2.8%) and increased AST (2.5%).

The most common ($\geq 10\%$ with a $\geq 2\%$ increase over placebo with docetaxel) adverse reactions are constipation, rash, decreased appetite, hemorrhage, increased weight, and hypertension. The most common laboratory test abnormalities ($\geq 30\%$) are anemia, hyperglycemia, decreased lymphocyte count, decreased neutrophil count, increased AST, increased ALT, and hypocalcemia.

Table 5 summarizes the adverse reactions in ARASENS.

Table 5: Adverse Reactions ($\geq 10\%$ with a $\geq 2\%$ increase compared to placebo with docetaxel) in ARASENS

Adverse Reaction	NUBEQA with docetaxel (N=652)		Placebo with docetaxel (N=650)	
	All Grades %	Grades 3 or 4 %	All Grades %	Grades 3 or 4 %
Constipation	23	0.3	20	0.3
Rash*	20	1.8	15	0.2
Decreased Appetite	19	0.2	13	0.6
Hemorrhage†	18	1.4	13	1.4
Weight Increased	18	2.1	16	1.2
Hypertension‡	14	7	10	3.6

* Rash includes rash, rash maculo-papular, palmar-plantar erythrodysesthesia syndrome, eczema, dermatitis, skin exfoliation, dermatitis acneiform, drug eruption, rash pruritic, rash erythematous, erythema multiforme, rash macular, dermatitis exfoliative generalized, penile rash, dyshidrotic eczema, rash papular, dermatitis bullous, rash follicular, rash pustular, rash vesicular, toxic skin eruption

† Hemorrhage includes hematuria, epistaxis, anal hemorrhage, hemorrhoidal hemorrhage, rectal hemorrhage, upper gastrointestinal hemorrhage, hemoptysis, hemorrhage urinary tract, hemorrhagic stroke, subarachnoid hemorrhage, lower gastrointestinal hemorrhage, cystitis hemorrhagic, gastrointestinal hemorrhage, hemorrhage subcutaneous, intra-abdominal hemorrhage, nail bed bleeding, subdural hemorrhage

‡ Hypertension includes hypertension, blood pressure increased, hypertensive emergency and hypertensive crisis.

Clinically relevant adverse reactions in < 10% of patients who received NUBEQA with docetaxel included fractures (8%), ischemic heart disease (3.2%), seizures (0.6%), and drug-induced liver injury (0.3%).

Table 6 summarizes laboratory test abnormalities in the ARASENS study.

Table 6: Laboratory Test Abnormalities ($\geq 30\%$) in ARASENS

Laboratory Abnormality	NUBEQA with docetaxel* (N=652)		Placebo with docetaxel* (N=650)	
	All Grades	Grade 3-4	All Grades	Grade 3-4

	%	%	%	%
Anemia	72	6	71	7
Hyperglycemia	57	7	53	10
Lymphocyte count decreased	52	12	49	13
Neutrophil count decreased	49	33	44	31
AST increased [†]	40	3.6	35	2.3
ALT increased [†]	37	3.7	31	2.9
Hypocalcemia	31	2.8	28	1.9

* The denominator used to calculate the rate varied from 470 to 648 based on the number of patients with a baseline value and at least one post-treatment value.

† ALT or AST increases to $\geq 5 \times$ upper limit of normal (ULN) occurred in 5.3% of patients who received NUBEQA with docetaxel. ALT or AST increases to $\geq 20 \times$ ULN occurred in 0.3% of patients who received NUBEQA with docetaxel. The median time to onset of any grade ALT or AST increases was 2.8 months (range: 0.03 to 46.9).

Clinically relevant laboratory test abnormalities in < 30% of patients who received NUBEQA with docetaxel included blood bilirubin increased (all grades 20%, Grade 3-4 0.5%) compared to placebo with docetaxel (all grades 10%, grades 3-4 0.3%).

7 DRUG INTERACTIONS

7.1 Effects of Other Drugs on NUBEQA

Combined P-gp and Strong or Moderate CYP3A4 Inducer

Concomitant use of NUBEQA with a combined P-gp and strong or moderate CYP3A4 inducer decreases darolutamide exposure which may decrease NUBEQA activity [see *Clinical Pharmacology (12.3)*]. Avoid concomitant use of NUBEQA with combined P-gp and strong or moderate CYP3A4 inducers.

Combined P-gp and Strong CYP3A4 Inhibitors

Concomitant use of NUBEQA with a combined P-gp and strong CYP3A4 inhibitor increases darolutamide exposure [see *Clinical Pharmacology (12.3)*] which may increase the risk of NUBEQA adverse reactions. Monitor patients more frequently for NUBEQA adverse reactions and modify NUBEQA dosage as needed [see *Dosage and Administration (2.2)*].

7.2 Effects of NUBEQA on Other Drugs

Breast Cancer Resistance Protein (BCRP) and Organic Anion Transporting Polypeptides (OATP) 1B1 and 1B3 Substrates

NUBEQA is an inhibitor of BCRP transporter. Concomitant use of NUBEQA increases the AUC and C_{max} of BCRP substrates [see *Clinical Pharmacology (12.3)*], which may increase the risk of BCRP substrate-related toxicities.

Avoid concomitant use with drugs that are BCRP substrates where possible. If used together, monitor patients more frequently for adverse reactions, and consider dose reduction of the BCRP substrate drug.

NUBEQA is an inhibitor of OATP1B1 and OATP1B3 transporters. Concomitant use of NUBEQA may increase the plasma concentrations of OATP1B1 or OATP1B3 substrates. Monitor patients more frequently for adverse reactions of these drugs and consider dose reduction while patients are taking NUBEQA [see *Clinical Pharmacology (12.3)*], Review the prescribing information of the BCRP, OATP1B1 and OATP1B3 substrates when used concomitantly with NUBEQA.

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

The safety and efficacy of NUBEQA have not been established in females. Based on its mechanism of action, NUBEQA can cause fetal harm and loss of pregnancy [see *Clinical Pharmacology (12.1)*]. Animal embryo-fetal developmental toxicology studies were not conducted with darolutamide. There are no human data on the use of NUBEQA in pregnant females.

8.2 Lactation

Risk Summary

The safety and efficacy of NUBEQA have not been established in females. There are no data on the presence of darolutamide or its metabolites in human milk, the effect on the breastfed child, or the effect on milk production.

8.3 Females and Males of Reproductive Potential

Contraception

Males

Based on the mechanism of action, advise male patients with female partners of reproductive potential to use effective contraception during treatment and for 1 week after the last dose of NUBEQA [see *Use in Specific Populations (8.1)*].

Infertility

Males

Based on animal studies, NUBEQA may impair fertility in males of reproductive potential [see *Nonclinical Toxicology (13.1)*].

8.4 Pediatric Use

Safety and effectiveness of NUBEQA in pediatric patients have not been established.

8.5 Geriatric Use

Of the 954 patients who received NUBEQA in ARAMIS, 88% of patients were 65 years and over, and 49% were 75 years and over. Of the 445 patients who received NUBEQA in ARANOTE, 74% of patients were 65 years and over, and 30% were 75 years and over. Of the 652 patients who received NUBEQA in ARASENS, 63% of patients were 65

years and over, and 16% were 75 years and over. No overall differences in safety or efficacy were observed between these patients and younger patients in both studies.

8.6 Renal Impairment

Patients with severe renal impairment (eGFR 15–29 mL/min/1.73 m²) who are not receiving hemodialysis have a higher exposure to NUBEQA and reduction of the dose is recommended [see *Dosage and Administration (2.3)* and *Clinical Pharmacology (12.3)*]. No dose reduction is needed for patients with mild or moderate renal impairment (eGFR 30–89 mL/min/1.73 m²). The effect of end stage renal disease (eGFR ≤15 mL/min/1.73 m²) on darolutamide pharmacokinetics is unknown.

8.7 Hepatic Impairment

Patients with moderate hepatic impairment (Child-Pugh Class B) have a higher exposure to NUBEQA and reduction of the dose is recommended [see *Dosage and Administration (2.4)* and *Clinical Pharmacology (12.3)*]. No dose reduction is needed for patients with mild hepatic impairment. The effect of severe hepatic impairment (Child-Pugh C) on darolutamide pharmacokinetics is unknown.

10 OVERDOSAGE

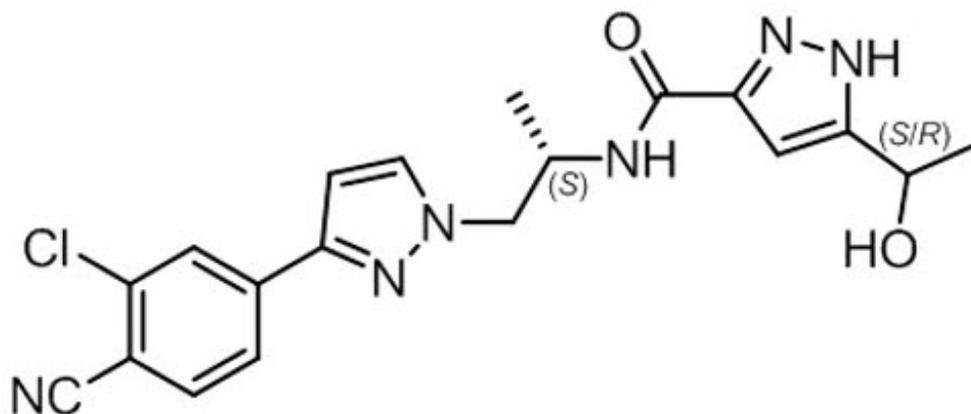
There is no known specific antidote for darolutamide overdose. The highest dose of NUBEQA studied clinically was 900 mg twice daily, equivalent to a total daily dose of 1800 mg. No dose limiting toxicities were observed with this dose.

Considering the saturable absorption and the absence of evidence for acute toxicity, an intake of a higher than recommended dose of darolutamide is not expected to lead to systemic toxicity in patients with intact hepatic and renal function [see *Clinical Pharmacology (12.3)*].

In the event of intake of a higher than recommended dose in patients with severe renal impairment or moderate hepatic impairment, if there is suspicion of toxicity, interrupt NUBEQA treatment and undertake general supportive measures until clinical toxicity has been diminished or resolved. If there is no suspicion of toxicity, NUBEQA treatment can be continued with the next dose as scheduled.

11 DESCRIPTION

NUBEQA is an androgen receptor inhibitor. The chemical name is N-{(2S)-1-[3-(3-chloro-4-cyanophenyl)-1H-pyrazol-1-yl]propan-2-yl}-5-(1-hydroxyethyl)-1H-pyrazole-3-carboxamide.



The molecular weight is 398.85 and the molecular formula is $C_{19}H_{19}ClN_6O_2$. The structural formula is:

Darolutamide is an optically active with a specific rotation value $[\alpha]_D^{20} = 72.2$ $^{\circ}mL/(dm \cdot g)$, white to greyish- or yellowish white crystalline powder, that is soluble in tetrahydrofuran, but practically insoluble in aqueous medium. Darolutamide has a pKa of 11.75.

NUBEQA (darolutamide) is supplied as film-coated tablets containing 300 mg of darolutamide for oral use. The inactive ingredients of the tablet are: calcium hydrogen phosphate, croscarmellose sodium, lactose monohydrate, magnesium stearate, povidone K 30, hypromellose 15 cP, macrogol 3350, and titanium dioxide.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Darolutamide is an androgen receptor (AR) inhibitor. Darolutamide competitively inhibits androgen binding, AR nuclear translocation, and AR-mediated transcription. A major metabolite, ketodarolutamide, exhibited similar *in vitro* activity to darolutamide. In addition, darolutamide functioned as a progesterone receptor (PR) antagonist *in vitro* (approximately 1% activity compared to AR). Darolutamide decreased prostate cancer cell proliferation *in vitro* and tumor volume in mouse xenograft models of prostate cancer.

12.2 Pharmacodynamics

PSA reduction was observed in CRPC patients receiving darolutamide doses of 100-900 mg twice a day, reaching a plateau of PSA reduction at the 600 mg twice daily dose.

Twice daily dosing of 600 mg darolutamide in nmCRPC patients resulted in undetectable PSA levels in 24% of patients at 12 months compared to 0.4% of patients in the placebo arm.

Twice daily dosing of 600 mg darolutamide in mCSPC patients resulted in undetectable PSA levels (<0.2 ng/mL) in 63% of patients compared to 19% of patients in the placebo arm.

Twice daily dosing of 600 mg darolutamide in combination with docetaxel in mCSPC patients resulted in undetectable PSA levels in 60% of patients at 12 months compared

to 26% of patients in the placebo with docetaxel arm.

Cardiac Electrophysiology

The effect of darolutamide (600 mg twice daily) on the QTc interval was evaluated in a subgroup of 500 patients in the ARAMIS study. No large mean increase in QTc (i.e., > 20 ms) was detected.

12.3 Pharmacokinetics

Following administration of 600 mg twice daily, darolutamide mean (%CV) steady-state peak plasma concentration (C_{max}) is 4.79 mg/L (30.9%) and area under the plasma concentration-time curve from time 0 to 12 hours (AUC_{12h}) is 52.82 h• μ g/mL (33.9%). Steady-state is reached 2–5 days after repeated dosing with food, with an approximate 2-fold accumulation.

The exposure (C_{max} and AUC_{12}) of the darolutamide and the active metabolite ketodarolutamide increase in a nearly dose-proportional manner in the dose range of 100 to 700 mg (0.17 to 1.17 times the approved recommended dosage). No further increase in darolutamide exposure was observed at 900 mg twice daily (1.5 times the approved recommended dosage).

Absorption

Darolutamide C_{max} is reached approximately 4 hours after administration of a single 600 mg oral dose.

The absolute bioavailability is approximately 30% following oral administration of a NUBEQA tablet containing 300 mg darolutamide under fasted conditions.

Food Effect

Bioavailability of darolutamide increased by 2.0- to 2.5-fold when administered with food. A similar increase of exposure was observed for the active metabolite ketodarolutamide.

Distribution

The apparent volume of distribution of darolutamide after intravenous administration is 119 L.

Protein binding is 92% for darolutamide and 99.8% for the active metabolite, ketodarolutamide. Serum albumin is the main binding protein for darolutamide and ketodarolutamide.

Elimination

The effective half-life of darolutamide and ketodarolutamide is approximately 20 hours in patients. The clearance (%CV) of darolutamide following intravenous administration is 116 mL/min (39.7%).

Metabolism

Darolutamide is primarily metabolized by CYP3A4, as well as by UGT1A9 and UGT1A1. Ketodarolutamide total exposure in plasma is 1.7-fold higher compared to darolutamide.

Excretion

After a single radiolabeled dose as an oral solution, a total of 63.4% of darolutamide-related material is excreted in the urine (approximately 7% unchanged)

and 32.4% (approximately 30% unchanged) in the feces. More than 95% of the dose was recovered within 7 days after administration.

Specific Populations

No clinically significant differences in the pharmacokinetics of darolutamide were observed based on age (41-95 years), race (White, Asian, Black or African American), mild to moderate renal impairment (eGFR 30–89 mL/min/1.73m²), or mild hepatic impairment.

In non-cancer subjects with severe renal impairment (eGFR 15–29 mL/min/1.73 m²) not receiving dialysis or with moderate hepatic impairment (Child-Pugh Class B), NUBEQA exposure increased by about 2.5- and 1.9-fold, respectively, compared to healthy subjects.

The effect of end-stage renal disease (eGFR <15 mL/min/1.73 m²) or severe hepatic impairment (Child-Pugh C) on darolutamide pharmacokinetics has not been studied.

Drug Interaction Studies

Clinical Studies

Combined P-gp and Strong CYP3A4 Inducers

Concomitant use of rifampicin (a combined P-gp and strong CYP3A4 inducer) decreased mean darolutamide AUC₀₋₇₂ by 72% and C_{max} by 52%. The decrease of darolutamide exposure by moderate CYP3A4 inducers is expected to be in the range of 36% – 58%.

Combined P-gp and Strong CYP3A4 Inhibitors

Itraconazole (a strong combined CYP3A4 and P-gp inhibitor) increased mean darolutamide AUC₀₋₇₂ by 1.7- and C_{max} by 1.4-fold.

CYP3A4 substrates

Concomitant use of darolutamide decreased the mean AUC and C_{max} of midazolam (CYP3A4 substrate) by 29% and 32%, respectively. No clinically relevant differences in the pharmacokinetics of midazolam were observed when used concomitantly with darolutamide.

BCRP, OATP1B1 and OATP1B3 Substrates

Concomitant use of darolutamide increased the mean AUC and C_{max} of rosuvastatin (BCRP, OATP1B1 and OATP1B3 substrate) by approximately 5-fold.

Docetaxel

Administration of darolutamide in combination with docetaxel resulted in no clinically relevant changes in the pharmacokinetics of docetaxel in mCSPC patients. There were no clinically relevant changes in the pharmacokinetics of darolutamide, when used in combination with docetaxel.

P-gp Substrates

No clinically relevant differences in the pharmacokinetics of dabigatran (P-gp substrate) were observed when used concomitantly with darolutamide.

In vitro, darolutamide did not inhibit the major CYP enzymes (CYP1A2, 2A6, 2B6, 2C8, 2C9, 2C19, 2D6, 2E1, and 3A4) or transporters (MRP2, BSEP, OATs, OCTs, MATEs,

OATP2B1, and NTCP) at clinically relevant concentrations.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Oral administration of darolutamide to male and female RasH2 transgenic mice for 26 weeks did not show carcinogenic potential at doses up to 1000 mg/kg/day.

Darolutamide was clastogenic in an *in vitro* chromosome aberration assay in human peripheral blood lymphocytes. Darolutamide did not induce mutations in the bacterial reverse mutation (Ames) assay and was not genotoxic in the *in vivo* combined bone marrow micronucleus assay and the Comet assay in the liver and duodenum of the rat.

Fertility studies in animals have not been conducted with darolutamide. In repeat-dose toxicity studies in male rats (up to 26 weeks) and dogs (up to 39 weeks), tubular dilatation of testes, hypospermia, and atrophy of seminal vesicles, testes, prostate gland and epididymides were observed at doses ≥ 100 mg/kg/day in rats (0.6 times the human exposure based on AUC) and ≥ 50 mg/kg/day in dogs (approximately 1 times the human exposure based on AUC).

14 CLINICAL STUDIES

Non-Metastatic Castration Resistant Prostate Cancer

ARAMIS (NCT02200614) was a multicenter, double-blind, placebo-controlled clinical trial in 1509 patients with non-metastatic castration resistant prostate cancer with a prostate-specific antigen doubling time (PSADT) of ≤ 10 months. Randomization was stratified by PSADT and use of bone-targeted therapy at study entry. Patients with pelvic lymph nodes less than 2 cm in short axis below the aortic bifurcation were eligible to enter the study. Absence or presence of metastasis was assessed by blinded independent central review (BICR). PSA results were not blinded and were not used for treatment discontinuation.

Patients were randomized 2:1 to receive either 600 mg NUBEQA orally twice daily (n=955) or matching placebo (n=554). Treatment continued until radiographic disease progression as assessed by CT, MRI, ^{99m}Tc bone scan by BICR, unacceptable toxicity or withdrawal. All patients received a gonadotropin-releasing hormone (GnRH) analog concurrently or had a bilateral orchiectomy.

The median age was 74 years (range 48–95) and 9% of patients were 85 years of age or older. The racial distribution included 79% White, 13% Asian, 3% Black or African American, while ethnicity included 3.1% Hispanic or Latino patients. A majority of patients (73%) had a Gleason score of 7 or higher at diagnosis. The median PSADT was 4.5 months. Forty-two percent of patients in both treatment arms had prior surgery or radiotherapy to the prostate. Eleven percent of patients had enlarged pelvic lymph nodes less than 2 cm at study entry. Six percent of patients were retrospectively identified by BICR as having metastases at baseline. Seventy-three percent of patients received prior treatment with an anti-androgen (bicalutamide or flutamide). All patients had an Eastern Cooperative Oncology Group Performance Status (ECOG PS) score of 0 or 1 at study entry. At baseline, 47% of patients reported no pain on the Brief Pain Inventory-Short Form (a 7-day diary average of the daily worst pain item). There were

12 patients (1.3%) enrolled on the NUBEQA arm with a history of seizure.

The major efficacy endpoint was metastasis free survival (MFS), defined as the time from randomization to the time of first evidence of BICR-confirmed distant metastasis or death from any cause within 33 weeks after the last evaluable scan, whichever occurred first. Distant metastasis was defined as new bone or soft tissue lesions or enlarged lymph nodes above the aortic bifurcation. Overall survival (OS), time to pain progression, and time to initiation of cytotoxic chemotherapy, were additional efficacy endpoints.

Treatment with NUBEQA resulted in a statistically significant improvement in MFS compared to placebo. At the protocol-specified final analysis of OS, treatment with NUBEQA resulted in a statistically significant improvement in OS compared to placebo. The final analysis of OS and time to initiation of cytotoxic chemotherapy was event-driven and conducted after 254 OS events had occurred. The efficacy results from ARAMIS are summarized in Table 7 and Figure 1.

Table 7: Efficacy Results from the ARAMIS Study

	NUBEQA (N=955)	Placebo (N=554)
Metastasis-free survival*		
Distant Metastasis or Death (%)	221 (23)	216 (39)
Median, months (95% CI)†	40.4 (34.3, NR)	18.4 (15.5, 22.3)
Hazard Ratio (95% CI)‡	0.41 (0.34, 0.50)	
P-value§	<0.0001	
Overall survival§		
Deaths (%)	148 (15)	106 (19)
Median, months (95% CI)†	NR (56.1, NR)	NR (46.9, NR)
Hazard Ratio (95% CI)‡	0.69 (0.53, 0.88)	
P-value¶	0.003	

NR: not reached

* Locoregional-only progression occurred in 6% of patients overall.

† Based on Kaplan-Meier estimates

‡ Hazard ratio is based on a stratified Cox regression model. Hazard ratio < 1 favors NUBEQA.

§ The pre-specified final OS analysis was event-driven and occurred 14 months after the MFS analysis

¶ P-value is based on a log-rank test stratified by PSADT (\leq 6 months vs. > 6 months) and use of osteoclast-targeted therapy (yes vs. no)

Figure 1: Kaplan-Meier Curve Metastasis Free Survival; Intent-To-Treat nmCRPC population (ARAMIS)

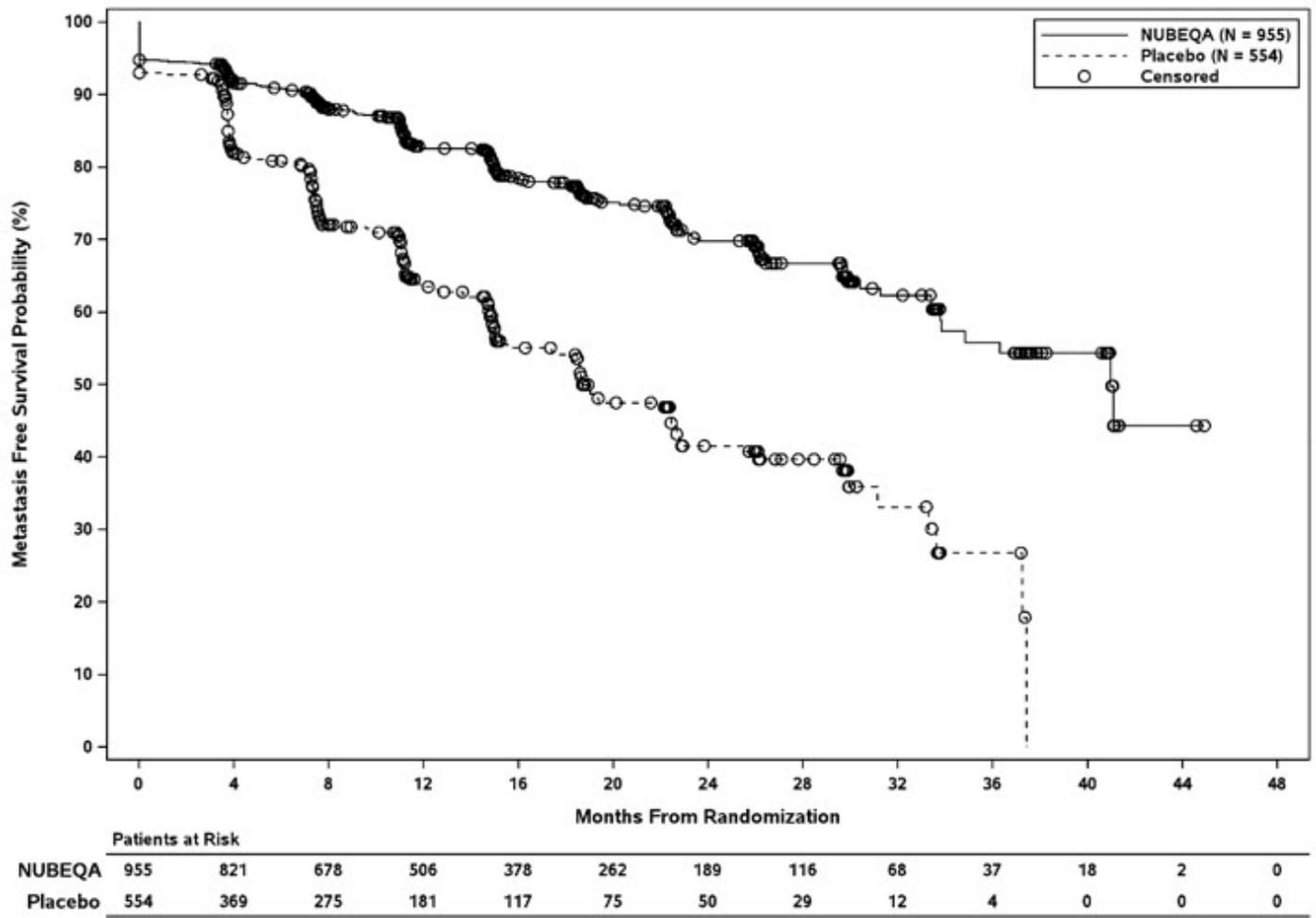
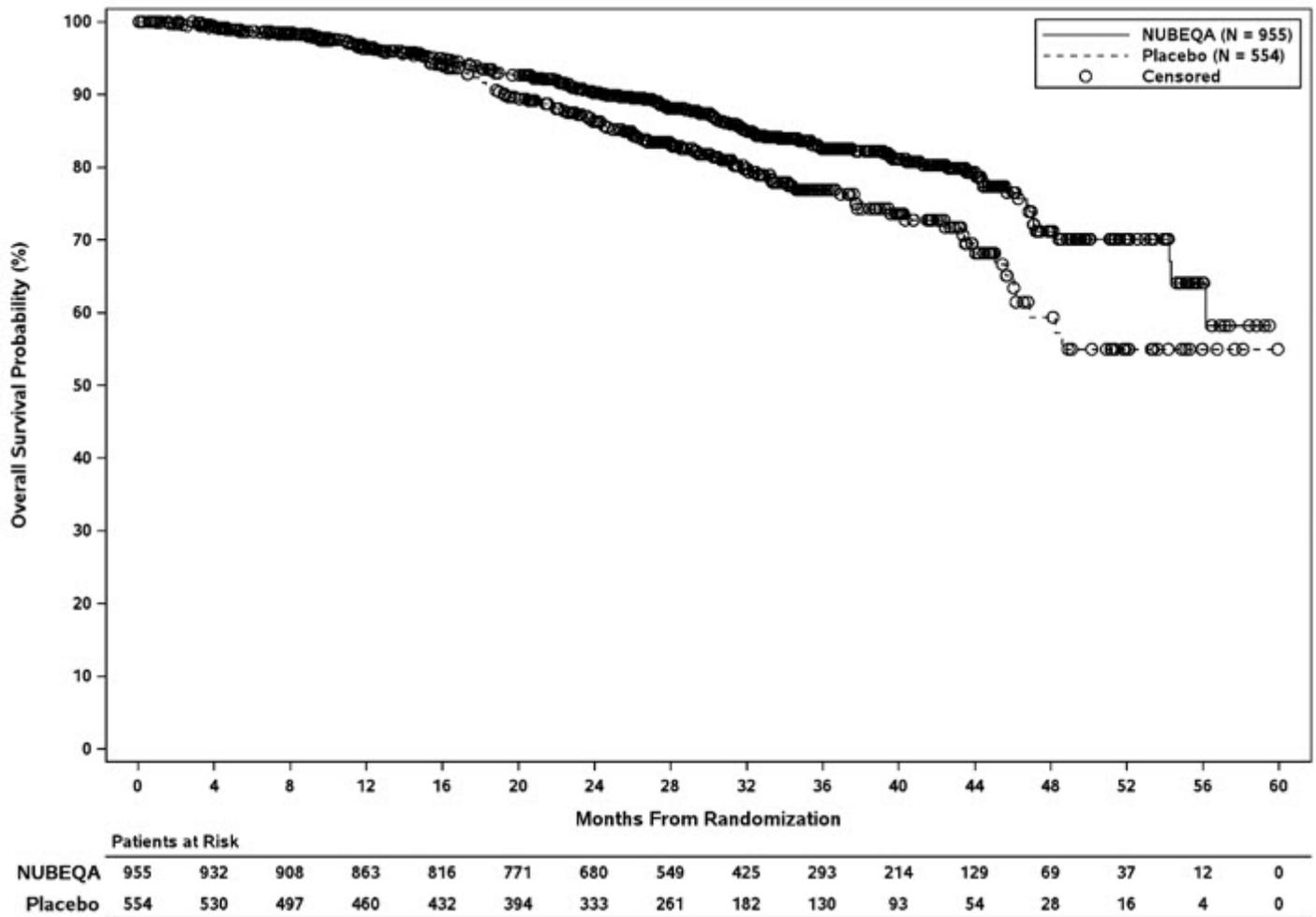


Figure 2: Kaplan-Meier curves of Overall Survival; Intent-To-Treat nmCRPC population (ARAMIS)



MFS results were consistent across patient subgroups for PSADT (≤ 6 months or > 6 months) or prior use of bone-targeting agents (yes or no).

Treatment with NUBEQA resulted in a statistically significant delay in time to pain progression (HR = 0.65, 95% CI = 0.53, 0.79; $p < 0.0001$). Time to pain progression was defined as at least a 2-point worsening from baseline of the pain score on Brief Pain Inventory Short Form or initiation of opioids and reported in 28% of all patients on study.

Treatment with NUBEQA resulted in a statistically significant delay in the initiation of cytotoxic chemotherapy (HR = 0.58, 95% CI = 0.44, 0.76; $p < 0.0001$).

Metastatic Castration Sensitive Prostate Cancer (mCSPC)

ARANOTE

ARANOTE (NCT02799602) was a multicenter, double-blind, placebo-controlled study in 669 patients with mCSPC. Patients were randomized 2:1 to receive 600 mg NUBEQA orally twice daily ($n=446$) or matching placebo ($n=223$). Patients with both high- and low-volume mCSPC were eligible for the study. High-volume disease was defined as presence of visceral metastases, or 4 or more bone lesions, with at least 1 metastasis beyond the vertebral column and pelvic bones. Randomization was stratified by presence of visceral metastasis and use of prior local therapy at study entry. All patients received a gonadotropin-releasing hormone (GnRH) agonist or antagonist concurrently or had a bilateral orchiectomy. Treatment with NUBEQA or placebo continued until progressive disease, change of antineoplastic therapy, unacceptable toxicity, death, or

withdrawal.

The median age was 70 years (range 43-93) and 29% of patients were 75 years of age or older. The racial distribution was 56% White, 31% Asian, and 10% Black; while ethnicity included 24% Hispanic patients. Sixty-eight percent (68%) of patients had a Gleason score of 8 or higher at diagnosis. At study entry, ECOG PS of 0, 1, and 2 were 50%, 47%, and 3%, respectively. Seventy-three percent (73%) of patients had *de novo* disease and 22% had recurrent disease. Seventy-one percent (71%) of patients had high-volume disease, including 12% with visceral metastases, and 29% had low-volume disease. The median PSA level at baseline was 21 µg/L. One patient (0.2%) with a medical history of seizure was treated with NUBEQA in ARANOTE.

The major efficacy outcome measure was radiographic progression-free survival (rPFS), defined as the time from randomization to radiological disease progression or death by central blinded review. Radiographic disease progression was defined by identification of 2 or more new bone lesions on a bone scan with confirmation (Prostate Cancer Working Group 3 criteria) and/or progression in soft tissue disease. Additional efficacy outcome measure was overall survival (OS).

Treatment with NUBEQA resulted in a statistically significant improvement in rPFS compared to placebo. There was no statistically significant improvement in OS.

Efficacy results for ARANOTE are shown in Table 8 and Figure 3.

Table 8: Efficacy Results in Patients with mCSPC in ARANOTE

Efficacy Endpoint	NUBEQA (N=446)	Placebo (N=223)
Radiographic Progression-Free Survival		
Patients with event, n (%)	128 (29)	94 (42)
Median, months (95% CI)	NR (NR, NR)	25 (19, NR)
Hazard ratio (95% CI)*	0.54 (0.41, 0.71)	
p-value†	<0.0001	
Overall Survival		
Patients with event, n (%)	115 (26)	70 (31)
Median, months (95% CI)	NR (NR, NR)	NR (NR, NR)
Hazard Ratio (95% CI)*	0.78 (0.58, 1.05)	
p-value†	NS	

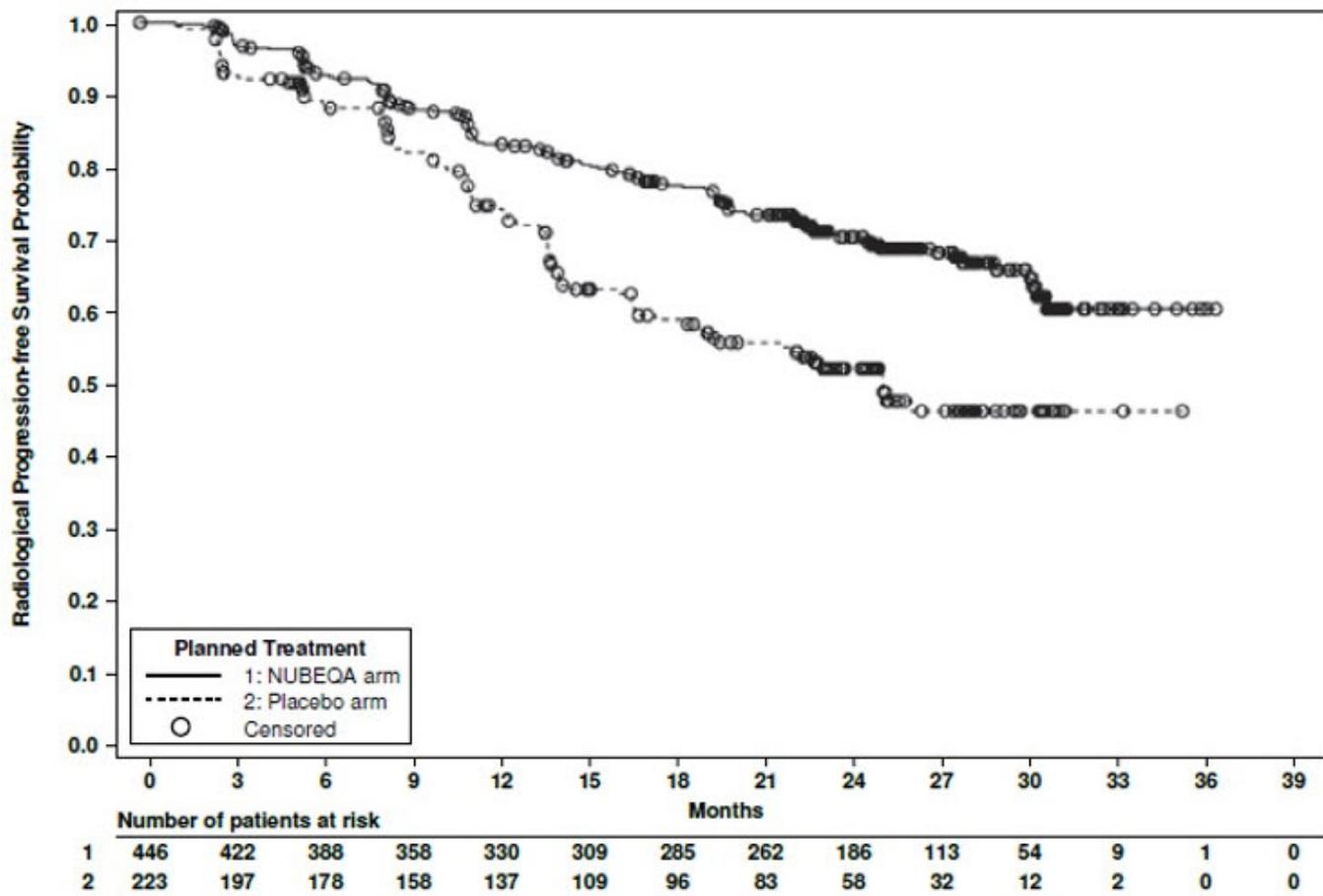
NR: not reached

NS: not statistically significant

* Based on stratified Cox regression model

† Based on stratified log-rank test

Figure 3: Kaplan-Meier Curves for Radiographic Progression-Free Survival; mCSPC population (ARANOTE)



ARASENS

ARASENS (NCT02799602) was a multicenter, double-blind, placebo-controlled clinical trial in 1306 patients with mCSPC. Patients were randomized 1:1 to receive 600 mg NUBEQA orally twice daily (n=651) or matching placebo (n=655), concomitantly with 75 mg/m² of docetaxel for 6 cycles. All patients received a gonadotropin-releasing hormone (GnRH) agonist or antagonist concurrently or had a bilateral orchiectomy. Treatment with NUBEQA or placebo continued until symptomatic progressive disease, change of antineoplastic therapy, or unacceptable toxicity. Patients with regional lymph node involvement only (M0) were excluded from the study. Patients were stratified by extent of disease (non-regional lymph nodes metastases only (M1a), bone metastases with or without lymph node metastases (M1b) or visceral metastases with or without lymph node metastases or with or without bone metastases (M1c)) and by alkaline phosphatase level (< or ≥ upper limit of normal) at study entry.

The median age was 67 years (range 41–89) and 17% of patients were 75 years of age or older. The racial distribution included 52% White, 36% Asian, 4% Black, while ethnicity included 7% Hispanic patients. Seventy-eight percent (78%) of patients had a Gleason score of ≥8 at diagnosis. Seventy one percent (71%) of patients had an ECOG performance status score of 0 and 29% patients had an ECOG performance status score of 1. There were 86% of patients with *de novo* and 13% with recurrent disease. At initial diagnosis of metastatic disease, 3% had M1a, 83% had M1b and 14% had M1c disease; alkaline phosphatase was < ULN in 45% of patients and ≥ ULN in 55% of patients; median PSA level at baseline was 30 µg/L and 24 µg/L for NUBEQA vs placebo

group, respectively. Four patients (0.6%) with a medical history of seizure were treated with NUBEQA with docetaxel in ARASENS.

The major efficacy outcome measure was overall survival (OS). Time to pain progression was an additional efficacy outcome measure.

Treatment with NUBEQA with docetaxel resulted in a statistically significant improvement in OS compared to placebo with docetaxel. OS results were consistent across stratified subgroups (extent of disease and alkaline phosphatase level).

Efficacy results are shown in Table 9 and Figure 4.

Table 9: Efficacy Results from the ARASENS study

	NUBEQA with docetaxel (N=651)	Placebo with docetaxel (N=654)*
Overall survival		
Deaths (%)	229 (35)	304 (46)
Median in months (95% CI)	NR (NR, NR)	48.9 (44.4, NR)
Hazard Ratio (95% CI) [†]	0.68 (0.57, 0.80)	
P-value [‡]	<0.0001	

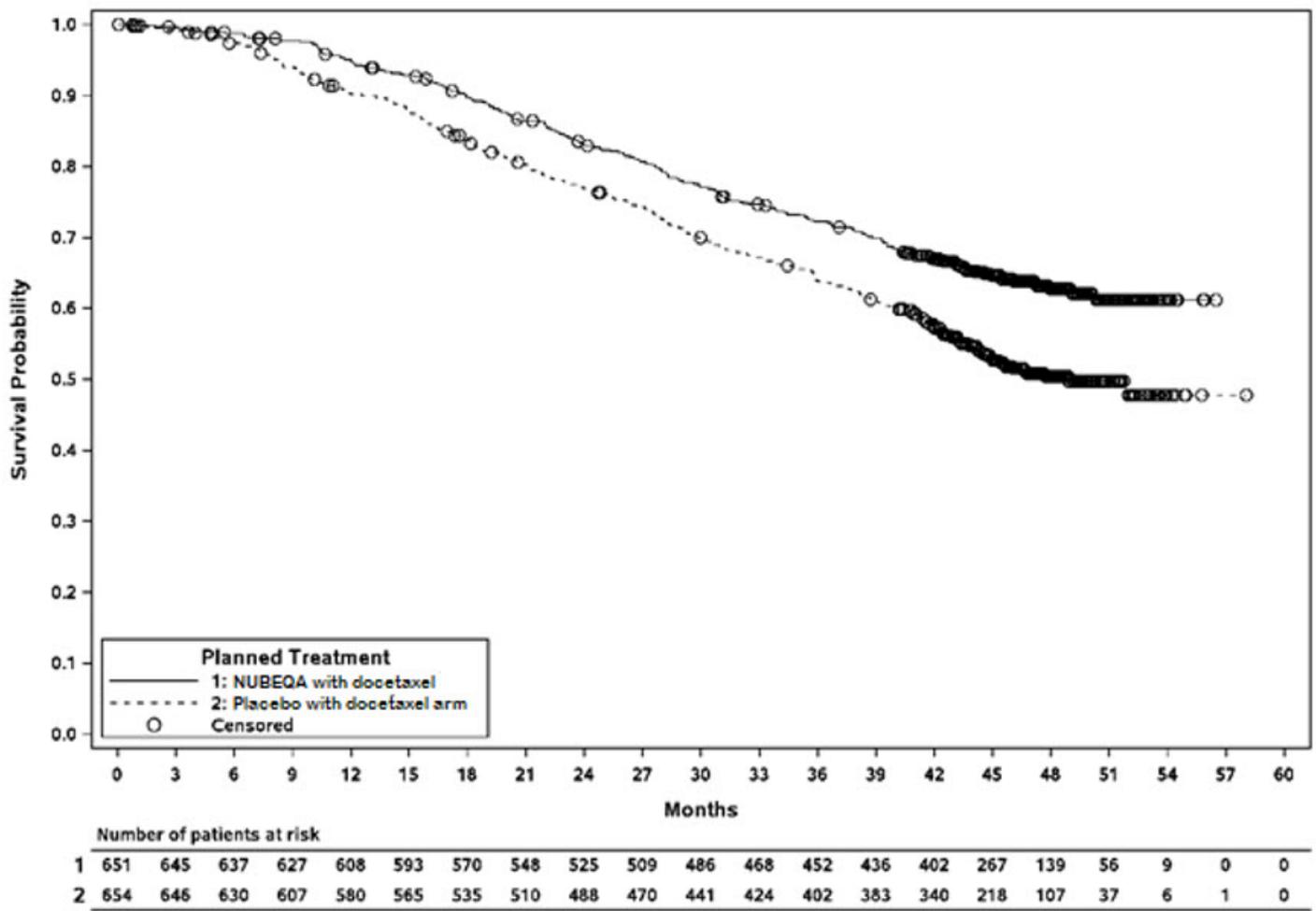
NR: not reached

* One patient in the placebo arm was excluded from all analyses due to the violation of Good Clinical Practices

† Hazard ratio < 1 favors NUBEQA

‡ P-value is one-sided, and based on a log-rank test stratified by extent of disease (non-regional lymph nodes metastases only or bone metastases with or without lymph node metastases or visceral metastases with or without lymph node metastases or with or without bone metastases) and Alkaline Phosphatase (ALP < ULN or ALP ≥ ULN)

Figure 4: Kaplan-Meier curves of Overall Survival; mCSPC population (ARASENS)



Treatment with NUBEQA with docetaxel resulted in a statistically significant delay in time to pain progression (HR = 0.79, 95% CI= 0.66, 0.95; 1-sided p value = 0.006). Time to pain progression was defined as the time from randomization to the time of pain progression. Pain progression is defined as:

- An increase of 2 or more points in the "worst pain in 24 hours" score (WPS) from nadir observed at 2 consecutive evaluations at least 4 weeks apart, or initiation of short- or long-acting opioid use for pain for at least 7 consecutive days, for asymptomatic patients (WPS=0) at baseline
- An increase of 2 or more points in the "worst pain in 24 hours" score (WPS) from nadir observed at 2 consecutive evaluations at least 4 weeks apart, and a WPS of 4 or greater, or initiation of short- or long-acting opioid use for pain for at least 7 consecutive days, for symptomatic patients (WPS ≥ 1) at baseline

Patients with opioid use within 4 weeks before randomization were censored at randomization in this analysis (125 patients (19%) in the NUBEQA with docetaxel arm and 118 patients (18%) in the placebo with docetaxel arm).

16 HOW SUPPLIED/STORAGE AND HANDLING

How Supplied

NUBEQA (darolutamide) 300 mg film-coated tablets are white to off-white, oval shaped tablets, marked with "300" on one side, and "BAYER" on the other side. NUBEQA 300 mg tablets are available in bottles of 120 tablets.

NDC 50419-395-01

Storage and Handling

Store at 20°C to 25°C (68°F to 77°F); excursions permitted between 15°C and 30°C (59°F and 86°F) [see USP Controlled Room Temperature].

Keep the bottle tightly closed after first opening.

17 PATIENT COUNSELING INFORMATION

Advise the patient to read the FDA-approved patient labeling (Patient Information)

Ischemic Heart Disease

Inform patients that NUBEQA has been associated with an increased risk of ischemic heart disease. Advise patients to seek immediate medical attention if any symptoms suggestive of an ischemic heart disease event occur [see *Warnings and Precautions (5.1)*].

Seizure

Inform patients that NUBEQA has been associated with an increased risk of seizure. Discuss conditions that may predispose to seizures and medications that may lower the seizure threshold. Advise patients of the risk of engaging in any activity where sudden loss of consciousness could cause serious harm to themselves or others. Inform patients to contact their healthcare provider right away if they have loss of consciousness or seizure [see *Warnings and Precautions (5.2)*].

Embryo-Fetal Toxicity

Inform patients that NUBEQA can be harmful to a developing fetus and can cause loss of pregnancy [see *Use in Specific Populations (8.1)*].

Advise male patients with female partners of reproductive potential to use effective contraception during treatment and for 1 week after the last dose of NUBEQA [see *Warnings and Precautions (5.3) and Use in Specific Populations (8.1, 8.3)*].

Dosage and Administration

Inform patients receiving concomitant gonadotropin-releasing hormone (GnRH) agonist or antagonist therapy that they need to maintain this treatment during the course of treatment with NUBEQA.

Instruct patients to take their dose of two tablets (twice daily). NUBEQA should be taken with food. Each tablet should be swallowed whole.

Inform patients that in the event of a missed daily dose of NUBEQA, to take any missed dose, as soon as they remember. A missed dose may be taken together with the next scheduled dose. Thereafter, resume the regularly scheduled twice daily dosing [see *Dosage and Administration (2.1)*].

Infertility

Advise male patients that NUBEQA may impair fertility [see *Use in Specific Populations (8.3)*].

Manufactured by: Orion Corporation, Orion Pharma, FI-02101 Espoo, Finland

Manufactured for: Bayer HealthCare Pharmaceuticals Inc., Whippany, NJ 07981 USA

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For more information, call Bayer HealthCare Pharmaceuticals Inc. at Bayer at 1-888-842-2937 or go to www.NUBEQA-us.com

PATIENT INFORMATION

NUBEQA® (NOO-bə-kə) (darolutamide) tablets

What is NUBEQA?

NUBEQA is a prescription medicine used to treat adults with prostate cancer:

- that has not spread to other parts of the body and no longer responds to a medical or surgical treatment that lowers testosterone
- that has spread to other parts of the body and responds to a medical or surgical treatment that lowers testosterone
- that has spread to other parts of the body and responds to a medical or surgical treatment that lowers testosterone in combination with docetaxel.

It is not known if NUBEQA is safe and effective in females or children.

Before taking NUBEQA, tell your healthcare provider about all your medical conditions, including if you:

- have heart problems
- have high blood pressure
- have diabetes
- have high amounts of fat or cholesterol in your blood (dyslipidemia)
- have a history of seizures, brain injury, stroke, or brain tumors
- have kidney or liver problems
- are pregnant or plan to become pregnant. NUBEQA can cause harm to your unborn baby and loss of pregnancy (miscarriage).
- have a partner who may become pregnant. Males who have female partners who may become pregnant should use effective birth control (contraception) during treatment and for 1 week after the last dose of NUBEQA. Talk with your healthcare provider about birth control methods.
- are breastfeeding or plan to breastfeed. It is not known if NUBEQA passes into breast milk.

Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements. NUBEQA may affect the way other medicines work and other medicines may affect how NUBEQA works.

You should not start or stop any medicine before you talk with the healthcare provider that prescribed NUBEQA.

Know the medicines you take. Keep a list of them with you to show to your healthcare provider and pharmacist when you get a new medicine.

How should I take NUBEQA?

- Take NUBEQA exactly as your healthcare provider tells you.
- Your healthcare provider may change your dose, temporarily stop, or completely stop your treatment with NUBEQA if you have certain side effects.

- Take your prescribed dose of NUBEQA 2 times a day with food.
- Swallow NUBEQA tablets whole.
- If you are receiving gonadotropin-releasing hormone (GnRH) therapy, you should continue with this treatment during your treatment with NUBEQA unless you have had a surgery to lower the amount of testosterone in your body (surgical castration).
- If you miss a dose of NUBEQA, take your prescribed dose as soon as you remember. A missed dose may be taken together with the next scheduled dose. Thereafter, resume the regularly scheduled twice daily dosing.
- If you take more NUBEQA than prescribed, call your healthcare provider right away.

What are the possible side effects of NUBEQA?

NUBEQA may cause serious side effects, including:

- **Heart disease.** Blockage of the arteries in the heart (ischemic heart disease) that can lead to death has happened in some people during treatment with NUBEQA. Your healthcare provider will monitor you for signs and symptoms of heart problems during your treatment with NUBEQA. Call your healthcare provider or get medical help right away if you get chest pain or discomfort at rest or with activity, or shortness of breath during your treatment with NUBEQA.
- **Seizure.** Treatment with NUBEQA may increase your risk of having a seizure. You should avoid activities where a sudden loss of consciousness could cause serious harm to yourself or others. Tell your healthcare provider right away if you have loss of consciousness or seizure during treatment with NUBEQA.

The most common side effects of NUBEQA include:

- increase in liver function tests
- decreased white blood cells (neutropenia)
- feeling more tired than usual

The most common side effects of NUBEQA in combination with docetaxel include:

- | | |
|-----------------------|--------------------------------------|
| • constipation | • decreased red blood cells (anemia) |
| • rash | • high blood sugar levels |
| • decreased appetite | • decreased white blood cells |
| • bleeding | • increase in liver function tests |
| • weight gain | • low blood calcium levels |
| • high blood pressure | |

NUBEQA may cause fertility problems in males, which may affect the ability to father children. Talk to your healthcare provider if you have concerns about fertility.

These are not all the possible side effects of NUBEQA.

Call your healthcare provider for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

How should I store NUBEQA?

- Store NUBEQA at room temperature between 68°F to 77°F (20°C to 25°C).
- Keep the bottle tightly closed after you first open it.

Keep NUBEQA and all medicines out of the reach of children.

General information about the safe and effective use of NUBEQA.

Medicines are sometimes prescribed for purposes other than those listed in a Patient Information leaflet. Do not use NUBEQA for a condition for which it was not prescribed. Do not give NUBEQA to other people, even if they have the same symptoms that you have. It may harm them. You can ask your healthcare provider or pharmacist for information about NUBEQA that is written for health professionals.

What are the ingredients in NUBEQA?

Active ingredient: darolutamide

Inactive ingredients: calcium hydrogen phosphate, croscarmellose sodium, lactose monohydrate, magnesium stearate, povidone K 30, hypromellose 15 cP, macrogol 3350, and titanium dioxide.

Manufactured by: Orion Corporation, Orion Pharma, FI-02101 Espoo, Finland

Manufactured for: Bayer HealthCare Pharmaceuticals Inc., Whippany, NJ 07981 USA

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For more information, call Bayer HealthCare Pharmaceuticals Inc. at Bayer at 1-888-842-2937 or go to www.NUBEQA-us.com

This Patient Information has been approved by the U.S. Food and Drug Administration.

Revised: X/202X

PACKAGE/LABEL PRINCIPAL DISPLAY PANEL

50419-395-01

Rx Only

NUBEQA

(darolutamide) tablets

300 mg

120 film-coated tablets



NUBEQA

darolutamide tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:50419-395
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DAROLUTAMIDE (UNII: X05U0N2RCO) (DAROLUTAMIDE - UNII:X05U0N2RCO)	DAROLUTAMIDE	300 mg

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS DIBASIC CALCIUM PHOSPHATE (UNII: L11K75P92J)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POVIDONE K30 (UNII: U725QWY32X)	
POLYETHYLENE GLYCOL 3350 (UNII: G2M7P15E5P)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	WHITE (to off white)	Score	no score
Shape	OVAL	Size	16mm
Flavor		Imprint Code	300;Bayer
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50419-395-01	1 in 1 CARTON	07/31/2019	
1		120 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:50419-395-72	1 in 1 CARTON	03/03/2021	
2		120 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA212099	07/31/2019	

Labeler - Bayer HealthCare Pharmaceuticals Inc. (005436809)**Establishment**

Name	Address	ID/FEI	Business Operations
Orion Corporation, Orion Pharma		539763727	MANUFACTURE(50419-395) , ANALYSIS(50419-395)

Establishment

Name	Address	ID/FEI	Business Operations
Sharp Corporation		143696495	PACK(50419-395) , LABEL(50419-395)

Establishment

Name	Address	ID/FEI	Business Operations
Orion Corporation, Orion Pharma		537940319	ANALYSIS(50419-395)

Establishment

Name	Address	ID/FEI	Business Operations
Orion Corporation, Orion Pharma		368380223	LABEL(50419-395) , PACK(50419-395)

Establishment

Name	Address	ID/FEI	Business Operations
Fermion Oy		537993060	API MANUFACTURE(50419-395) , ANALYSIS(50419-395)

Establishment

Name	Address	ID/FEI	Business Operations
Quinta-Analytica s.r.o.		495722555	ANALYSIS(50419-395)

Establishment

Name	Address	ID/FEI	Business Operations
Fermion Oy		537994803	ANALYSIS(50419-395)

Revised: 3/2026

Bayer HealthCare Pharmaceuticals Inc.