Ordering Information

ELAHERE®
mirvetuximab soravtansine-gynx
injection 100 mg

100 mg/20 mL (5 mg/mL) in a single-dose vial
NDC: 72903-853-01
Products shown are not actual size.

SPECIALTY DISTRIBUTORS

AmerisourceBergen ASD Healthcare
800-746-6273
800-547-9413
www.asdhealthcare.com
8 AM–7:30 PM ET, Monday–Thursday
8 AM–7 PM ET, Friday

Cardinal Health Specialty Pharmaceutical Distribution
For Hospital
855-855-0708
614-553-6301
GMB-SPD-CSORDERENTRY@cardinalhealth.com

For Physician Office
877-453-3972
877-274-9897
GMB-SPDOncologySalesTeam@cardinalhealth.com

Ordering Portal
specialtyonline.cardinalhealth.com
orderexpress.cardinalhealth.com

Biologics by McKesson
800-850-4306
800-823-4506
biologics.mckesson.com
8 AM–8 PM ET 7-days a week.
Registered pharmacists available 24/7.

AmerisourceBergen Oncology Supply
800-633-7555
800-248-8205
www.oncologysupply.com
7 AM–11 PM ET, Monday–Friday,
8:30 AM–8 PM ET, Saturday–Sunday

McKesson Plasma and Biologics (Hospitals, IDNs, VA)
877-625-2566
888-752-7626
MPBOrders@mckesson.com
9 AM–7:30 PM ET, Monday–Friday

McKesson Specialty Health (MD Offices)
800-482-6700
800-289-9285
mscs.mckesson.com
8 AM–8 PM ET, Monday–Friday

Axium Healthcare Puerto Rico (for Puerto Rico ONLY)
844-355-4191
800-546-2163
www.axiumpr.com
8 AM–6 PM AT, Monday–Friday

INDICATION[1]

ELAHERE is indicated for the treatment of adult patients with folate receptor-alpha (FRα) positive, platinum-resistant epithelial ovarian, fallopian tube, or primary peritoneal cancer, who have received one to three prior systemic treatment regimens. Select patients for therapy based on an FDA-approved test.

IMPORTANT SAFETY INFORMATION[1]

WARNING: OCULAR TOXICITY

- **ELAHERE can cause severe ocular toxicities, including visual impairment, keratopathy, dry eye, photophobia, eye pain, and uveitis.**
- **Conduct an ophthalmic exam including visual acuity and slit lamp exam prior to initiation of ELAHERE, every other cycle for the first 8 cycles, and as clinically indicated.**
- **Administer prophylactic artificial tears and ophthalmic topical steroids.**
- **Withhold ELAHERE for ocular toxicities until improvement and resume at the same or reduced dose.**
- **Discontinue ELAHERE for Grade 4 ocular toxicities.**

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Product returns
Credit for returns is subject to the AbbVie Return Goods Policy. For more information, please contact your Specialty Distributor.

IDN=integrated delivery networks; NDC=National Drug Code; VA=US Department of Veterans Affairs.

Please see full Prescribing Information, including Boxed Warning.

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ELAHERE Support Services (ESS) is committed to helping appropriate patients **start on ELAHERE** by offering **access** and **reimbursement** support, **affordability** assistance, and dedicated **nursing support** for patient questions based on the Prescribing Information.

ENROLL YOUR PATIENT IN ELAHERE SUPPORT SERVICES

Visit ELAHEREhcp.com to download and complete the enrollment form.

WHAT ELAHERE SUPPORT SERVICES OFFERS

Once enrolled, ESS offers the following services and programs for patients:

Access & reimbursement

- Benefits investigation
- Prior authorization assistance
- Appeals assistance

Copay assistance*

- Support for commercially eligible patients with out-of-pocket costs
- Patients could **pay as little as $0** for their medication

Patient assistance

- Support for uninsured or underinsured patients who meet eligibility requirements to access medication at no charge†

Nurse Navigators

- A resource available to patients and their caregivers to answer questions about their treatment based on the Prescribing Information

*Terms and conditions apply. Patients are eligible for copay assistance if enrolled in private commercial health insurance and are not covered by state or federal healthcare programs, and who meet the eligibility criteria. Patients will be enrolled for 12 months. There are no income requirements to participate in the program.
†Criteria include patients who are uninsured or have insurance that excludes coverage for ELAHERE (including patients on Medicare or Medicaid), residents of the United States or Puerto Rico, and patients who meet the financial eligibility requirements. Terms and conditions apply.

GET IN TOUCH WITH ELAHERE SUPPORT SERVICES

For questions, connect with an ELAHERE Support Services Program specialist by calling 1-833–ELAHERE (1-833-352-4373) Monday to Friday, 8:00 AM to 8:00 PM ET or email to ELAHERESupport@cardinalhealth.com

Reference: 1. ELAHERE. Package insert. AbbVie, Inc.; 2024.

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IMPORTANT SAFETY INFORMATION[1] (CONTINUED)

WARNINGS and PRECAUTIONS

Ocular Disorders

ELAHERE can cause severe ocular adverse reactions, including visual impairment, keratopathy (corneal disorders), dry eye, photophobia, eye pain, and uveitis.

Ocular adverse reactions occurred in 59% of patients with ovarian cancer treated with ELAHERE. Eleven percent (11%) of patients experienced Grade 3 ocular adverse reactions, including blurred vision, keratopathy (corneal disorders), dry eye, cataract, photophobia, and eye pain; two patients (0.3%) experienced Grade 4 events (keratopathy and cataract). The most common (≥5%) ocular adverse reactions were blurred vision (48%), keratopathy (36%), dry eye (27%), cataract (16%), photophobia (14%), and eye pain (10%).

The median time to onset for first ocular adverse reaction was 5.1 weeks (range: 0.1 to 68.6). Of the patients who experienced ocular events, 53% had complete resolution; 38% had partial improvement (defined as a decrease in severity by one or more grades from the worst grade at last follow up). Ocular adverse reactions led to permanent discontinuation of ELAHERE in 1% of patients.

Premedication and use of lubricating and ophthalmic topical steroid eye drops during treatment with ELAHERE are recommended. Advise patients to avoid use of contact lenses during treatment with ELAHERE unless directed by a healthcare provider.

Refer patients to an eye care professional for an ophthalmic exam including visual acuity and slit lamp exam prior to treatment initiation, every other cycle for the first 8 cycles, and as clinically indicated. Promptly refer patients to an eye care professional for any new or worsening ocular signs and symptoms.

Monitor for ocular toxicity and withhold, reduce, or permanently discontinue ELAHERE based on severity and persistence of ocular adverse reactions.

Pneumonitis

Severe, life-threatening, or fatal interstitial lung disease (ILD), including pneumonitis, can occur in patients treated with ELAHERE.

Pneumonitis occurred in 10% of patients treated with ELAHERE, including 1% with Grade 3 events and 1 patient (0.1%) with a Grade 4 event. One patient (0.1%) died due to respiratory failure in the setting of pneumonitis and lung metastases. One patient (0.1%) died due to respiratory failure of unknown etiology. Pneumonitis led to permanent discontinuation of ELAHERE in 3% of patients.

Monitor patients for pulmonary signs and symptoms of pneumonitis, which may include hypoxia, cough, dyspnea, or interstitial infiltrates on radiologic exams. Infectious, neoplastic, and other causes for such symptoms should be excluded through appropriate investigations. Withhold ELAHERE for patients who develop persistent or recurrent Grade 2 pneumonitis until symptoms resolve to ≤ Grade 1 and consider dose reduction. Permanently discontinue ELAHERE in all patients with Grade 3 or 4 pneumonitis. Patients who are asymptomatic may continue dosing of ELAHERE with close monitoring.

Peripheral Neuropathy (PN)

Peripheral neuropathy occurred in 36% of patients with ovarian cancer treated with ELAHERE across clinical trials; 3% of patients experienced Grade 3 peripheral neuropathy. Peripheral neuropathy adverse reactions included peripheral neuropathy (20%), peripheral sensory neuropathy (9%), paraesthesia (6%), neurotoxicity (3%), hypoaesthesia (1%), peripheral motor neuropathy (0.9%), polyneuropathy (0.3%), and peripheral sensorimotor neuropathy (0.1%). Monitor patients for signs and symptoms of neuropathy, such as paresthesia, tingling or a burning sensation, neuropathic pain, muscle weakness, or dysesthesia. For patients experiencing new or worsening PN, withhold dosage, dose reduce, or permanently discontinue ELAHERE based on the severity of PN.

Embryo-Fetal Toxicity

Based on its mechanism of action, ELAHERE can cause embryo-fetal harm when administered to a pregnant woman because it contains a genotoxic compound (DM4) and affects actively dividing cells.

Advise pregnant women of the potential risk to a fetus. Advise females of reproductive potential to use effective contraception during treatment with ELAHERE and for 7 months after the last dose.

ADVERSE REACTIONS

The most common (≥20%) adverse reactions, including lab abnormalities, were increased aspartate aminotransferase, fatigue, increased alanine aminotransferase, blurred vision, nausea, increased alkaline phosphatase, diarrhea, abdominal pain, keratopathy, peripheral neuropathy, musculoskeletal pain, decreased lymphocytes, decreased platelets, decreased magnesium, decreased hemoglobin, dry eye, constipation, decreased leukocytes, vomiting, decreased albumin, decreased appetite, and decreased neutrophils.

DRUG INTERACTIONS

DM4 is a CYP3A4 substrate. Closely monitor patients for adverse reactions with ELAHERE when used concomitantly with strong CYP3A4 inhibitors.

USE IN SPECIFIC POPULATIONS

Lactation

Advise women not to breastfeed during treatment with ELAHERE and for 1 month after the last dose.

Hepatic Impairment

Avoid use of ELAHERE in patients with moderate or severe hepatic impairment (total bilirubin >1.5 ULN).

Please see Full Prescribing Information, including Boxed WARNING.

References: 1. ELAHERE. Package insert. AbbVie, Inc.; 2024.

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