

WINREVAIR- sotatercept-csrk

Merck Sharp & Dohme LLC

HIGHLIGHTS OF PRESCRIBING INFORMATION

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

WINREVAIR™ (sotatercept-csrk) for injection, for subcutaneous use
Initial U.S. Approval: 2024

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

Indications and Usage (1)

10/2025

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

WINREVAIR is an activin signaling inhibitor indicated for the treatment of adults with pulmonary arterial hypertension (PAH, WHO Group 1 pulmonary hypertension) to improve exercise capacity and WHO functional class (FC), and reduce the risk of clinical worsening events, including hospitalization for PAH, lung transplantation and death. (1)

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

- The recommended starting dose is 0.3 mg/kg by subcutaneous injection. (2.1)
- The recommended target dose is 0.7 mg/kg every 3 weeks by subcutaneous injection. (2.2)
- Dosage modifications due to increased hemoglobin (Hgb) and decreased platelets may be necessary. Check Hgb and platelets before each dose for the first 5 doses, or longer if values are unstable, and monitor periodically thereafter. (2.3)
- See full prescribing information for preparation and administration instructions. (2.4)

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

- For injection: 45 mg lyophilized cake or powder in a single-dose vial (3)
- For injection: 60 mg lyophilized cake or powder in a single-dose vial (3)

-----CONTRAINDICATIONS-----

None (4)

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

- Erythrocytosis: If severe, may increase the risk of thromboembolic events and hyperviscosity syndrome. Monitor Hgb before each dose for the first 5 doses, or longer if values are unstable, and periodically thereafter to determine if dose adjustments are required. (5.1)
- Severe Thrombocytopenia: May increase the risk of bleeding. Monitor platelets before each dose for the first 5 doses, or longer if values are unstable, and periodically thereafter to determine if dose adjustments are required. (5.2)
- Serious Bleeding: Serious bleeding events were reported and were more likely with concomitant prostacyclin and/or antithrombotic agents, or with low platelet counts. Do not administer WINREVAIR if the patient is experiencing serious bleeding. (5.3)
- Embryo-Fetal Toxicity: May cause fetal harm. Advise females of reproductive potential of the potential risk to a fetus and use of effective contraception. (5.4, 8.1, 8.3)
- Impaired Fertility: May impair female and male fertility. (5.5, 8.3, 13.1)

-----ADVERSE REACTIONS-----

The most common ($\geq 10\%$ in patients receiving WINREVAIR and 5% more than placebo) adverse reactions were infections, epistaxis, telangiectasia, diarrhea, headache, rash, increased hemoglobin, dizziness, erythema, and gingival bleeding. (6.1)

To report SUSPECTED ADVERSE REACTIONS, contact Merck Sharp & Dohme LLC at 1-877-888-4231 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch

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

Lactation: Breastfeeding not recommended. (8.2)

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

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FULL PRESCRIBING INFORMATION

1 INDICATIONS AND USAGE

WINREVAIR™ is indicated for the treatment of adults with pulmonary arterial hypertension (PAH, Group 1 pulmonary hypertension) to improve exercise capacity and World Health Organization (WHO) functional class (FC), and reduce the risk of clinical worsening events including hospitalization for PAH, lung transplantation and death [see *Clinical Studies (14.1)*].

2 DOSAGE AND ADMINISTRATION

2.1 Recommended Starting Dosage

WINREVAIR is administered once every 3 weeks by subcutaneous injection according to patient body weight. The starting dose of WINREVAIR is 0.3 mg/kg.

Obtain hemoglobin (Hgb) and platelet count prior to the first dose of WINREVAIR. Do not initiate treatment if platelet count is $<50,000/\text{mm}^3$ ($<50 \times 10^9/\text{L}$) [see *Dosage and Administration (2.3)*].

Injection volume for starting dose is calculated based on patient weight as follows:

$$\text{Injection Volume (mL)} = \frac{\text{Weight (kg)} \times 0.3 \text{ mg/kg}}{50 \text{ mg/mL}}$$

Injection volume should be rounded to the nearest 0.1 mL.

For example: $(70 \text{ kg} \times 0.3 \text{ mg/kg}) \div 50 \text{ mg/mL} = 0.42 \text{ mL}$, rounds to 0.4 mL

See Table 1 for selecting the appropriate kit based on calculated injection volume for starting dose.

Table 1: Kit Type Based on Injection Volume for Dose of 0.3 mg/kg

Injection Volume (mL)	Kit Type
0.2 to 0.9	45 mg kit (containing 1 x 45 mg vial)
1 to 1.1	60 mg kit (containing 1 x 60 mg vial)

2.2 Recommended Target Dosage

After verifying acceptable Hgb and platelet count, increase to the target dose of 0.7 mg/kg. Continue treatment at 0.7 mg/kg every 3 weeks unless dosage adjustments are required [see *Dosage and Administration (2.3)*].

Injection volume for target dose is calculated based on patient weight as follows:

$$\text{Weight (kg)} \times 0.7 \text{ mg/kg}$$

$$\text{Injection Volume (mL)} = \frac{\text{Weight (kg)} \times 0.7 \text{ mg/kg}}{50 \text{ mg/mL}}$$

Injection volume should be rounded to the nearest 0.1 mL.

For example: (70 kg x 0.7 mg/kg) ÷ 50 mg/mL = 0.98 mL, rounds to 1 mL

See Table 2 for selecting the appropriate kit based on calculated injection volume for target dose.

Table 2: Kit Type Based on Injection Volume for Dose of 0.7 mg/kg

Injection Volume (mL)	Kit Type
0.4 to 0.9	45 mg kit (containing 1 x 45 mg vial)
1 to 1.2	60 mg kit (containing 1 x 60 mg vial)
1.3 to 1.8	90 mg kit (containing 2 x 45 mg vials)
1.9 to 2.4	120 mg kit (containing 2 x 60 mg vials)

Missed Dose, Overdose, and Underdose

If a dose of WINREVAIR is missed, administer as soon as possible. If the missed dose of WINREVAIR is not administered within 3 days of the scheduled date, adjust the schedule to maintain 3-week dosing intervals. In case of an overdose, monitor for erythrocytosis [see *Overdosage (10)*].

2.3 Dosage Modifications Due to Hemoglobin Increase or Platelet Count Decrease

Check Hgb and platelet count before each dose for the first 5 doses, or longer if values are unstable. Thereafter, monitor Hgb and platelet count periodically [see *Warnings and Precautions (5.1, 5.2)*].

Delay treatment for at least 3 weeks if any of the following occur:

- Hgb increases >2.0 g/dL from the previous dose and is above ULN.
- Hgb increases >4.0 g/dL from baseline.
- Hgb increases >2.0 g/dL above ULN.
- Platelet count decreases to <50,000/mm³ (<50 x 10⁹/L).

Recheck Hgb and platelet count before reinitiating treatment. For treatment delays lasting >9 weeks, restart treatment at 0.3 mg/kg, and escalate to 0.7 mg/kg after verifying acceptable Hgb and platelet count.

2.4 Preparation and Administration

Administration is subject to monitoring of hemoglobin and platelet count [see *Dosage and Administration (2.3)*, *Warnings and Precautions (5.1, 5.2)*].

WINREVAIR is intended for use under the guidance of a healthcare professional. Patients and caregivers may administer WINREVAIR when considered appropriate and when they receive training and follow-up from the healthcare provider (HCP) on how to

reconstitute, prepare, measure, and inject WINREVAIR [see *Patient Counseling Information (17)*].

Confirm at subsequent visits that the patient and/or caregiver can correctly prepare and administer WINREVAIR, particularly if the dose changes or the patient requires a different kit [see *Warnings and Precautions (5.1)*].

Refer to the Instructions for Use (IFU) for detailed instructions on the proper preparation and administration of WINREVAIR.

Selecting the Appropriate Product Kit

If a patient's body weight requires the use of two 45 mg vials or two 60 mg vials of lyophilized product, use a 2-vial kit instead of two individual 1-vial kits. A 2-vial kit includes instructions to combine the contents of two vials, which aids in measuring the proper dosage and eliminates the need for multiple injections [see *How Supplied/Storage and Handling (16.1)*].

Reconstitution Instructions

- Remove the injection kit from the refrigerator and wait 15 minutes to allow the prefilled syringe(s) and drug product to come to room temperature prior to preparation.
- Attach the vial adapter to the vial.
- Visually inspect the pre-filled syringe for any damage or leaks and the Sterile Water for Injection inside to ensure there are no visible particles.
- Snap off the cap of the pre-filled syringe and attach the syringe to the vial adapter.
- Inject all of the Sterile Water for Injection from the attached syringe into the vial containing the lyophilized powder. This will provide a final concentration of 50 mg/mL.
- Gently swirl the vial to reconstitute the drug product. DO NOT shake or vigorously agitate.
- Allow the vial to stand for up to 3 minutes to allow bubbles to disappear.
- Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permit.
- When properly mixed, WINREVAIR should be clear to opalescent and colorless to slightly brownish-yellow and does not have clumps or powder.
- If prescribed a 2-vial presentation, repeat the steps within this section to prepare the second vial.
- Use the reconstituted solution as soon as possible, but no later than 4 hours after reconstitution. Discard unused reconstituted solution.

Syringe Preparation

- Turn the syringe and vial upside-down and withdraw the appropriate volume for injection, based on the patient's weight.
 - If the dose amount requires the use of two vials, withdraw the entire contents of the first vial and slowly transfer full contents into the second vial.
 - Turn the syringe and vial upside-down and withdraw the required amount of drug product.
 - If necessary, remove excess drug product.
- If necessary, remove excess air from the syringe.

Administration Instructions

WINREVAIR is for subcutaneous injection.

- Select the injection site on the abdomen (at least 2 inches away from navel), upper thigh, or upper arm, and swab with an alcohol wipe. Select a new site for each injection that is not scarred, tender, or bruised.
 - For administration by the patient or caregiver, use only the abdomen and upper thigh (see IFU).
- Perform subcutaneous injection.

3 DOSAGE FORMS AND STRENGTHS

- For injection: 45 mg white to off-white lyophilized cake or powder appearance in a single-dose vial.
- For injection: 60 mg white to off-white lyophilized cake or powder appearance in a single-dose vial.

4 CONTRAINDICATIONS

None.

5 WARNINGS AND PRECAUTIONS

5.1 Erythrocytosis

WINREVAIR may increase hemoglobin. Severe erythrocytosis may increase the risk of thromboembolic events or hyperviscosity syndrome. In clinical studies, moderate elevations in Hgb (>2 g/dL above ULN) occurred in 15% of patients taking WINREVAIR while no elevations ≥ 4 g/dL above ULN were observed. Monitor Hgb before each dose for the first 5 doses, or longer if values are unstable, and periodically thereafter, to determine if dose adjustments are required [see *Dosage and Administration (2.3)*, *Adverse Reactions (6.1)*].

5.2 Severe Thrombocytopenia

WINREVAIR may decrease platelet count. Severe thrombocytopenia may increase the risk of bleeding. In clinical studies, severe thrombocytopenia (platelet count $< 50,000/\text{mm}^3$ [$< 50 \times 10^9/\text{L}$]) occurred in 3% to 6% of patients taking WINREVAIR. Thrombocytopenia occurred more frequently in patients also receiving prostacyclin infusion.

Do not initiate treatment if platelet count is $< 50,000/\text{mm}^3$ [see *Dosage and Administration (2.3)*].

Monitor platelets before each dose for the first 5 doses, or longer if values are unstable, and periodically thereafter to determine whether dose adjustments are required. [see *Dosage and Administration (2.3)*, *Adverse Reactions (6.1)*].

5.3 Serious Bleeding

In clinical studies, serious bleeding (e.g., gastrointestinal, intracranial hemorrhage) was reported in 4% vs 1% (STELLAR) and 7% vs 5% (ZENITH) of patients taking WINREVAIR vs placebo, respectively [see *Clinical Studies (14.1)*]. Patients with serious bleeding were

more likely to be on prostacyclin background therapy and/or antithrombotic agents, or have low platelet counts. Advise patients about signs and symptoms of blood loss. Evaluate and treat bleeding accordingly. Do not administer WINREVAIR if the patient is experiencing serious bleeding [see *Warnings and Precautions (5.2)*, *Adverse Reactions (6.1)*].

5.4 Embryo-Fetal Toxicity

Based on findings in animal reproduction studies, WINREVAIR may cause fetal harm when administered to a pregnant woman. In animal reproduction studies, administration of WINREVAIR to pregnant rats and rabbits during organogenesis resulted in adverse developmental outcomes, including increased embryo-fetal mortality, alterations to growth, and structural variations at exposures 4-fold and 0.6-fold (based on area under the curve [AUC]) those occurring at the maximum recommended human dose (MRHD), respectively. Advise pregnant women of the potential risk to a fetus. Advise females of reproductive potential to use an effective method of contraception during treatment with WINREVAIR and for at least 4 months after the final dose [see *Use in Specific Populations (8.1, 8.3)*].

5.5 Impaired Fertility

Based on findings in animals, WINREVAIR may impair female and male fertility. Advise patients on the potential effects on fertility [see *Use in Specific Populations (8.3)*, *Nonclinical Toxicology (13.1)*].

6 ADVERSE REACTIONS

The following clinically significant adverse reactions are described elsewhere in the labeling:

- Erythrocytosis [see *Warnings and Precautions (5.1)*]
- Severe Thrombocytopenia [see *Warnings and Precautions (5.2)*]
- Serious Bleeding [see *Warnings and Precautions (5.3)*]
- Embryo-Fetal Toxicity [see *Warnings and Precautions (5.4)*]
- Impaired Fertility [see *Warnings and Precautions (5.5)*]

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.

STELLAR

The following data reflect exposure to WINREVAIR in the STELLAR trial. Adult PAH patients with WHO FC II or III (n=323) were randomized in a 1:1 ratio to receive WINREVAIR or placebo in combination with background standard of care therapies. Patients received a starting dose of 0.3 mg/kg via SC injection and the dose was increased to the target dose of 0.7 mg/kg administered once every 3 weeks for 24 weeks. After completing the primary 24-week treatment phase, patients continued into a long-term double-blind (LTDB) treatment period, maintaining their randomized treatment assignment, until all patients completed the primary treatment period. The median duration of treatment was 273 days in the placebo group and 313 days in the

WINREVAIR group [see *Clinical Studies (14.1)*].

The most common adverse reactions occurring in STELLAR ($\geq 10\%$ for WINREVAIR and at least 5% more than placebo) are shown in Table 3.

Table 3: Adverse Reactions $\geq 10\%$ in Patients Receiving WINREVAIR and at least 5% More Than Placebo in STELLAR*

Adverse reaction	WINREVAIR N=163	Placebo N=160
Headache	40 (24.5)	28 (17.5)
Epistaxis	36 (22.1)	3 (1.9)
Rash	33 (20.2)	13 (8.1)
Telangiectasia	27 (16.6)	7 (4.4)
Diarrhea	25 (15.3)	16 (10.0)
Dizziness	24 (14.7)	10 (6.3)
Erythema	22 (13.5)	5 (3.1)

* Double-blind placebo-controlled period + Long-term double-blind period of STELLAR

Increased Hemoglobin

Increases in Hgb were managed by dose delays (10%), dose reductions (6%), or both (5%). Shifts in Hgb from normal to above normal levels occurred in 87 (53%) patients receiving WINREVAIR and in 23 (14%) patients receiving placebo.

Thrombocytopenia

Decreases in platelets were managed by dose delays (2%), dose reductions (2%), or both (2%). Shifts in platelet count from normal to below normal occurred in 40 (25%) patients receiving WINREVAIR and in 26 (16%) patients receiving placebo.

Telangiectasia

In patients exposed to WINREVAIR who experienced telangiectasia, the median time to onset was 36.1 weeks.

Increased Blood Pressure

In patients taking WINREVAIR, mean systolic/diastolic blood pressure increased from baseline by 2.2/4.9 mmHg at 24 weeks. In patients taking placebo, the change from baseline in mean blood pressure was -1.6/-0.6 mmHg.

Treatment Discontinuation

The incidences of treatment discontinuations due to an adverse reaction were 4% in the WINREVAIR group and 7% in the placebo group. No specific adverse reactions causing treatment discontinuations occurred with a frequency greater than 1% and more often in the WINREVAIR group.

ZENITH

The following data reflect exposure to WINREVAIR in the ZENITH trial. Adult PAH patients with WHO FC III or IV at high risk of mortality (n=172) were randomized in a 1:1

ratio to treatment with WINREVAIR or placebo in combination with background standard of care therapies. Patients who did not experience a primary endpoint event remained in the Double-Blind Placebo-Controlled (DBPC) Treatment Period, while patients who experienced an event of PAH worsening-related hospitalization of ≥ 24 hours were eligible to enroll into the open-label, long-term follow-up (LTFU) study SOTERIA. The median duration of exposure was longer in the WINREVAIR group (435 days) than in the placebo group (268 days) [see *Clinical Studies (14.1)*].

The overall incidences of adverse reactions in both arms were higher in the ZENITH trial than in the STELLAR trial. Severe reduction in platelet count $< 50,000/\text{mm}^3$ ($< 50.0 \times 10^9/\text{L}$) occurred in 6% of patients taking WINREVAIR. In the WINREVAIR group, 1 patient (1%) discontinued study intervention due to an adverse event, compared with 4 patients (5%) in the placebo group.

Table 4: Adverse Reactions $\geq 10\%$ in Patients Receiving WINREVAIR and at least 5% More Than Placebo in ZENITH

Adverse reaction	WINREVAIR N=86	Placebo N=86
Infections	58 (67.4)	38 (44.2)
Epistaxis	39 (45.3)	8 (9.3)
Diarrhea	22 (25.6)	15 (17.4)
Telangiectasia	22 (25.6)	3 (3.5)
Increased hemoglobin	13 (15.1)	1 (1.2)
Rash	9 (10.5)	4 (4.7)
Erythema	9 (10.5)	3 (3.5)
Gingival bleeding	9 (10.5)	2 (2.3)

Uncontrolled Long-term Safety Data

PULSAR was a multicenter, randomized, double-blind, phase 2 trial that included a 24-week placebo-controlled treatment period, followed by an 18-month active-drug extension period. The safety profile in the long-term uncontrolled extension period of the PULSAR study was generally similar to that observed in the STELLAR study. Patients were treated with WINREVAIR 0.3 mg/kg or 0.7 mg/kg (n=104) and had a mean duration of exposure of 151 weeks (maximum 218 weeks).

Intrapulmonary Right-to-Left Shunting: Cases of intrapulmonary right-to-left shunting have been reported in a clinical trial with WINREVAIR. In SOTERIA, an ongoing open-label study of the long-term safety and efficacy of WINREVAIR, right-to-left intrapulmonary shunting has been reported in 2 participants ($< 0.5\%$) who developed worsening hypoxemia despite improved PAH hemodynamics. Post-marketing cases have also been reported.

Partial to complete improvement in oxygenation has been observed following discontinuation of WINREVAIR.

6.2 Post-marketing Experience

The following adverse reaction has been reported during post-approval use of WINREVAIR. Because these reactions are reported voluntarily from a population of

uncertain size, it is generally not possible to reliably estimate their frequency or establish a causal relationship to drug exposure.

Cardiac disorders: pericardial effusion

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

Based on findings in animal reproduction studies, WINREVAIR may cause fetal harm when administered to a pregnant woman. There are risks to the mother and the fetus associated with pulmonary arterial hypertension in pregnancy (*see Clinical Considerations*). There are no available data on WINREVAIR use in pregnant women to inform a drug-associated risk of major birth defects, miscarriage, or adverse maternal or fetal outcomes.

In animal reproduction studies, administration of WINREVAIR to pregnant rats and rabbits during the period of organogenesis resulted in adverse developmental outcomes, including embryo-fetal mortality, alterations to growth, and structural variations at exposures 4-fold and 0.6-fold (based on area under the curve [AUC]) above those occurring at the maximum recommended human dose (MRHD), respectively (*see Data*). Advise pregnant women of the potential risk to a fetus [*see Use in Specific Populations (8.3)*].

The background risk of major birth defects and miscarriage for the indicated population is not known. All pregnancies have a background risk of birth defect, loss, or other adverse outcomes. In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2% to 4% and 15% to 20%, respectively.

Report exposure during pregnancy or lactation to the Merck Sharp & Dohme, LLC Adverse Event reporting line at 1-877-888-4231.

Clinical Considerations

Disease-Associated Maternal and/or Embryo/Fetal Risk

In patients with pulmonary arterial hypertension, pregnancy is associated with an increased rate of maternal and fetal morbidity and mortality, including spontaneous abortion, intrauterine growth restriction, and premature labor.

Data

Animal Data

In embryo-fetal developmental toxicity studies, pregnant animals were dosed subcutaneously with sotatercept-csrk during the period of organogenesis. Sotatercept-csrk was administered to rats on gestation days 6 and 13 at doses of 5, 15, or 50 mg/kg and to rabbits on gestation days 7 and 14 at doses of 0.5, 1.5, or 5 mg/kg. Effects in both species included reductions in numbers of live fetuses and fetal body weights, delays in ossification, and increases in resorptions and post-implantation losses. In rats and rabbits, these effects were observed at exposures (based on area under the curve [AUC]) approximately 4-fold and 0.6-fold the maximum recommended

human dose (MRHD), respectively. In rats only, skeletal variations (increased number of supernumerary ribs and changes in the number of thoracic or lumbar vertebrae) occurred at an exposure 15-fold the human exposure at the MRHD.

In a prenatal and postnatal development study in rats, sotatercept-csrk was administered subcutaneously at doses of 1.5 and 5 mg/kg on gestation days 6 and 13, or at dosages of 1.5, 5, or 10 mg/kg during lactation on days 1, 8, and 15. There were no adverse effects in first filial generation (F1) pups from dams dosed during gestation at estimated exposures up to 2-fold the MRHD. In F1 pups from dams dosed during lactation, decreases in pup weight correlated with delays in sexual maturation at estimated exposures (based on AUC) ≥ 2 -fold the MRHD.

8.2 Lactation

Risk Summary

There are no data on the presence of sotatercept-csrk in human milk, the effects on the breastfed infant, or the effects on milk production. Because of the potential for serious adverse reactions in the breastfed child, advise patients that breastfeeding is not recommended during treatment with WINREVAIR, and for 4 months after the final dose.

8.3 Females and Males of Reproductive Potential

WINREVAIR may cause fetal harm when administered to pregnant women [*see Use in Specific Populations (8.1)*].

Pregnancy Testing

Pregnancy testing is recommended for females of reproductive potential before starting WINREVAIR treatment.

Contraception

Females

Advise female patients of reproductive potential to use effective contraception during treatment with WINREVAIR and for at least 4 months after the final dose if treatment is discontinued [*see Use in Specific Populations (8.1)*].

Infertility

Based on findings in animals, sotatercept-csrk may impair female and male fertility [*see Nonclinical Toxicology (13.1)*]. In male rats, although adverse histologic changes in reproductive organs were not reversible after a 13-week period, functional fertility demonstrated reversibility.

8.4 Pediatric Use

The safety and effectiveness of WINREVAIR have not been established in patients less than 18 years of age.

8.5 Geriatric Use

A total of 127 patients ≥ 65 years of age participated in clinical studies for PAH, of which 99 (78%) were treated with WINREVAIR. No differences in efficacy of WINREVAIR were observed between the < 65 -year-old and ≥ 65 -year-old subgroups.

With the exception of bleeding events (a collective group of adverse events of clinical interest), there were no differences in safety between the <65-year-old and ≥65-year-old subgroups. Bleeding events occurred more commonly in the older WINREVAIR subgroup, but with no imbalance between age subgroups for any specific bleeding event.

Clinical studies of WINREVAIR did not include sufficient numbers of patients aged 75 and older to determine whether they respond differently from younger patients.

10 OVERDOSAGE

In healthy volunteers, WINREVAIR dosed at 1 mg/kg resulted in increases in Hgb associated with hypertension; both improved with phlebotomy. In the event of overdose, monitor closely for increases in Hgb and blood pressure, and provide supportive care as appropriate. WINREVAIR is not dialyzable.

11 DESCRIPTION

Sotatercept-csrk is a homodimeric recombinant fusion protein consisting of the extracellular domain of the human activin receptor type IIA (ActRIIA) linked to the human IgG1 Fc domain. The molecular weight based on the amino acid sequence of sotatercept-csrk is approximately 78 kDa as a homodimer.

Sotatercept-csrk for injection is a sterile, preservative-free, white to off-white lyophilized cake or powder appearance in single-dose vials for subcutaneous administration after reconstitution.

Each 45 mg single-dose vial provides 45 mg of sotatercept-csrk and citric acid monohydrate (0.40 mg), polysorbate 80 (0.18 mg), sodium citrate (1.84 mg), and sucrose (72 mg) at pH 5.8. After reconstitution with 1 mL Sterile Water for Injection, the resulting concentration is 50 mg/mL of sotatercept-csrk and the nominal deliverable volume is 0.9 mL.

Each 60 mg single-dose vial provides 60 mg of sotatercept-csrk and citric acid monohydrate (0.53 mg), polysorbate 80 (0.24 mg), sodium citrate (2.45 mg), and sucrose (96 mg) at pH 5.8. After reconstitution with 1.3 mL Sterile Water for Injection, the resulting concentration is 50 mg/mL of sotatercept-csrk and the nominal deliverable volume is 1.2 mL.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Sotatercept-csrk, a recombinant activin receptor type IIA-Fc (ActRIIA-Fc) fusion protein, is an activin signaling inhibitor that binds to activin A and other TGF- β superfamily ligands. As a result, sotatercept-csrk improves the balance between the pro-proliferative (ActRIIA/Smad2/3-mediated) and anti-proliferative (BMPRII/Smad1/5/8-mediated) signaling to modulate vascular proliferation. In rat models of PAH, a sotatercept-csrk analog reduced inflammation and inhibited proliferation of endothelial and smooth muscle cells in diseased vasculature. These cellular changes were associated with thinner vessel walls, partial reversal of right ventricular remodeling, and improved

hemodynamics.

12.2 Pharmacodynamics

STELLAR

A greater decrease from baseline in pulmonary vascular resistance (PVR) was observed in the WINREVAIR group compared to the placebo group. The median treatment difference in PVR between sotatercept-csrk and placebo was -235 dynes*sec/cm⁵ (95% CI: -288, -181). Sotatercept-csrk steady state exposure at 0.7 mg/kg dose was associated with near maximal reduction in PVR based on exposure-response analysis. A greater decrease from baseline in NT-proBNP was observed in the WINREVAIR group compared to the placebo group. The median treatment difference in NT-proBNP between the sotatercept-csrk and placebo was -442 pg/mL (95% CI: -574, -310).

ZENITH

The median treatment difference in change in PVR from baseline between the sotatercept and placebo groups after 24 weeks was -340 dynes*sec/cm⁵ (95% CI: -511, -168). The median treatment difference in change in NT-proBNP from baseline between the sotatercept-csrk and placebo groups after 24 weeks was -2339 pg/mL (95% CI: -3379, -1299). The median treatment difference in change from baseline in mean pulmonary artery pressure (mPAP) between the sotatercept-csrk and placebo groups after 24 weeks was -21.2 mm Hg (95% CI: -27.8, -14.6).

12.3 Pharmacokinetics

Following subcutaneous administration of 0.7 mg/kg WINREVAIR every three weeks to PAH patients (PULSAR, SPECTRA, and STELLAR), the steady state geometric mean (%CV) area under the time concentration curve (AUC) is 171.3 mcg×d/mL (34.2%), and peak concentration (C_{max}) is 9.7 mcg/mL (30%). Sotatercept-csrk AUC and C_{max} increased proportionally with dose. Steady state is achieved after approximately 15 weeks following initiation of multiple dosing. The accumulation ratio of sotatercept-csrk AUC is approximately 2.2. Sotatercept-csrk pharmacokinetics were similar in PAH participants in ZENITH.

Absorption

Following subcutaneous administration, the absolute bioavailability of sotatercept-csrk is approximately 66%. The sotatercept-csrk median time to peak drug concentration (T_{max}) is approximately 7 days (range from 2 to 8 days) following multiple SC administration every 4 weeks.

Distribution

The population PK model estimated volume of distribution (%CV) of sotatercept-csrk at steady state is approximately 5.3 L (27.3%) in patients with PAH.

Elimination

The sotatercept-csrk effective half-life is approximately 24 days and its clearance is approximately 0.18 L/day.

Metabolism

Sotatercept-csrk is expected to be metabolized into small peptides by catabolic

pathways.

Specific Populations

No clinically significant differences in sotatercept-csrk pharmacokinetics (PK) were observed based on age (18 to 81 years of age), sex, race, mild to moderate (eGFR ranging from 30 to 89 mL/min) renal impairment (PAH patients), or end-stage kidney disease (eGFR <15 mL/min) with dialysis. Based on limited data, severe renal impairment (eGFR ranging from 15 to 30 mL/min, n=3) had no impact on the PK of sotatercept-csrk. Sotatercept-csrk is not dialyzable. The effect of hepatic impairment on the PK of sotatercept-csrk has not been studied.

Body Weight

The clearance (CL) and central volume of distribution (Vc) increase with increase in body weight. This effect is not clinically significant when sotatercept-csrk is administered using weight-based dosing as recommended.

12.6 Immunogenicity

The observed incidence of anti-drug antibodies is highly dependent on the sensitivity and specificity of the assay. Differences in assay methods preclude meaningful comparisons of the incidence of anti-drug antibodies in the studies described below with the incidence of anti-drug antibodies in other studies, including those of WINREVAIR or of other sotatercept-csrk products.

During the 24-week treatment period in STELLAR, 27% (44/163) of sotatercept-csrk-treated patients developed anti-sotatercept-csrk antibodies (ADA). Among the 44 ADA-positive patients, 12 (27%) tested positive for neutralizing antibodies against sotatercept-csrk.

During the ZENITH trial, with a median sotatercept treatment duration of 435 days, 43% (36/84) of patients developed anti-sotatercept antibodies. Among these 36 patients, 20 (56%) tested positive for neutralizing antibodies against sotatercept.

There were no identified clinical effects of anti-sotatercept-csrk antibodies on pharmacokinetics, pharmacodynamics, safety, or effectiveness of sotatercept-csrk over the treatment duration of 24 weeks at the recommended dosage in these studies.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

No carcinogenicity or mutagenicity studies have been conducted with sotatercept-csrk.

In a fertility and early embryonic development study in female rats, sotatercept-csrk was administered SC once weekly at doses of 5, 15, and 50 mg/kg beginning 2 weeks prior to mating and through gestation day 7. At doses ≥ 15 mg/kg (≥ 9 fold the MRHD, based on estimated AUC), pregnancy rates were decreased and there were increases in preimplantation and postimplantation loss and reductions in live litter size. Increased estrous cycle duration occurred at 50 mg/kg only (21-fold the MRHD, based on estimated AUC).

In a fertility study in male rats, sotatercept-csrk was administered SC once weekly at

doses of 0.3, 3, and 30 mg/kg for 13 weeks (beginning 10 weeks prior to mating). A subset of animals was examined after a 13-week recovery period. At ≥ 0.3 mg/kg (0.5-fold the MRHD, based on estimated AUC) there were non-reversible histologic changes in the efferent ducts, testes, and epididymides. Reversible decreases in functional fertility endpoints occurred at 30 mg/kg (20-fold the MRHD, based on estimated AUC).

14 CLINICAL STUDIES

14.1 Pulmonary Arterial Hypertension

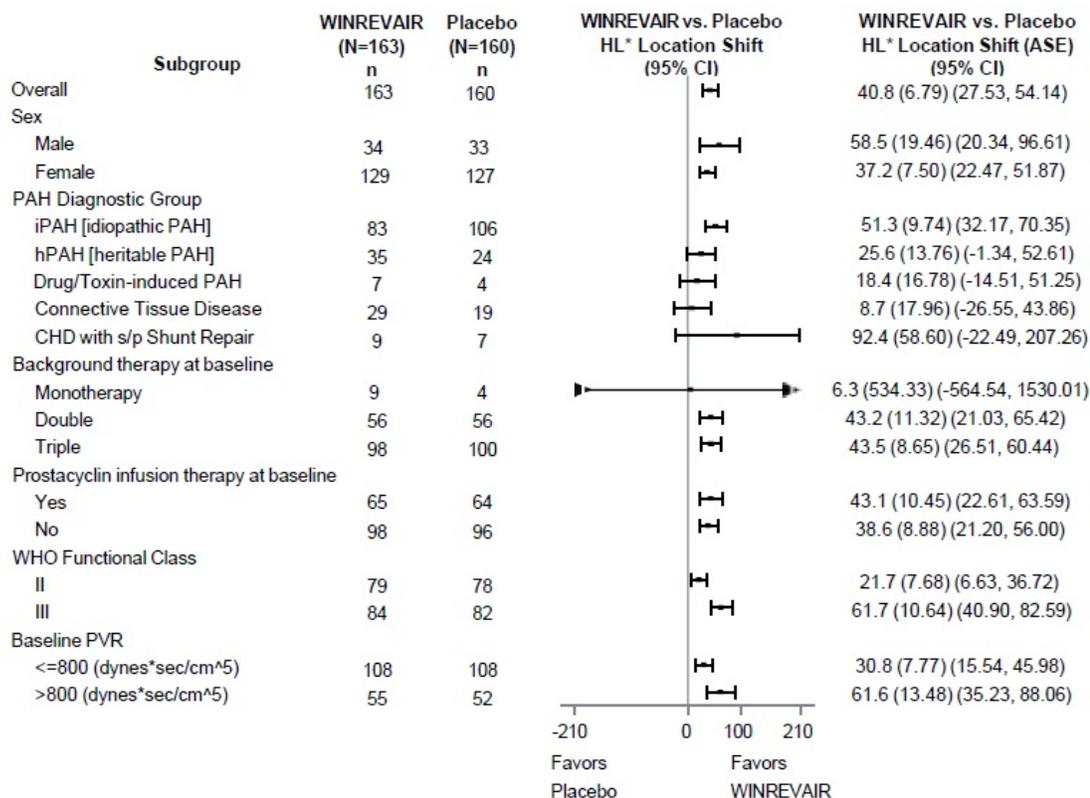
STELLAR

The efficacy of WINREVAIR was evaluated in adult patients with PAH in the STELLAR trial (NCT04576988). STELLAR was a global, double-blind, placebo-controlled, multicenter, parallel-group clinical trial in which 323 patients with PAH (WHO Group 1, FC II or III) were randomized 1:1 to WINREVAIR (target dose 0.7 mg/kg) (n=163) or placebo (n=160) administered subcutaneously once every 3 weeks.

Participants were: 79% female; had a median age of 48 years (range: 18 to 82 years), and median body weight of 68 kg (range 38 to 141 kg); and 89% White, 2% Black/African American, 2% Asian, 0.3% American Indian or Alaska Native, 0.3% Native Hawaiian or Other Pacific Islander, 6% Missing/Other races. The most common PAH etiologies were idiopathic PAH (59%), heritable PAH (18%), and PAH associated with connective tissue diseases (CTD) (15%). STELLAR excluded patients with human immunodeficiency virus (HIV)-associated PAH, PAH associated with portal hypertension, schistosomiasis-associated PAH, and pulmonary veno occlusive disease. The mean time from PAH diagnosis to screening was 8.8 years. Most participants were receiving either three (61%) or two (35%) background drugs for PAH, and 40% were receiving prostacyclin infusions. Patients had a WHO FC II (49%) or III (51%) at baseline.

The primary efficacy endpoint was the change from baseline at Week 24 in 6-Minute Walk Distance (6 MWD). In the WINREVAIR group, the placebo-adjusted median increase in 6 MWD was 41 meters (95% CI: 28, 54; $p < 0.001$). Figure 1 displays placebo-adjusted changes in 6 MWD at Week 24 in relevant subgroups.

Figure 1: Change from Baseline in 6-Minute Walk Distance (meters) at Week 24 in Subgroups in STELLAR



*Hodges-Lehmann location shift from placebo estimate (median of all paired differences). ASE = asymptotic standard error.

Change from baseline in 6 MWD at Week 24 for subjects who died was imputed to -2000 meters to receive the worst rank. Change from baseline in 6 MWD at Week 24 for subjects who had missing data due to a non-fatal clinical worsening event was imputed to -1000 meters to receive the next-worst rank.

Treatment with WINREVAIR led to an improvement from baseline by at least 1 WHO FC at Week 24 in 29% of patients compared to 14% of patients treated with placebo (p<0.001).

Treatment with WINREVAIR resulted in an 84% reduction in the occurrence of death from any cause or PAH clinical worsening events compared to placebo (see Table 5 and Figure 2). These outcomes were captured until the last patient completed the Week 24 visit (data up to the data cutoff; median duration of exposure 33.6 weeks).

Table 5: Death from Any Cause or PAH Clinical Worsening Events in STELLAR

	WINREVAIR (N=163) n (%)	Placebo (N=160) n (%)	Hazard Ratio (95% CI)
Number of subjects who experienced death or at least one clinical worsening event	9 (5.5)	42 (26.3)	0.16 (0.08, 0.35) p<0.001
Assessment of clinical worsening events*			
Death	2 (1.2)	7 (4.4)	
Worsening related listing for			

worsening-related listing for lung and/or heart transplant	1 (0.6)	2 (1.3)
Need to initiate rescue therapy with an approved PAH therapy or the need to increase the dose of infusion prostacyclin by 10% or more	2 (1.2)	17 (10.6)
Need for atrial septostomy [†]	0 (0.0)	0 (0.0)
PAH-specific hospitalization (≥24 hours)	0 (0.0)	8 (5.0)
Deterioration of PAH [‡]	4 (2.5)	15 (9.4)

N = number of subjects in the category.

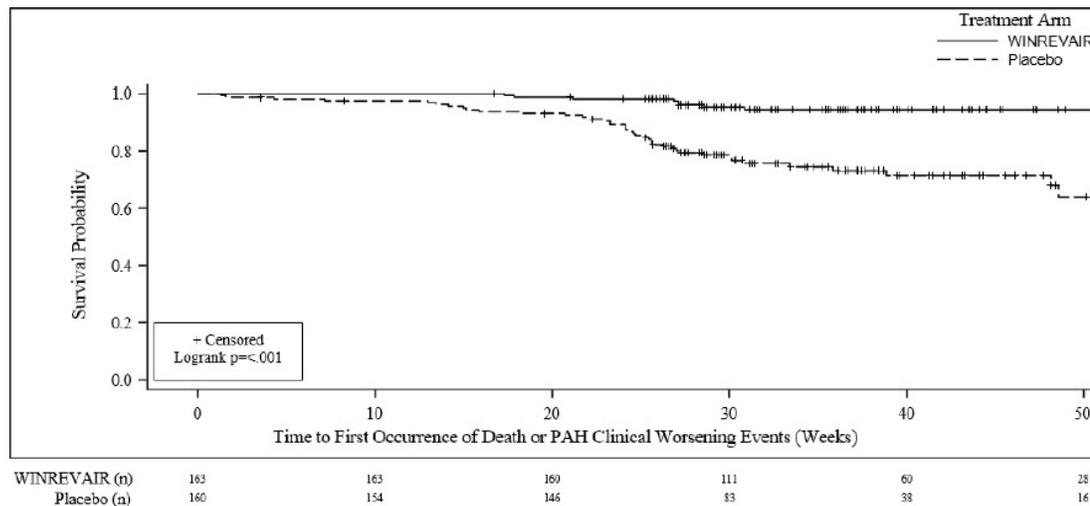
6 MWT = 6-Minute Walking Test

* A subject can have more than one assessment recorded for their clinical worsening.

† There were no events of atrial septostomy.

‡ Deterioration of PAH is defined by both of the following events occurring at any time, even if they began at different times, as compared to their baseline values: (a) Worsened WHO functional class (II to III, III to IV, II to IV, etc.); and (b) Decrease in 6 MWD by ≥15% (confirmed by two 6 MWTs at least 4 hours apart but no more than one week).

Figure 2: Time to Death from Any Cause or First Occurrence of PAH Clinical Worsening Event Kaplan-Meier Plot in STELLAR



ZENITH

The efficacy of WINREVAIR was evaluated in adult PAH patients with WHO FC III or IV at high risk of mortality in the ZENITH trial (NCT04896008). ZENITH was a global, double-blind, placebo-controlled, multicenter, parallel-group clinical trial in which 172 patients were randomized 1:1 to WINREVAIR (target dose 0.7 mg/kg) (n=86) or placebo (n=86) administered subcutaneously once every 3 weeks. Efficacy was evaluated at the pre-specified interim analysis which occurred when 61 patients experienced a primary endpoint event and median patient time on study was 273 days.

The demographic and baseline clinical characteristics were similar between the WINREVAIR and placebo groups. Participants were: 77% female; had a median age of 58

years (range: 18 to 75 years); and 87% White, 5% Black/African American, 4% Asian, 1% American Indian or Alaska Native, 3% Missing/Other race, 74% were FC III and 26% were FC IV. The most common PAH etiologies were idiopathic PAH (50%), PAH associated with connective tissue diseases (CTD) (28%), and heritable PAH (11%). The mean time since PAH diagnosis to screening was 8 years. Participants were on background PAH treatment, 72% on triple therapy, 28% on double therapy, and 59% on prostacyclin infusion therapy. The REVEAL Lite 2 risk score was <9 for 2% of participants, 9 to 10 for 67% of participants, and ≥ 11 for 30% of participants. The ZENITH trial excluded patients diagnosed with human immunodeficiency virus (HIV)-associated PAH, PAH associated with portal hypertension, pulmonary veno-occlusive disease, or pulmonary capillary hemangiomatosis or overt signs of capillary and/or venous involvement.

The primary efficacy endpoint was time to first event of all-cause death, lung transplantation, or PAH worsening-related hospitalization of ≥ 24 hours. In the WINREVAIR treatment group, the risk of a first event of all-cause death, lung transplantation, or PAH worsening-related hospitalization of ≥ 24 hours was 76% lower compared with the placebo group (HR: 0.24; 95% CI: 0.13, 0.43; $p < 0.0001$) (see Table 6). Fewer participants in the WINREVAIR group (15 [17%]) than in the placebo group (47 [55%]) had a primary endpoint event as of the data cutoff. Based on the primary endpoint result, the study was stopped for favorable efficacy at the interim analysis.

The treatment effect of WINREVAIR was consistent across the prespecified subgroups (see Figure 4).

Table 6: Primary Endpoint Results in ZENITH

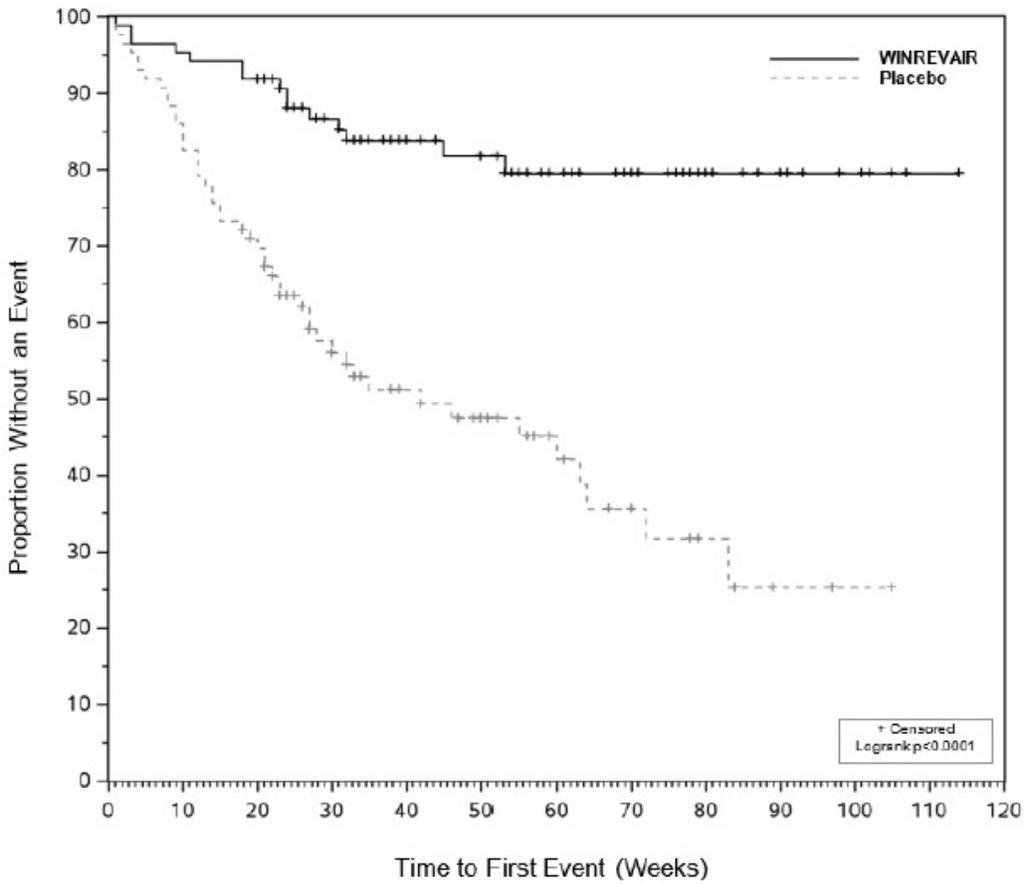
	WINREVAIR (N=86) n (%)	Placebo (N=86) n (%)	Hazard Ratio (95% CI) p-value
Number of participants with at least 1 primary event*	15 (17.4)	47 (54.7)	0.24 (0.13, 0.43) <0.0001
Components of primary endpoint†			
All-cause death	7 (8.1)	13 (15.1)	
Lung transplantation	1 (1.2)	6 (7.0)	
PAH worsening-related to hospitalization (≥ 24 hours)	8 (9.3)	43 (50.0)	

* The primary composite endpoint analysis includes the first occurrence of an adjudicated morbidity-mortality event up to the data cutoff. All deaths up to the data cutoff are included, regardless of adjudication and regardless of whether they occurred during or post-ZENITH, except for those occurring after lung transplantation or enrollment in SOTERIA.

† Shows each component of the composite primary endpoint as a standalone outcome. A participant is included in more than one row if multiple events meeting primary endpoint definition were observed.

Figure 3: Time to First Event in ZENITH of All-cause Death, Lung Transplantation, or PAH Worsening-related

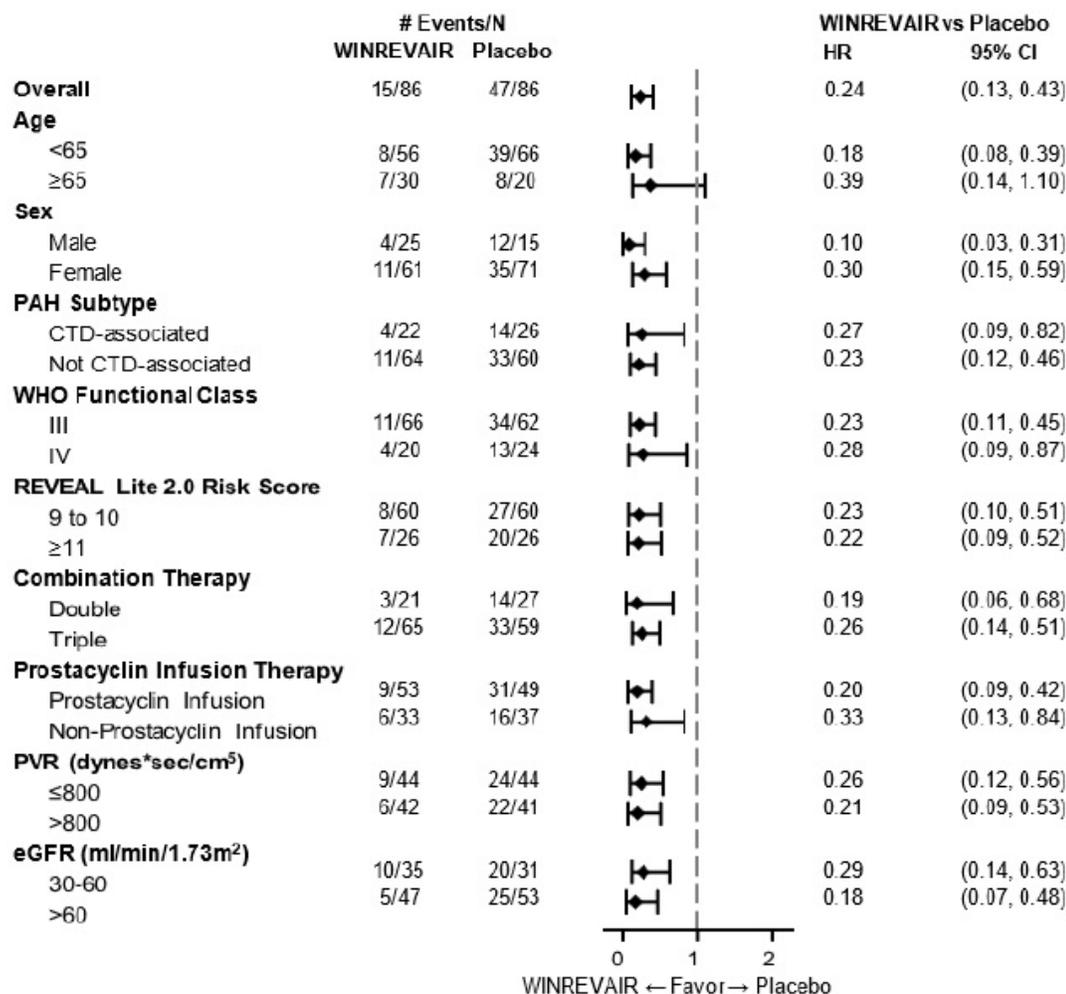
Hospitalization of ≥ 24 Hours Kaplan-Meier Plot



		0	10	20	30	40	50	60	70	80	90	100	110	120
WINREVAIR (n)	86	82	79	61	51	40	28	21	13	9	5	1	0	0
Placebo (n)	86	74	59	38	28	23	15	10	5	2	1	0	0	0

n= Number of subjects at Risk

Figure 4: Hazard Ratio for the Primary Endpoint in Subgroups in ZENITH



Subgroup analyses were not displayed if the number of participants in subgroup category was less than 10% of FAS.

For participants with REVEAL Lite 2.0 risk score <9 at screening, they were grouped under “9 to 10” for the analyses.

The secondary endpoint of overall survival (OS) included all deaths up to the data cutoff, except for those occurring after lung transplantation or enrollment in a long-term follow-up study. Twenty OS events were observed (7 deaths in the WINREVAIR treatment group and 13 deaths in the placebo group). The point estimate for the OS HR favored the WINREVAIR treatment group over the placebo group.

Subsequent secondary endpoints were not eligible to be tested due to the hierarchical testing strategy.

16 HOW SUPPLIED/STORAGE AND HANDLING

16.1 How Supplied

WINREVAIR (sotatercept-csrk) for injection is a white to off-white lyophilized cake or powder appearance supplied in single-dose vials (45 mg or 60 mg) packaged in kits that contain one measuring syringe and one safety needle. Each kit also contains Sterile Water for Injection in prefilled syringes necessary to reconstitute the product, vial adapter(s), and alcohol pads, as shown in Table 7.

Table 7: Kit Contents

Kit	Vial Adapters	Alcohol Pads	Sterile Water for Injection in prefilled syringes	NDC
(1) 45 mg vial	1	4	(1) 1 mL syringe	NDC # 0006-5090-01
(1) 60 mg vial	1	4	(1) 1.3 mL syringe	NDC # 0006-5091-01
(2) 45 mg vials	2	8	(2) 1 mL syringes	NDC # 0006-5087-01
(2) 60 mg vials	2	8	(2) 1.3 mL syringes	NDC # 0006-5088-01

16.2 Storage and Handling

Store vials refrigerated at 2°C to 8°C (36°F to 46°F) in original carton to protect from light. Do not freeze.

The kit should remain in the refrigerator until ready for use. The unused kit can be out of the refrigerator for (up to 25°C/77°F) up to 24 hours. For additional information on temperature excursions, call Merck Sharp & Dohme LLC at 1-800-672-6372.

17 PATIENT COUNSELING INFORMATION

Advise patients to read the FDA-approved patient labeling (Patient Information and IFU).

Discuss the following with patients prior to and during treatment with WINREVAIR.

Erythrocytosis

Caution patients that WINREVAIR may raise Hgb to levels that increase their risk of thrombotic events. Inform patients that Hgb levels will be assessed before at least the first 5 doses and then periodically, as dosage may need to be adjusted [see *Warnings and Precautions (5.1)*].

Severe Thrombocytopenia

Caution patients that WINREVAIR may cause platelet count to decrease, which if severe could cause bleeding. Inform patients that platelet count will be assessed before at least the first 5 doses and then periodically, as dosage may need to be adjusted [see *Warnings and Precautions (5.2)*].

Serious Bleeding

Inform patients of the possibility of serious bleeding, which is more likely to occur if they have low platelet counts or while on prostacyclin background therapy and/or antithrombotic agents. Advise patients to notify their healthcare provider about signs and symptoms of bleeding [see *Warnings and Precautions (5.3)*].

Embryo-Fetal Toxicity

Advise females of reproductive potential of the potential risk to a fetus. Advise females

of reproductive potential to use effective contraception while receiving WINREVAIR and for at least 4 months after the final dose. Advise females to contact their healthcare provider if they become pregnant, or if pregnancy is suspected during treatment with WINREVAIR [see *Warnings and Precautions (5.4)*, *Use in Specific Populations (8.1)*]. Report exposure during pregnancy or lactation to the Merck Sharp & Dohme, LLC Adverse Event reporting line at 1-877-888-4231.

Lactation

Advise females not to breastfeed during treatment with WINREVAIR and for 4 months after the final dose [see *Use in Specific Populations (8.2)*].

Females and Males of Reproductive Potential

Advise females and males of reproductive potential that WINREVAIR may impair fertility [see *Use in Specific Populations (8.3)*, *Nonclinical Toxicology (13.1)*].

Administration by Patient or Caregiver

Review the IFU with the patient or caregiver step-by-step. Provide training to the patient or caregiver regarding proper preparation and administration of WINREVAIR and decide whether a patient or caregiver is capable of preparing and administering WINREVAIR independently [see *Dosage and Administration (2.4)*].

Make sure the patient or caregiver can do the following correctly:

- reconstitute the medicine,
- measure the correct amount of medicine according to the patient's prescription,
- select and prepare a proper injection site, and
- inject the medicine subcutaneously.

Incorrect Dose or Missed Dose

Inform patients to call their healthcare provider for further instruction if they take more than or less than the correct dose. Advise them about signs/symptoms to monitor for and what to do if any of these signs/symptoms should occur. Advise them that additional laboratory tests may be required prior to the next scheduled dose to ensure that the next dose can be safely administered.

Instruct the patient that if they miss the prescribed dose of WINREVAIR, they should take it within 3 days and maintain the original schedule for the next dose. If not taken within 3 days, instruct them to call their healthcare provider for guidance [see *Dosage and Administration (2.2)*].

Manufactured by:

Merck Sharp & Dohme LLC, Rahway, NJ 07065, USA

U.S. license number 0002

For patent information: www.msd.com/research/patent

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uspi-mk7962-i-2512r004

WINREVAIR™ (WIN-reh-vair)
(sotatercept-csrk)
for injection, for subcutaneous use

What is WINREVAIR?

WINREVAIR is a prescription medicine used to treat adults with pulmonary arterial hypertension (PAH). PAH is a type of high blood pressure in the arteries of your lungs. WINREVAIR can:

- improve your ability to exercise and perform normal activities with fewer symptoms, and
- reduce the risk of your physical condition and symptoms worsening, including lowering your risk of hospitalization for PAH, lung transplant, and death.

It is not known if WINREVAIR is safe and effective in children under 18 years of age.

What should I tell my healthcare provider before taking WINREVAIR?

- Tell your healthcare provider about:
 - **All of your medical conditions**
 - **All of the medicines you take, including** prescriptions and over-the-counter medicines, vitamins, or herbal supplements

If you are able to get pregnant:

- WINREVAIR may harm your unborn baby. Tell your healthcare provider if you are pregnant or planning to get pregnant.
- Tell your healthcare provider right away if you become pregnant or think you may be pregnant during treatment with WINREVAIR.
- Your healthcare provider should do a pregnancy test before you start taking WINREVAIR.
- You should use effective birth control during treatment with WINREVAIR and for at least 4 months after your final dose. Ask your healthcare provider about birth control methods that may be right for you.
- You or your healthcare provider should report any exposure to WINREVAIR during pregnancy by calling 1-877-888-4231.

If you are breastfeeding or planning to breastfeed:

- Tell your healthcare provider if you are breastfeeding or planning to breastfeed. It is not known if WINREVAIR passes into breast milk.
- Do not breastfeed during treatment with WINREVAIR and for 4 months after your final dose. Talk to your healthcare provider about the best way to feed your baby.
- You or your healthcare provider should report any exposure to WINREVAIR during breastfeeding by calling 1-877-888-4231.

Before you take WINREVAIR:

- If your healthcare provider decides that you or a caregiver can give your injections of WINREVAIR at home, you or your caregiver should receive training from your healthcare provider on the right way to prepare and inject WINREVAIR. Do not try to inject WINREVAIR until you have been shown how to inject WINREVAIR by your healthcare provider.
- Your healthcare provider will do a blood test before your first 5 doses of WINREVAIR, longer if needed, and then from time to time to check your levels of hemoglobin (a protein in red blood cells that carries oxygen) and platelets (blood cells that help blood clot). After each of these blood tests, your healthcare provider may delay

treatment or change your dose if needed.

- **See the detailed “Instructions for Use” booklet that comes with WINREVAIR for information on how to prepare and inject a dose of WINREVAIR.**

How should I take WINREVAIR?

- Use WINREVAIR exactly as your healthcare provider tells you to.
- You will give WINREVAIR every 3 weeks as an injection just under your skin (subcutaneous) in your:
 - **stomach** (abdomen) at least 2 inches away from the belly button, **or**
 - **upper thigh**
- You should inject WINREVAIR right away after mixing the medicine powder with the sterile water for injection, but no later than 4 hours after mixing.

Your prescribed dose:

- Your healthcare provider will tell you how much WINREVAIR to inject and when to inject it. This is because your prescribed dose depends on your body weight and blood tests.
- Do not change your dose or stop taking WINREVAIR without talking to your healthcare provider.
- Do not take WINREVAIR more often than your healthcare provider tells you to. If you are not sure when to take WINREVAIR, call your healthcare provider.
- Your healthcare provider may delay treatment, change your dose, or stop treatment depending on how you respond to WINREVAIR.

What if I take the wrong dose or miss my dose?

- If you give more than or less than your prescribed dose, call your healthcare provider right away.
- If you miss your dose of WINREVAIR:
 - If it has been 3 days or fewer, give it as soon as possible. You can then follow your usual schedule for your next dose.
 - If it has been more than 3 days, call your healthcare provider for what to do.

What are the possible side effects of WINREVAIR?

WINREVAIR may cause serious side effects including:

- **High level of hemoglobin in your blood.** High levels of hemoglobin are common with WINREVAIR and can be severe and increase your risk for blood clots. Your healthcare provider will do blood tests to check your hemoglobin levels before starting and regularly during treatment with WINREVAIR.
- **Severely low number of platelets in your blood.** Low platelet counts are common with WINREVAIR and can be severe and increase your risk of bleeding. Your healthcare provider will do blood tests to check your platelet levels before starting and regularly during treatment with WINREVAIR. Tell your healthcare provider if you develop easy bruising or bleeding, bleeding that does not stop, or nosebleeds.
- **Serious bleeding.** You are more likely to have serious bleeding if you have low platelet counts, take prostacyclin medicines to treat PAH, or take medicines to help keep blood clots from forming. Tell your healthcare provider if you develop any signs or symptoms of bleeding, including:

- vomiting blood or your vomit looks like coffee-grounds
- pink or brown urine
- red or black stools that look like tar
- coughing up blood or blood clots
- persistent headaches
- nausea
- dizziness or feeling weak
- persistent abdominal cramps
- severe back pain
- abnormally heavy menstrual bleeding

- **Decreased fertility.** WINREVAIR may affect female and male fertility, which may affect your ability to have children. Talk to your healthcare provider if this is a concern for you.

The most common side effects of WINREVAIR include:

- infections
- headache
- nose bleeds
- rash
- tiny blood vessels that look like pink or red lines on the skin (spider veins)
- diarrhea
- dizziness
- redness
- bleeding gums

These are not all the possible side effects of WINREVAIR. Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

How should I store WINREVAIR?

- Store WINREVAIR in the refrigerator at 36°F to 46°F (2°C to 8°C). Do not freeze.
- Store in the original carton to protect from light.

Keep WINREVAIR and all medicines out of the reach of children and pets.

General information about the safe and effective use of WINREVAIR.

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

What are the ingredients in WINREVAIR?

Active ingredient: sotatercept-csrk

Inactive ingredient(s): citric acid monohydrate, polysorbate 80, sodium citrate, and sucrose.

Manufactured by: Merck Sharp & Dohme LLC, Rahway, NJ 07065, USA.

US License Number 0002

For patent information: www.msd.com/research/patent

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For more information, call 1-877-888-4231, or go to www.WINREVAIR.com.

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

Revised: October 2025

Important: Read this booklet first Important

INSTRUCTIONS FOR USE Mix

WINREVAIR™ (WIN-reh-vair) Withdraw
(sotatercept-csrk)

for injection, for subcutaneous use
How to use your WINREVAIR injection kit



For package containing 1 vial

Inject

For subcutaneous injection only (inject directly under the skin)

In this booklet:

Before you start

Important information for patients or caregivers

Get to know the parts of your kit

Important information you need to know before injecting WINREVAIR

Storing your injection kit

Get started

● **Mix** powdered medicine into liquid form

● **Withdraw** your prescribed dose

● **Inject** your medicine

How to throw away WINREVAIR

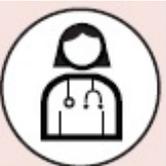
Frequently asked questions

Before you start:



Read this booklet

Read this Instructions for Use from start to finish before using WINREVAIR and each time you get a refill. There may be new information.



Start with your healthcare provider

Do not use WINREVAIR until your healthcare provider has shown you or your caregiver the right way to prepare and inject it. Your healthcare provider should show you how to inject WINREVAIR before you use it for the first time.

Questions?



If you have questions about how to give WINREVAIR the right way or need more information, call your pharmacy or healthcare provider.

For general product information visit www.WINREVAIR.com or call Merck Sharp & Dohme LLC at 1-800-672-6372.



Important information for patients or caregivers

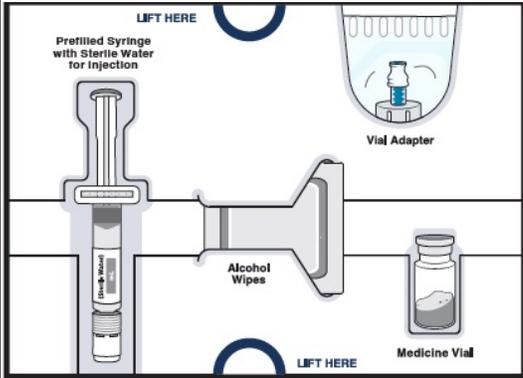
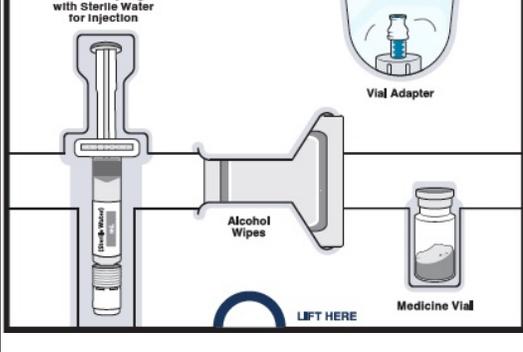
Before you prepare and inject WINREVAIR, you must be:

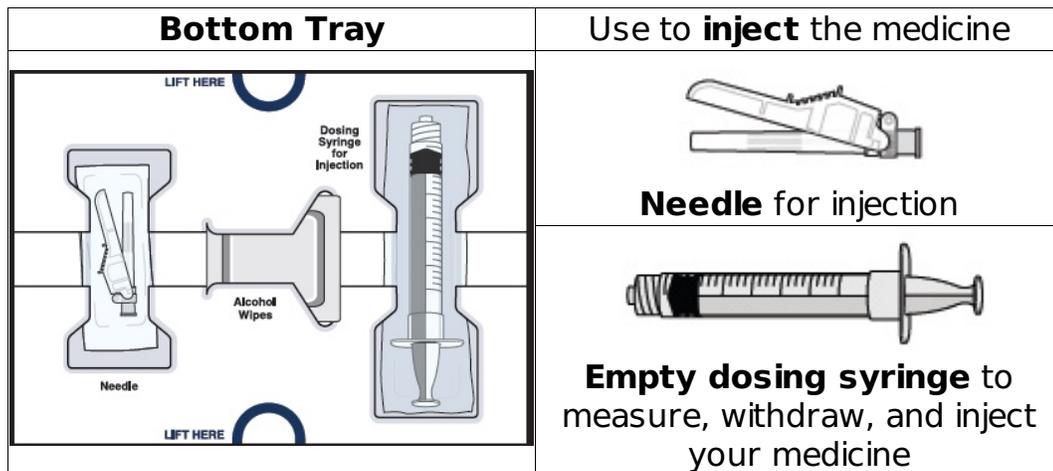
- trained by a healthcare provider following this Instructions for Use booklet step-by-step, and
- capable of independent administration of WINREVAIR as determined by a healthcare provider

Before using WINREVAIR, make sure you are trained to do the following correctly:

1. Reconstitute (mix) the medicine
2. Measure the correct amount of medicine according to your prescription
3. Select and prepare a proper injection site
4. Inject the medicine subcutaneously

Get to know the parts of your kit

Top Tray	Use to mix the medicine
	 <p>Vial Medicine powder</p> <p>1 Vial of WINREVAIR medicine powder</p>
	 <p>1 Prefilled syringe with sterile water for injection to mix medicine powder into liquid form</p>
	 <p>1 Vial adapter to connect the vial and syringe</p>

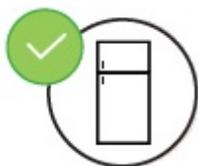


Important information you need to know before injecting WINREVAIR

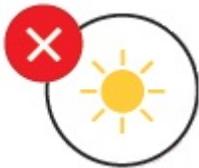
- You must mix WINREVAIR before using it. Make sure the medicine powder in the vial is completely dissolved before you inject.
- **Check your prescribed dose (amount in 'mL') each time you use WINREVAIR. Your prescribed dose may change.** Your prescribed dose is on the prescription label on your kit.
- Use only the supplies that are in the kit to prepare your prescribed dose.
- Do not open the package or mix the medicine until you are ready to use it.
- **Do not reuse any of the supplies.** After your injection, throw away the used vials with any remaining WINREVAIR medicine, needle, caps, and syringes in a sharps container. See pages 36-37 for more information.



Storing your injection kit



Store WINREVAIR injection kit in the refrigerator at 36°F to 46°F (2°C to 8°C). Do not freeze.



Store in the original carton to protect from light.



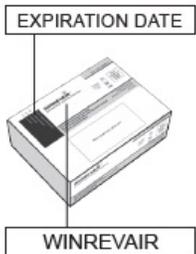
Keep injection kit out of the reach of children and pets.

Get started

Before you can prepare and inject WINREVAIR, you must first be trained and determined to be capable of independent administration of WINREVAIR by a healthcare provider.

1 Check WINREVAIR injection kit and expiration date

Remove the WINREVAIR injection kit from the refrigerator.



✓ **Check the expiration date and look for any signs of damage** on the kit or on the supplies.

⚠ If expired or damaged, do not use.
Call your pharmacy to get a new kit.

✓ **Check that you have the medicine** that your healthcare provider prescribed.

2 Let your kit come to room temperature, gather supplies, and wash your hands

Wait 15 minutes to allow your kit to warm to room temperature. Cold medicine is more painful to inject.



Along with your kit, gather these items and find a clean, flat surface where you will prepare and inject your dose.



Sharps disposal container



Gauze,

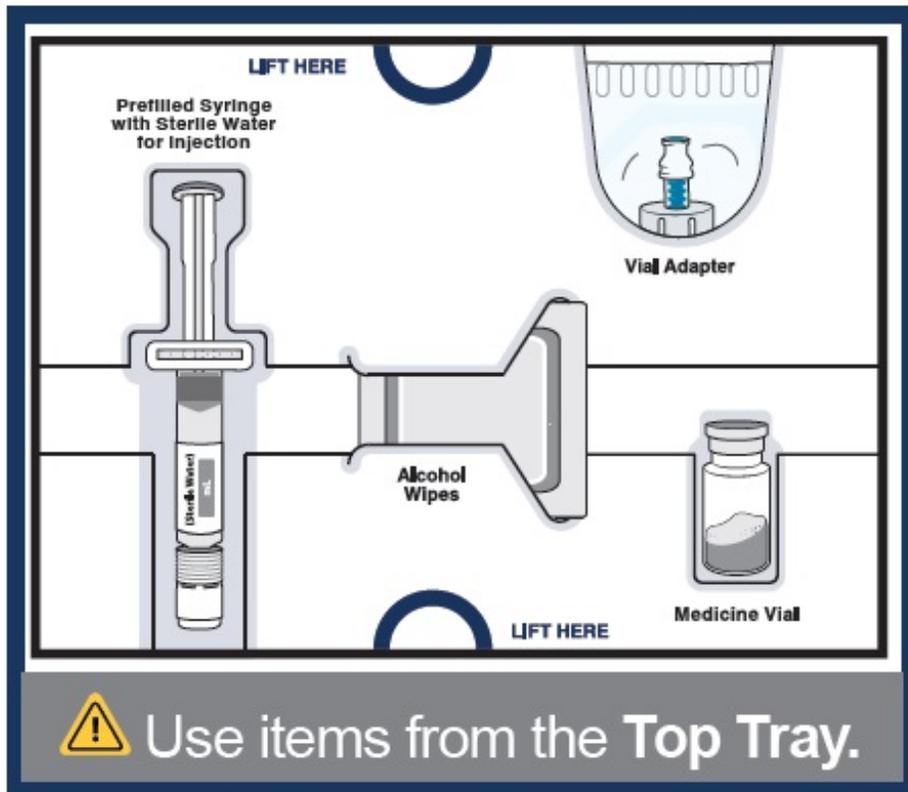
Wash your hands with soap and water.



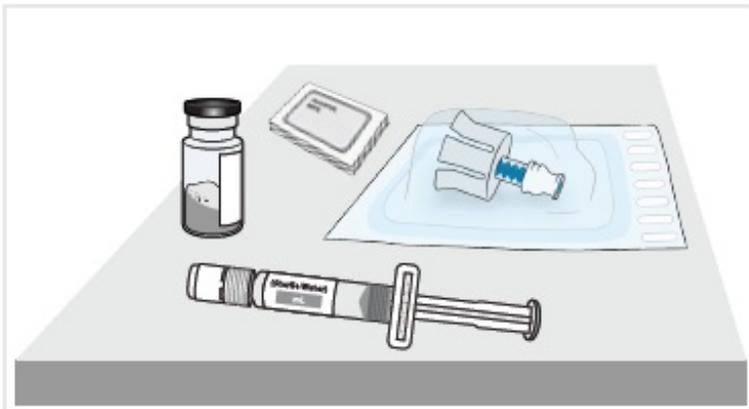
cotton ball, or
bandage

Mix powdered medicine into liquid form

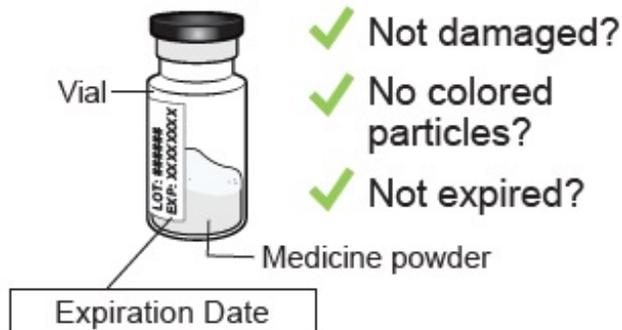
Start with the Top Tray



3 Remove medicine vial, prefilled syringe with sterile water for injection, alcohol wipes, and vial adapter from the kit.



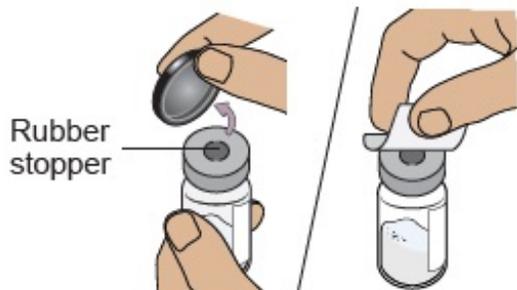
4a Check medicine vial



Check the vial label to **confirm the medicine is not expired**.
 Inspect the medicine powder.
 It should be white to off-white.

⚠ Do not use if expired, damaged, discolored, or you can see particles in it.

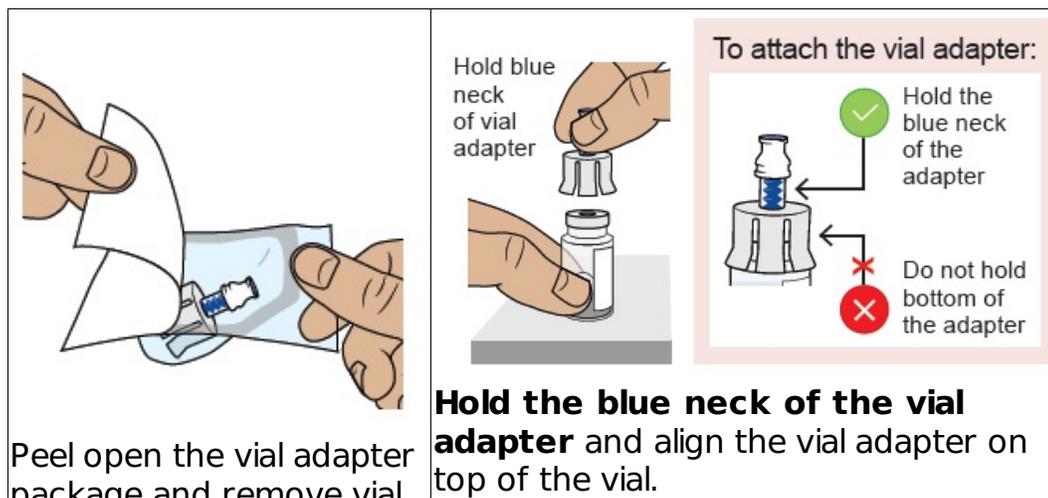
4b Remove plastic cap and clean vial



Flip off the plastic cap and **clean the rubber stopper on top of the vial** with an alcohol wipe.

⚠ Do not use if vial cap is missing.
 ⚠ Do not touch the cleaned rubber stopper.
 Set vial aside on a clean, flat surface.

5a Align vial adapter to vial

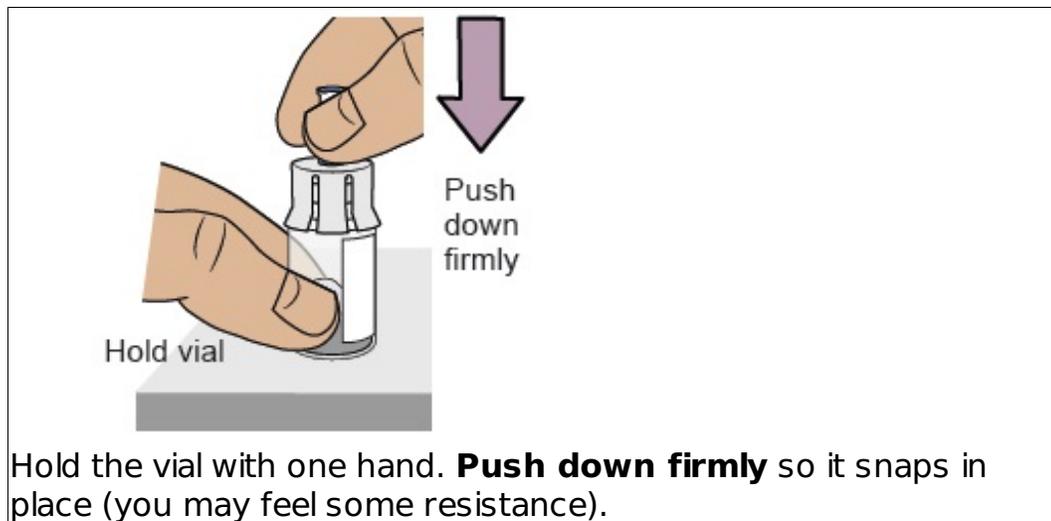


Peel open the vial adapter package and remove vial

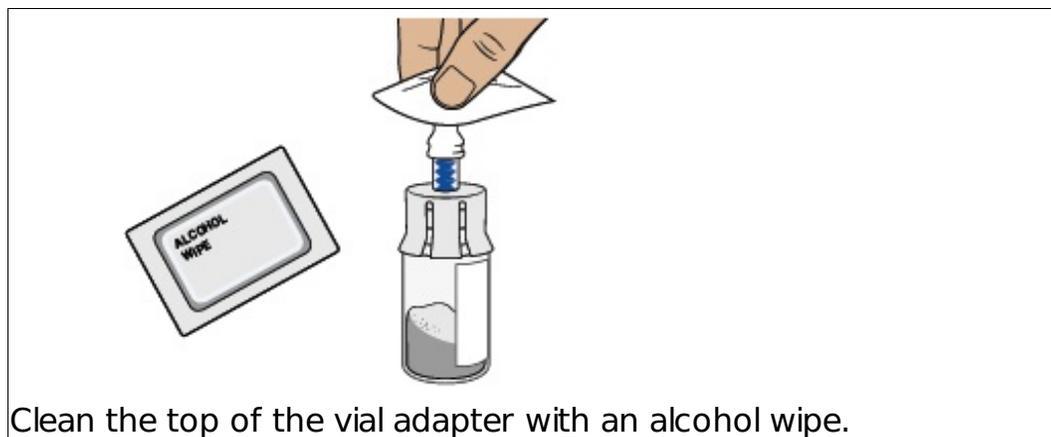
Hold the blue neck of the vial adapter and align the vial adapter on top of the vial.

package and remove vial adapter from its package. ⚠ Do not touch the inside of the vial adapter.

5b Attach vial adapter to vial

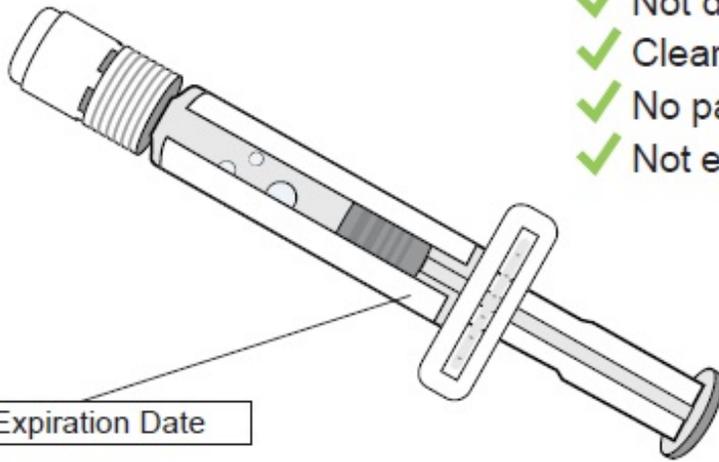


5c Clean vial adapter



6 Check prefilled syringe

Confirm the sterile water inside the prefilled syringe is clear and the product is not expired or leaking.



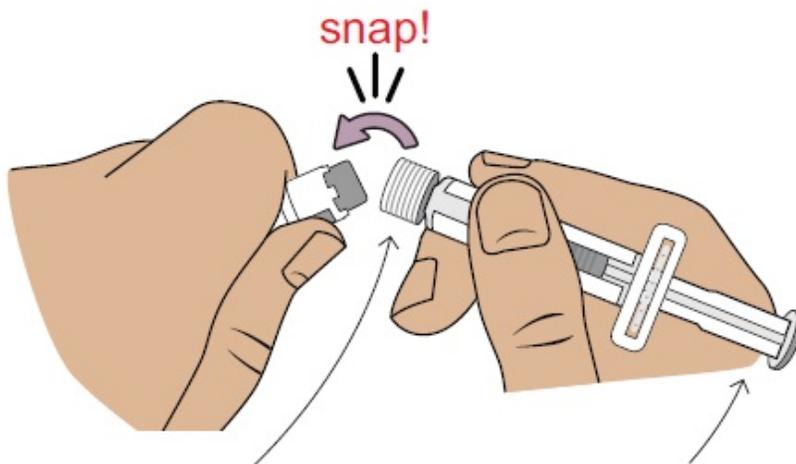
- ✓ Not damaged?
- ✓ Clear?
- ✓ No particles?
- ✓ Not expired?

Expiration Date

⚠ Do not use if you see any clumps, particles, discoloration, or product is expired. You might see air bubbles. This is okay.

7 Snap off prefilled syringe white cap

Snap off the prefilled syringe cap along the perforation.



⚠ Do not touch the tip of the prefilled syringe.

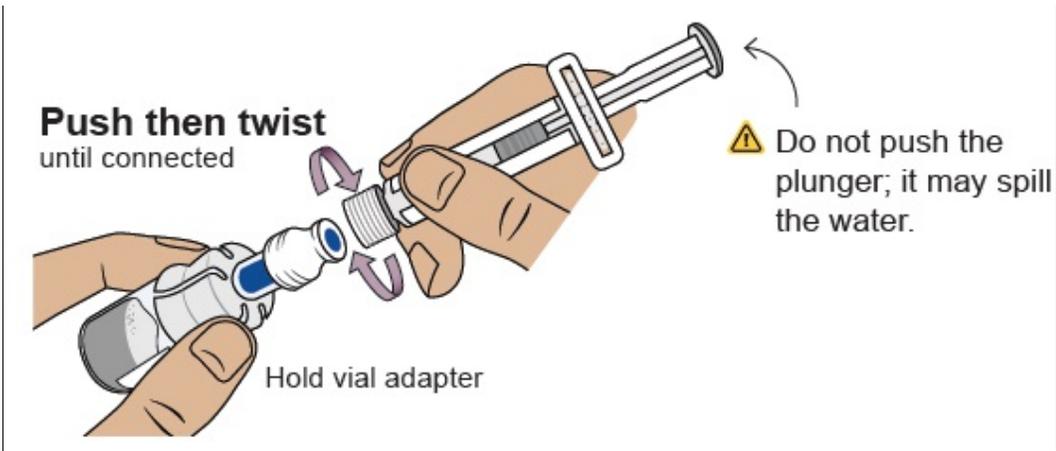
⚠ Do not push the plunger; it may spill the water.

8 Connect prefilled syringe to vial adapter

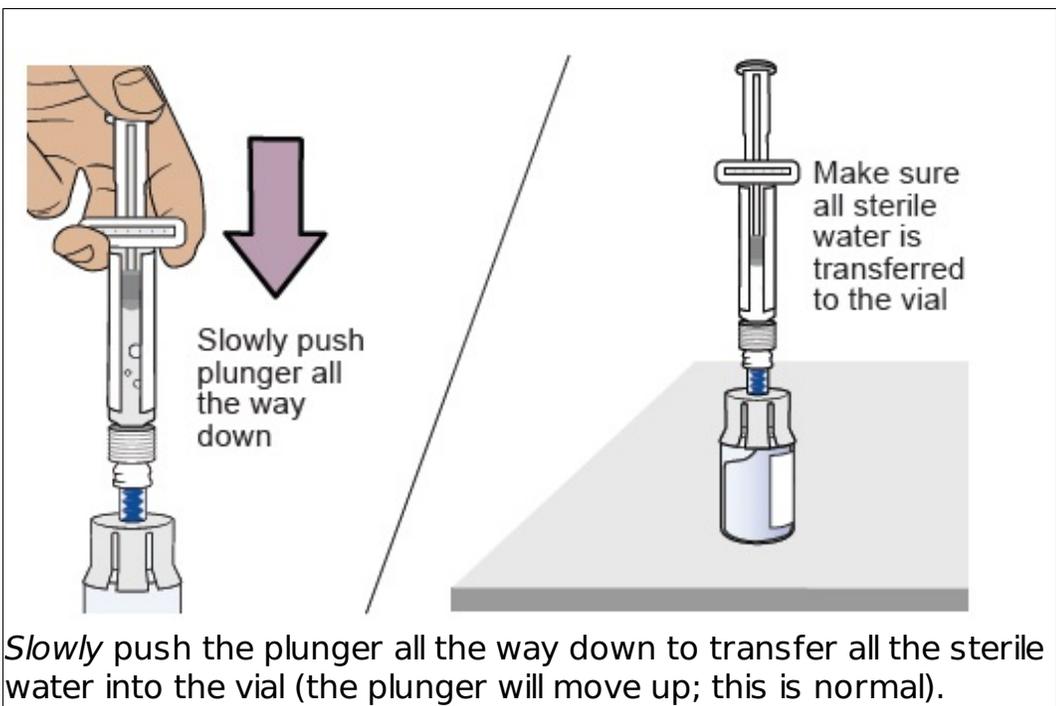
Pick up the vial with the vial adapter attached.

Align the prefilled syringe tip on the blue circle of the vial adapter.

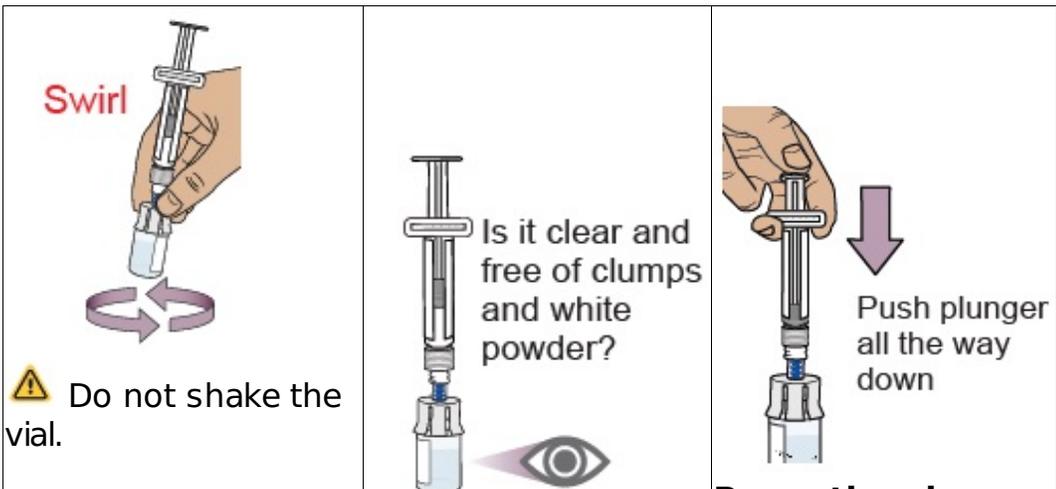
Push then twist the prefilled syringe onto the vial adapter until you cannot turn further. While twisting, be sure to hold onto the vial adapter.



9 Transfer sterile water from prefilled syringe to vial



10 Swirl to mix medicine



Hold the prefilled syringe and **gently swirl the vial in a circular motion** until the powder is fully dissolved. This may take up to **2 minutes**.

When the medicine is mixed well, it should be clear. If not, repeat this step until it is clear.

Press the plunger down to make sure all the liquid is in the vial (the plunger will move up; this is normal).

11 Wait for large bubbles in the vial to go away



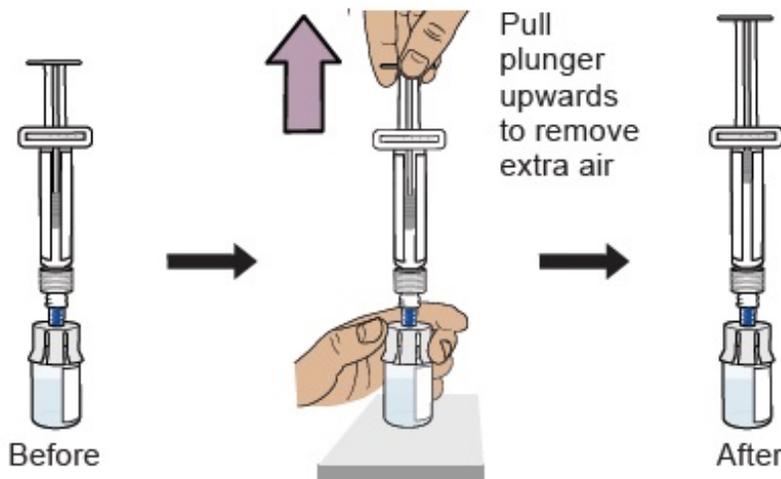
Set vial aside for large bubbles to go away.

This may take up to 3 minutes.

- ⚠ Before you continue, make sure the medicine in the vial:
- ✓ Is clear to slightly pearly and colorless to slightly brownish-yellow
 - ✓ Does not have clumps or powder
 - ✓ Does not have large bubbles

It is okay to have slight foam around edges of the vial

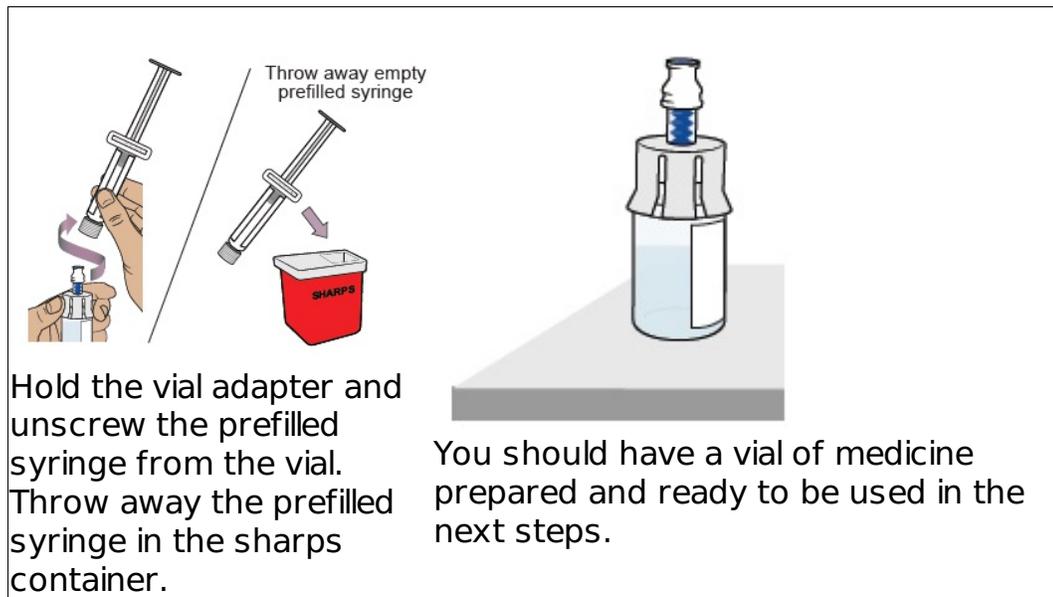
12 Prepare vial by removing extra air



With the vial standing upright, **gently pull the plunger upwards** to the top of the barrel. Do not pull the plunger out of the syringe.

Note: Pulling extra air out of the vial will help to make sure you have the right dose.

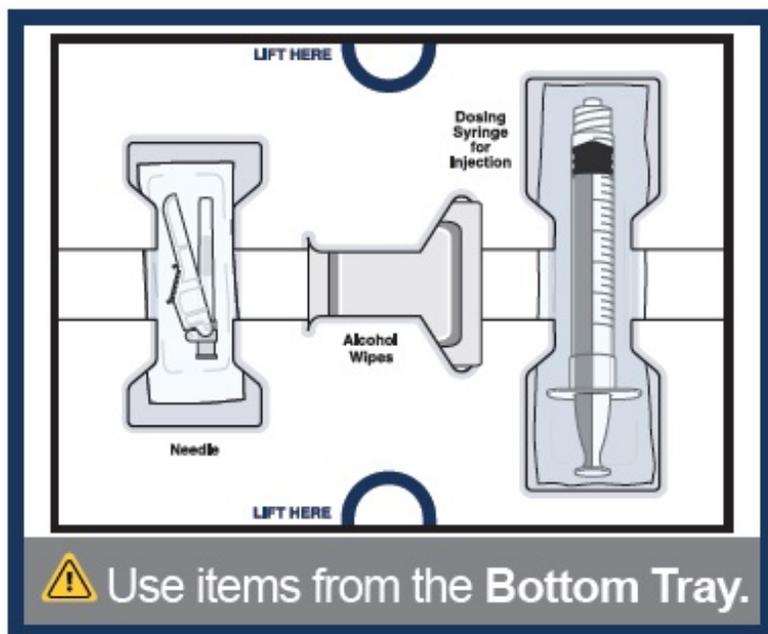
13 Remove prefilled syringe from vial



Withdraw your prescribed dose

For the next steps, you will need:

- Mixed vial of medicine
- Items from the Bottom Tray (needle, dosing syringe for injection, and alcohol wipes).



14 Clean top of vial adapter

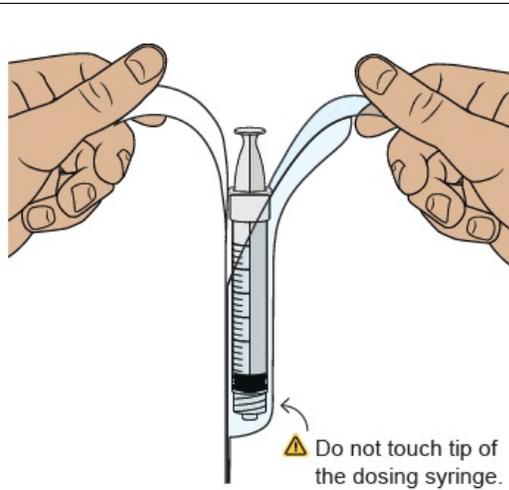
With a new alcohol wipe from the bottom tray, clean the top of the vial adapter.



15 Remove empty dosing syringe from its package

Find the empty dosing syringe in the bottom tray and remove it from its package.

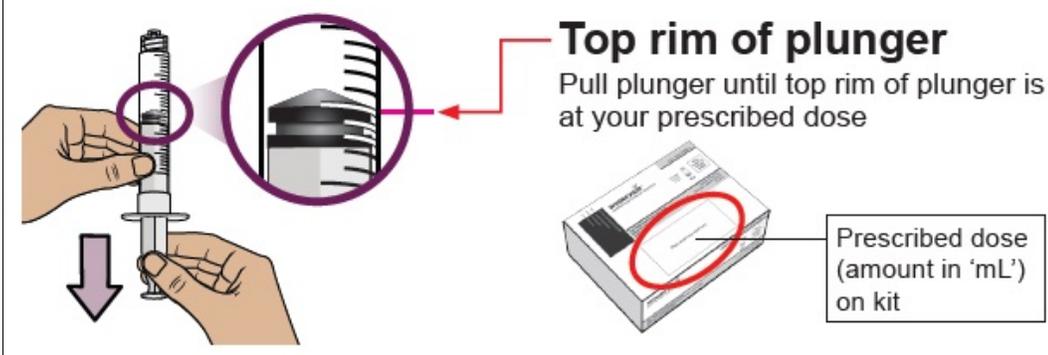
You will use this dosing syringe to measure out the medicine you need (based on your prescribed dose).



16 Pull air into the dosing syringe

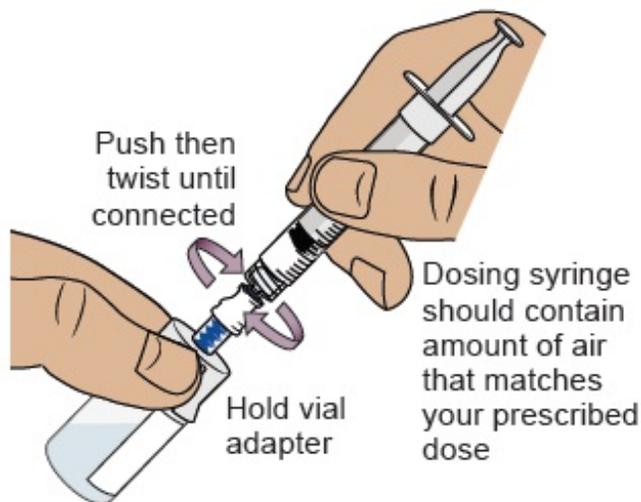
⚠️ You must do this to make sure pressure in the vial is even and you get an accurate dose.

Hold the dosing syringe upright and pull down the plunger to **draw air into the dosing syringe. Stop when top rim of plunger is at the amount in 'mL' that matches your prescribed dose.** Your prescribed dose is on the prescription label on your kit.



17 Connect dosing syringe to the vial

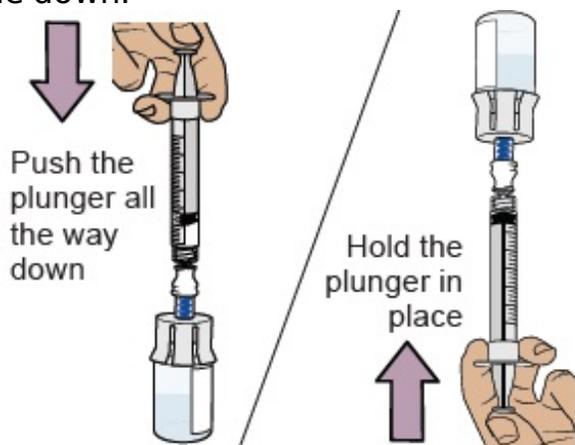
While holding the vial adapter, push then twist the dosing syringe onto the vial adapter until it stops.



18 Push air into vial, then flip vial upside down

Push the plunger all the way down to transfer all the air into the vial.

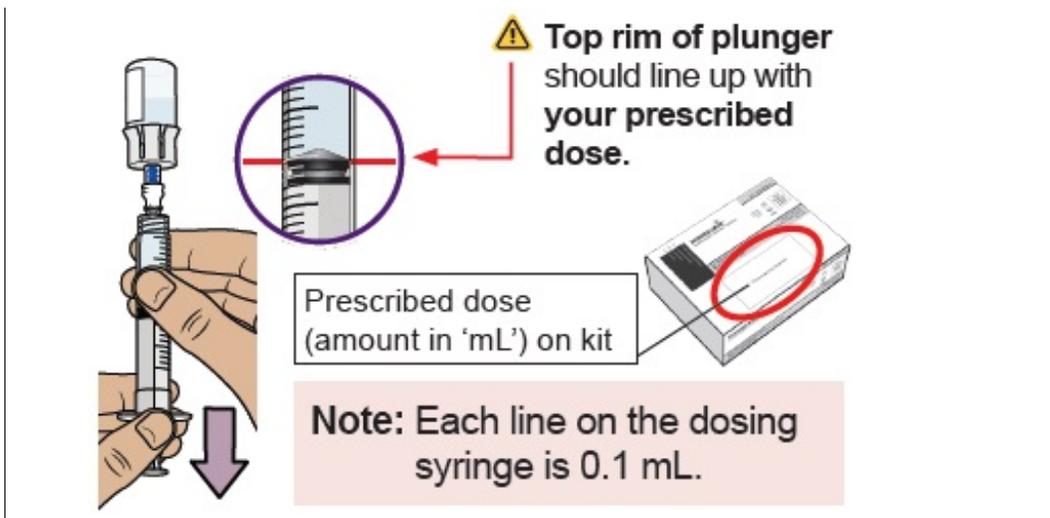
Then **hold the plunger in place** with your thumb and flip the vial upside down.



19 Pull plunger back to withdraw your prescribed dose

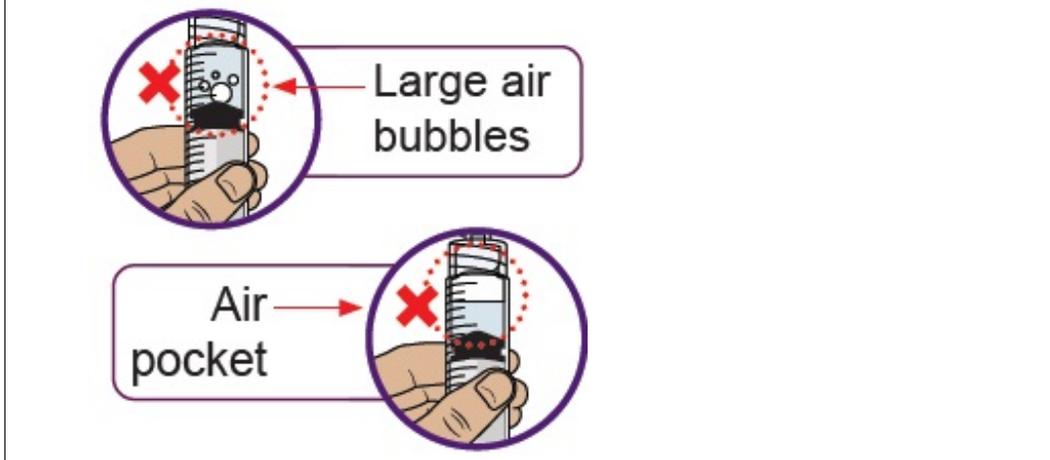
With the vial and dosing syringe upside down, **slowly pull the plunger back**.

Stop when you get to the **amount in 'mL' that matches your prescribed dose**.



20 Check for large air bubbles and air pockets

Check to see if there are large air bubbles or an air pocket in the syringe. You will remove extra air in the next steps.



21 Remove air bubbles and air pockets from syringe

If you see large air bubbles or an air pocket, tap the side of the dosing syringe to move the air to the top.

Slowly push the plunger up to remove extra air.



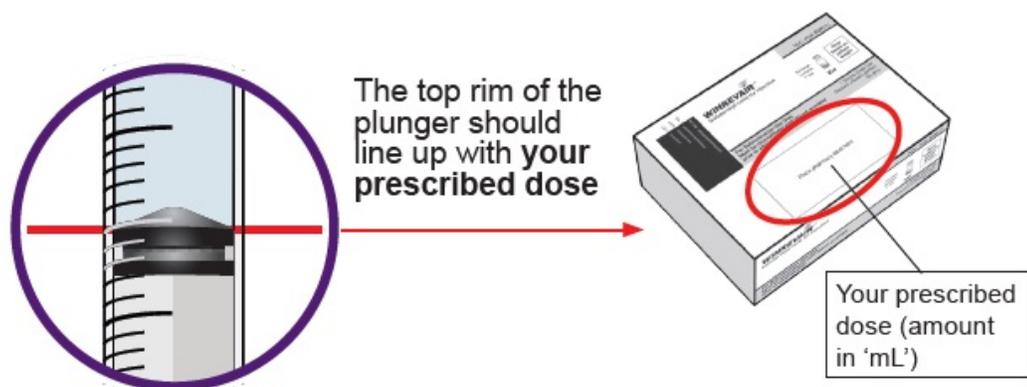
22 Compare amount to prescribed dose

After removing all extra air, **compare the amount in your syringe to your prescribed dose.** If you do not have your prescribed dose in your syringe, **slowly** pull the plunger back again to withdraw more medicine from the vial. Repeat Steps 19 to 21 until you get **your prescribed dose** and no large bubbles or air pockets remain.



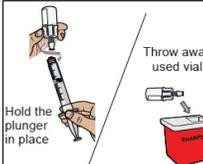
23 Confirm your prescribed dose

Before you continue, check to make sure you have the prescribed dose in the dosing syringe.



⚠️ If the amount does not match your prescribed dose, repeat Steps 19 to 22.

24 Remove the dosing syringe from the vial and set dosing syringe aside



Hold the plunger in place with one hand. With the other hand, hold the vial adapter and unscrew the filled dosing syringe from the vial.

Throw away used vial

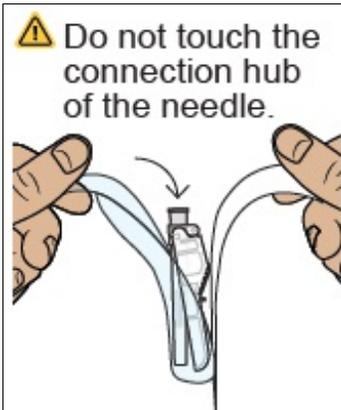
Throw away the used vial into the sharps container.



Place the filled dosing syringe on a clean, flat surface.

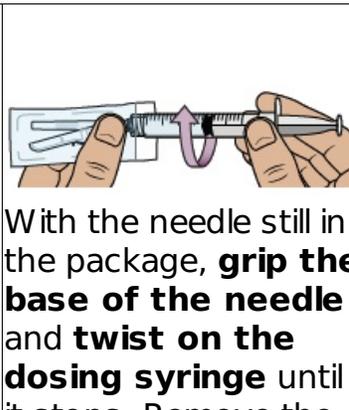
⚠ Do not touch the dosing syringe tip or let it to touch any surfaces.

25 Attach injection needle

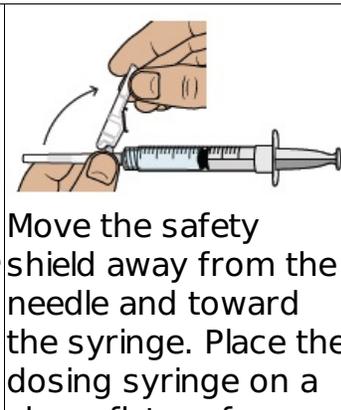


⚠ Do not touch the connection hub of the needle.

Find the needle in the bottom tray and open its package.



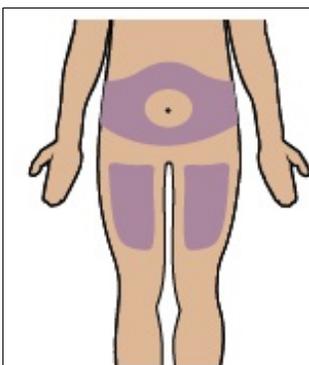
With the needle still in the package, **grip the base of the needle** and **twist on the dosing syringe** until it stops. Remove the needle package.



Move the safety shield away from the needle and toward the syringe. Place the dosing syringe on a clean, flat surface.

⚠ Do not uncap the needle.

26 Choose and clean your injection site



Select an injection site on your stomach (abdomen) or your upper thigh. If injecting on your stomach, avoid a 2-inch area around your belly button.

Choose a different site every time you inject.

⚠ Do not inject into skin that is damaged, sore, tender, bruised, or has red patches.

⚠ Do not inject through clothes.



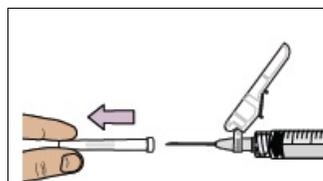
Clean the injection site with a new alcohol wipe.

⚠ Do not touch the cleaned injection site again.

Now, you are ready to inject the medicine.

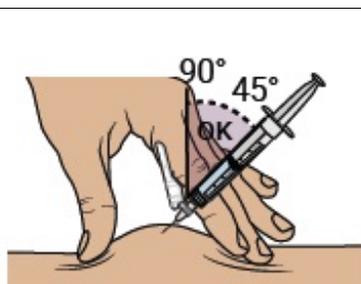
Inject your medicine

27 Inject your medicine



Pull the cap straight off the needle. Throw away the cap.

⚠ Do not touch the plunger until ready to inject so you do not lose any medicine.



Gently **pinch and hold a fold of skin** where you will inject. Insert the needle with a **dartlike motion at a 45° to 90° angle**.



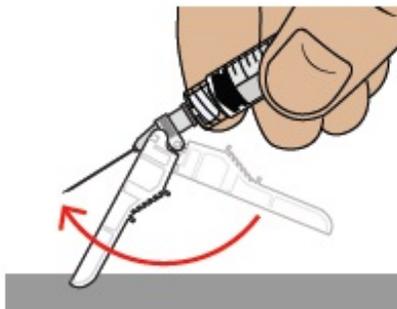
Push the plunger with slow, steady pressure all the way down until the dosing syringe is empty. **Confirm all the medicine has been injected.** You can let go of the skin fold now.



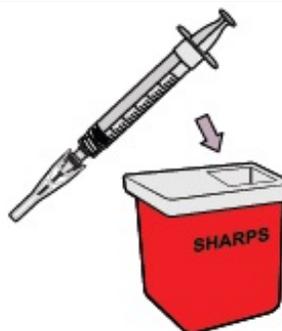
Keep your fingers away from the needle at all times.



While keeping the plunger pushed in, **remove the needle from your skin** at the same angle you inserted it.



To reapply the safety shield, push the shield against a flat surface until you hear a “click” and see that the needle is covered.



Throw away the dosing syringe and used items in a sharps disposal container.

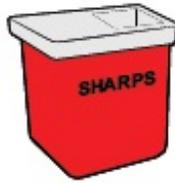
⚠ Do not remove the needle from the dosing syringe.

How to throw away WINREVAIR

- Throw away any used vials (including any remaining WINREVAIR liquid), needle, vial and needle caps, and used syringes in an FDA-cleared sharps

disposal container.

- Do not throw away the WINREVAIR vials, syringes, or needle in your household trash.
- **Do not reuse any of the supplies.** WINREVAIR is for one time use only.
- **Important:** Always keep the sharps disposal container out of reach of children and pets.



If you do not have an FDA-cleared sharps disposal container, you may use a household container that is:

- made of a heavy-duty plastic,
- can be closed with a tight-fitting, puncture-resistant lid, without sharp objects being able to come out,
- upright and stable during use,
- leak resistant, and
- properly labeled to warn of hazardous waste inside the container.

When your sharps disposal container is almost full, you will need to follow your community guidelines for the right way to throw away your sharps disposal container.

There may be state or local laws about how you should throw away used needles, dosing syringes, and prefilled syringes.

For more information about safe sharps disposal, and for specific information about sharps disposal in the state that you live in, go to the FDA's website at:
<http://www.fda.gov/safesharpsdisposal>.

 Do not recycle your used sharps container.

Frequently asked questions

What should I do if I am bleeding at the injection site?

Place a cotton ball or bandage on your skin right away and apply a small amount of pressure. If the bleeding does not stop, call your healthcare provider right away.

Where can I find my prescribed dose?

Your prescribed dose in 'mL' is on your prescription label on your kit. Contact your healthcare provider if you cannot find your prescribed dose.

What should I do if I accidentally get some medicine on my skin or my work surface?

Wash the area thoroughly with soap and water right away.

What should I do if I am not sure I administered my prescribed dose correctly?

Call your pharmacy or healthcare provider.

What should I do if the plunger of my dosing syringe moves automatically when I try to withdraw medicine from the vial?

Do not worry if your plunger moves slightly on its own when you are filling your dosing syringe with medicine.

With one hand, **hold the plunger in place to stop the plunger from moving.** With the other hand, unscrew the vial from the dosing syringe.

After unscrewed, it is safe to let go of the plunger.

You can avoid this automatic plunger movement by pushing air into the vial before filling your dosing syringe with medicine. Refer to Steps 16 to 23 for detailed instructions.

What should I do if my kit parts are damaged or are discolored, cloudy, or have particles?

Do not use your kit. Call your pharmacy to get a new kit.

What should I do if my medicine does not turn clear after mixing and swirling?

Do not use the medicine if you have swirled the medicine vial for about 2 minutes and let it stand for another 3 minutes but your medicine vial remains cloudy or has clumps, powder, or foreign particles. Call your pharmacy to get a new kit.

What should I do if the sterile water will not come out of the prefilled syringe?

Check that the vial adapter is attached to the vial securely. If not, hold the vial and press the vial adapter down firmly to make sure the vial adapter punctures the vial rubber stopper.

What should I do if I dropped my kit components?

Do not use if any items are damaged. If you are unsure, call your pharmacy to get a new kit.

Can I use my kit that has been left out of the refrigerator?

Store the injection kit in the refrigerator until ready to use it. If the unused injection kit has been out of the refrigerator up to 77°F (25°C) for more than 24 hours, please contact your pharmacy or healthcare provider.

Do I need to use mixed medicine right away?

We recommend you inject the medicine right away after mixing but no later than 4 hours after mixing. If it has been more than 4 hours, throw away unused mixed medicine. If you have questions, please contact your pharmacy or healthcare provider.

How can I get help preparing and giving my injection?

If you have questions about how to give WINREVAIR the correct way or need more information, you can call your pharmacy or healthcare provider.

Call your doctor for advice about side effects.

To report side effects or product quality complaints, please call Merck Sharp & Dohme LLC at 1-800-672-6372. You can also report side effects to FDA at 1-800-FDA-1088.

Manufactured by:
Merck Sharp & Dohme LLC
Rahway, NJ 07065, USA
U.S. License Number 0002

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This Instructions for Use has been approved by the U.S. Food and Drug Administration
Approved: March 2024

Important: Read this booklet first Important

INSTRUCTIONS FOR USE Mix

WINREVAIR™ (WIN-reh-vair) Combine
(sotatercept-csrk)

for injection, for subcutaneous use

How to use your WINREVAIR injection kit



Withdraw

For package containing 2 vials

For subcutaneous injection only (inject Inject
directly under the skin)

In this booklet:

Important information for patients or caregivers

Before you start

Get to know the parts of your kit

Important information you need to know before injecting WINREVAIR

Storing your injection kit

Get started

● **Mix** powdered medicine into liquid form

● **Combine** medicine from both vials

● **Withdraw** your prescribed dose

○ **Inject** your medicine

How to throw away WINREVAIR

Frequently asked questions



Important information for patients or caregivers

Before you prepare and inject WINREVAIR, you must be:

- trained by a healthcare provider following this Instructions for Use booklet step-by-step, and
- capable of independent administration of WINREVAIR as determined by a healthcare provider

Before using WINREVAIR, make sure you are trained to do the following correctly:

1. Reconstitute (mix) the medicine
2. Combine medicine from both vials
3. Measure the correct amount of medicine according to your prescription
4. Select and prepare a proper injection site
5. Inject the medicine subcutaneously

Before you start:



Read this booklet

Read this Instructions for Use from start to finish before using WINREVAIR and each time you get a refill. There may be new information.



Start with your healthcare provider

Do not use WINREVAIR until your healthcare provider has shown you or your caregiver the right way to prepare and inject it. Your healthcare provider should show you how to inject WINREVAIR before you use it for the first time.



Questions?

If you have questions about how to give WINREVAIR the right way or need more information, call your pharmacy or healthcare provider.

For general product information visit www.WINREVAIR.com or call Merck Sharp & Dohme LLC at 1-800-672-6372.

How to use WINREVAIR

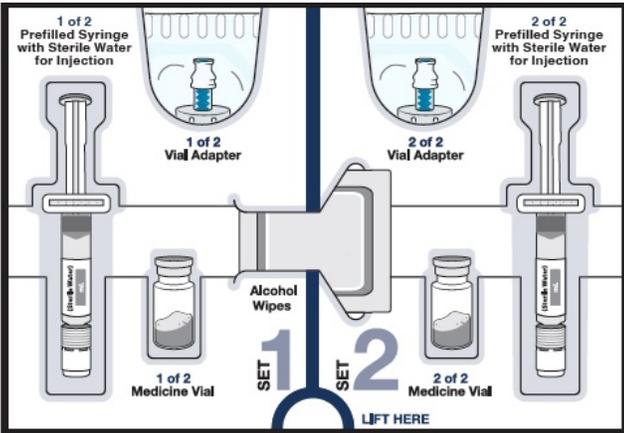
Before injecting WINREVAIR, make sure you read the full instructions in this booklet to get your prescribed dose.

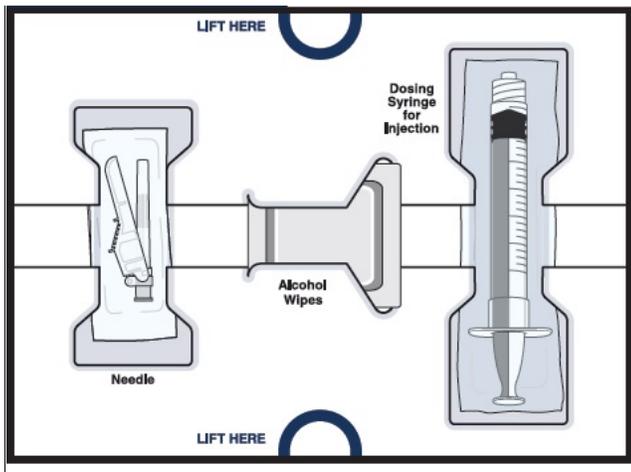
● **Mix** medicine in vial 1

- **Mix** medicine in vial 2
- **Combine** medicine mixture from vial 1 to vial 2
- **Withdraw** your prescribed dose from vial 2
- Then, **inject** your medicine

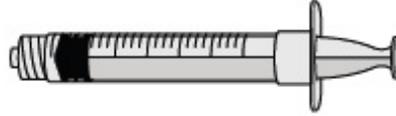
Proceed to the next pages for step-by-step directions.

Get to know the parts of your kit

Top Tray	Use to mix the medicine
	 <p>Vial Medicine powder</p> <p>2 Vials of WINREVAIR medicine powder</p>  <p>2 Prefilled syringes with sterile water for injection to mix medicine powder into liquid form</p>
	 <p>2 Vial adapters to connect the vial and syringe</p>
Bottom Tray	Use to inject the medicine



Needle for injection



Empty dosing syringe to measure, withdraw, and inject your medicine

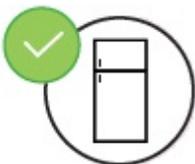
Important information you need to know before injecting WINREVAIR

- You must mix WINREVAIR before using it. Make sure the medicine powder in the vial is completely dissolved before you inject.
- **Check your prescribed dose (amount in 'mL') each time you use WINREVAIR. Your prescribed dose may change.** Your prescribed dose is on the prescription label on your kit.
- Use only the supplies that are in the kit to prepare your prescribed dose.
- Do not open the package or mix the medicine until you are ready to use it.
- **Do not reuse any of the supplies.** After your injection, throw away the used vials with any remaining WINREVAIR medicine, needle, caps, and syringes in a sharps container. See pages 38-39 for more information.

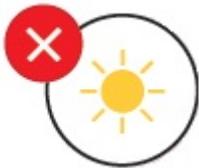


Prescribed dose

Storing your injection kit



Store WINREVAIR injection kit in the refrigerator at 36°F to 46°F (2°C to 8°C). Do not freeze.



Store in the original carton to protect from light.



Keep injection kit out of the reach of children and pets.

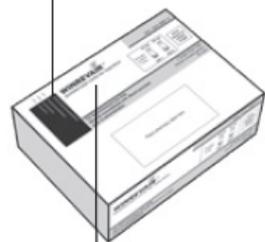
Get started

Before you can prepare and inject WINREVAIR, you must first be trained and determined to be capable of independent administration of WINREVAIR by a healthcare provider.

1 Check WINREVAIR injection kit and expiration date

Remove the WINREVAIR injection kit from the refrigerator.

EXPIRATION DATE



WINREVAIR

✓ **Check the expiration date and look for any signs of damage** on the kit or on the supplies.

⚠ If expired or damaged, do not use. Call your pharmacy to get a new kit.

✓ **Check that you have the medicine** that your healthcare provider prescribed.

2 Let your kit come to room temperature, gather supplies, and wash your hands

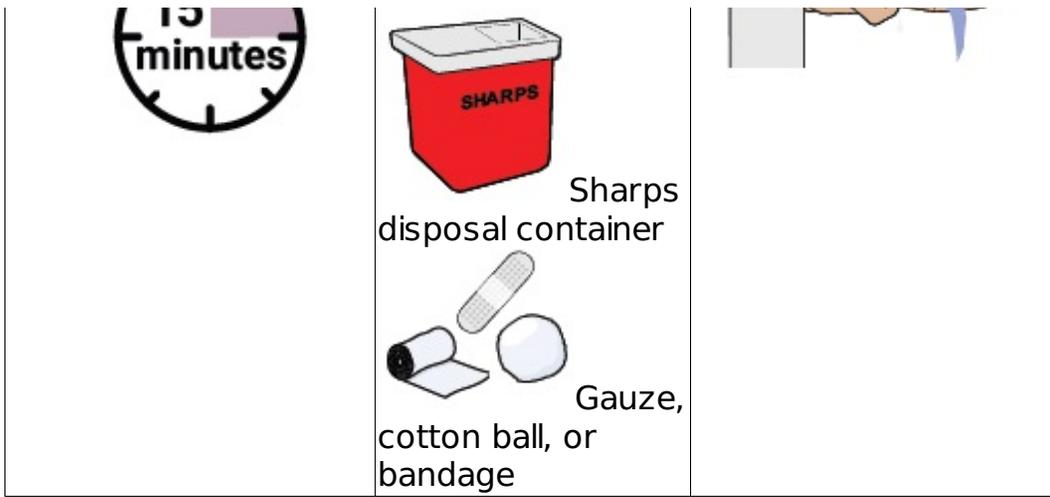
Wait 15 minutes to allow your kit to warm to room temperature. Cold medicine is more painful to inject.



Along with your kit, gather these items and find a clean, flat surface where you will prepare and inject your dose.

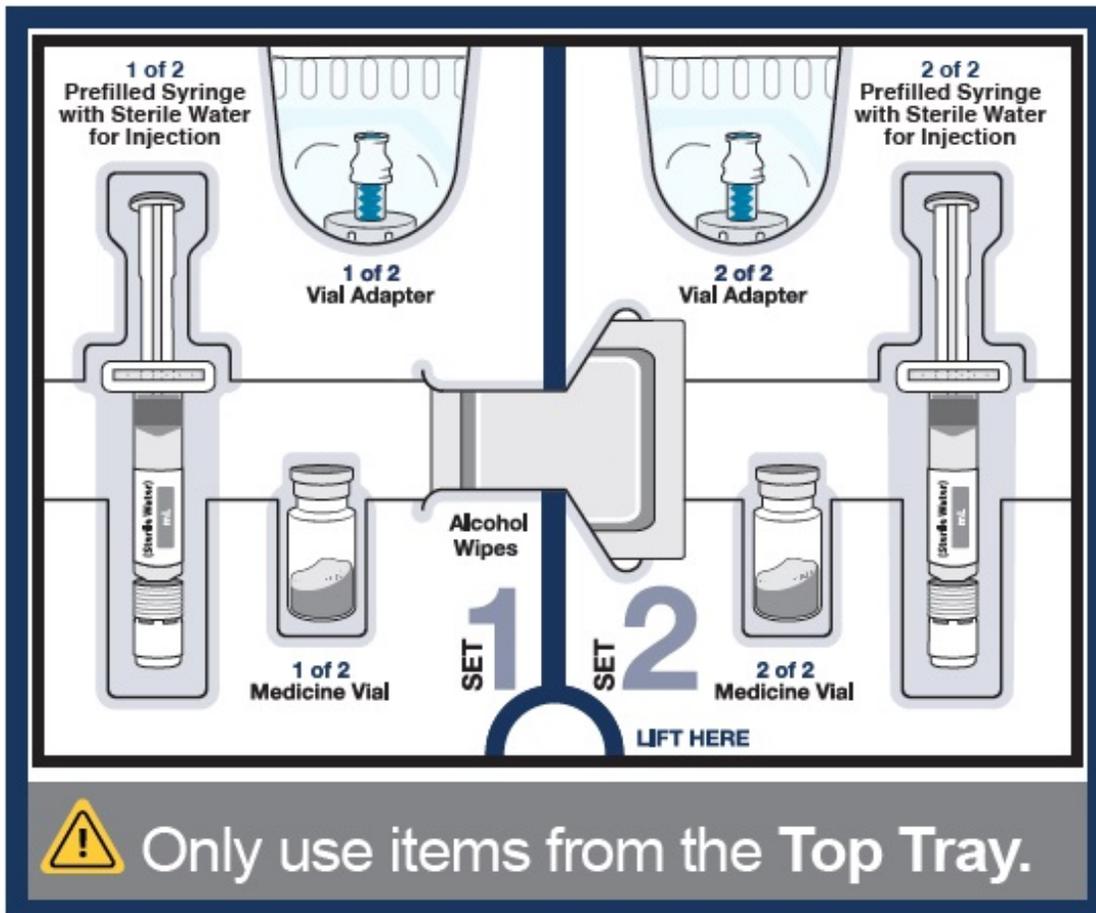
Wash your hands with soap and water.



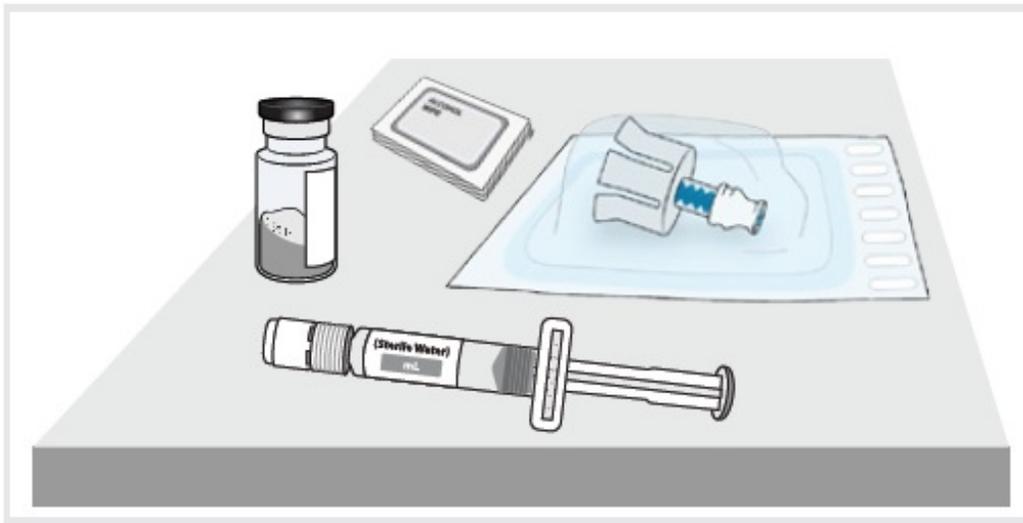


Mix powdered medicine into liquid form

Start with the Top Tray



3 Remove medicine vial 1, prefilled syringe with sterile water for injection 1, alcohol wipes, and vial adapter 1 from the kit.



4a Check medicine vial

Vial
Medicine powder

✓ Not damaged?
✓ No colored particles?
✓ Not expired?

EXPIRATION DATE

Check the vial label to **confirm the medicine is not expired**.
Inspect the medicine powder.
It should be white to off-white.

⚠ Do not use if expired, damaged, discolored, or you can see particles in it.

4b Remove plastic cap and clean vial

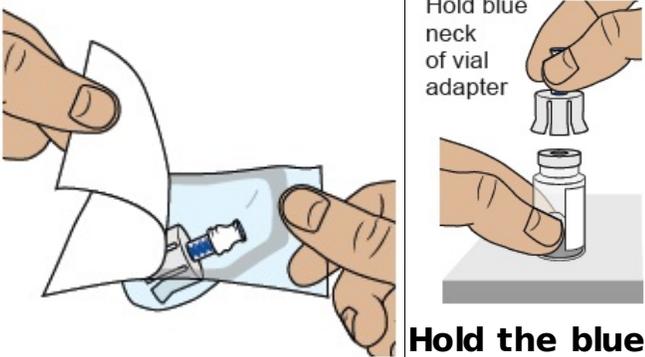
Rubber stopper

Flip off the plastic cap and **clean the rubber stopper on top of the vial** with an alcohol wipe.

⚠ Do not use if vial cap is missing.
⚠ Do not touch the cleaned rubber stopper.

Set vial aside on a clean, flat surface.

5a Align vial adapter to vial



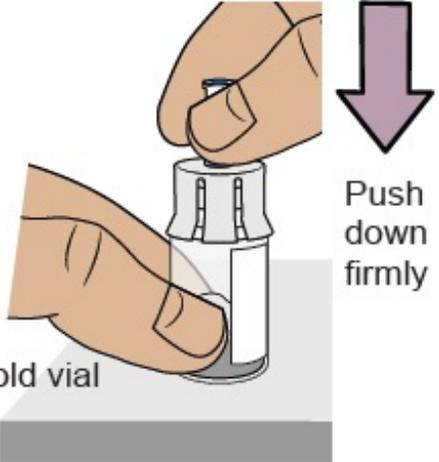
Peel open the vial adapter package and remove vial adapter from its package.

Hold blue neck of vial adapter

To attach the vial adapter:
Hold the blue neck of the adapter
Do not hold bottom of the adapter

Hold the blue neck of the vial adapter and align the vial adapter on top of the vial. ⚠️ Do not touch the inside of the vial adapter.

5b Attach vial adapter to vial

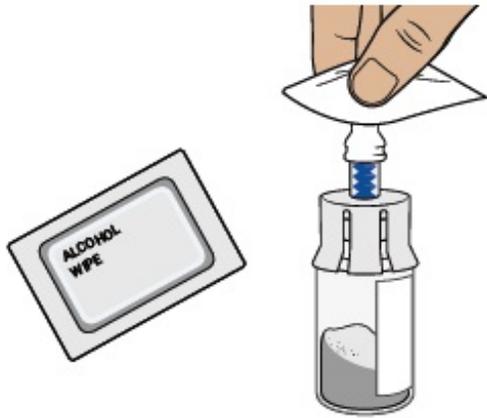


Hold vial

Push down firmly

Hold the vial with one hand. **Push down firmly** so it snaps in place (you may feel some resistance).

5c Clean vial adapter



Clean the top of the vial adapter with an alcohol wipe.

6 Check prefilled syringe

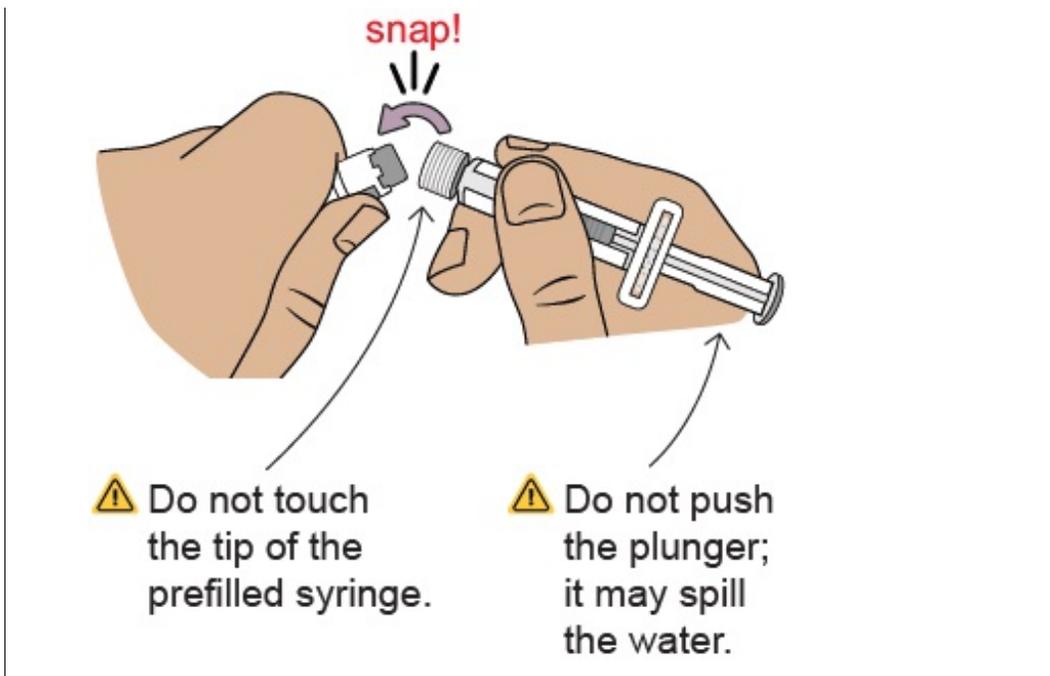
Confirm the sterile water inside the prefilled syringe is clear and the product is not expired or leaking.



 Do not use if you see any clumps, particles, discoloration, or product is expired. You might see air bubbles. This is okay.

7 Snap off prefilled syringe white cap

Snap off the prefilled syringe cap along the perforation.

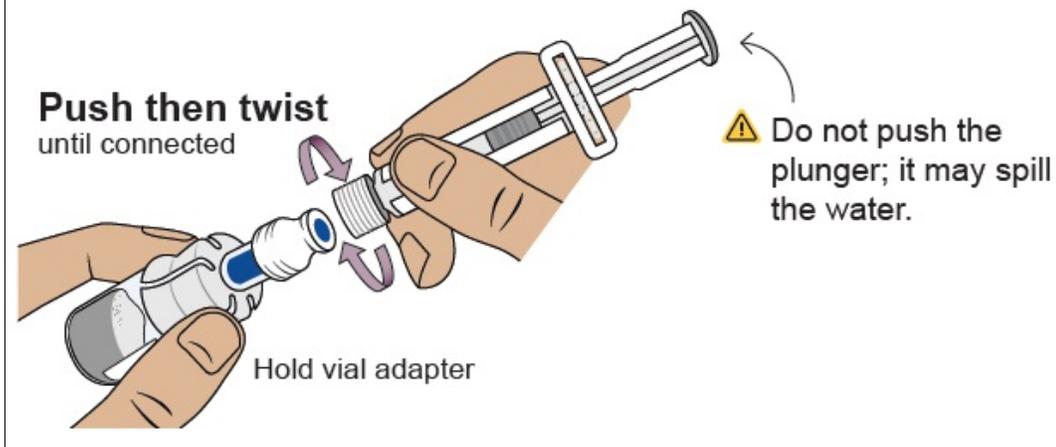


8 Connect prefilled syringe to vial adapter

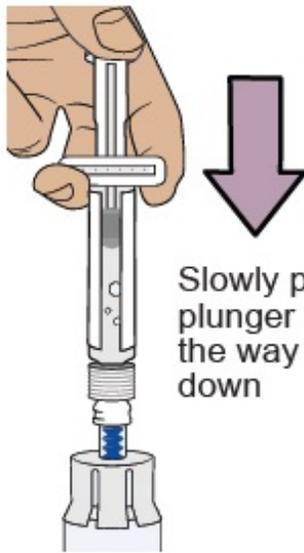
Pick up the vial with the vial adapter attached.

Align the prefilled syringe tip on the blue circle of the vial adapter.

Push then twist the prefilled syringe onto the vial adapter until you cannot turn further. While twisting, be sure to hold onto the vial adapter.



9 Transfer sterile water from prefilled syringe to vial



Slowly push plunger all the way down



Make sure all sterile water is transferred to the vial

Slowly push the plunger all the way down to transfer all the sterile water into the vial (the plunger will move up; this is normal).

Leave the prefilled syringe attached to the vial and set aside on a flat surface.

10 Prepare vial 2, prefilled syringe 2, and vial adapter 2

Remove vial 2, prefilled syringe 2, and vial adapter 2 from Set 2 in the kit.



Important:

Turn back to Page 13 and repeat Steps **4** to **9** to prepare vial 2.

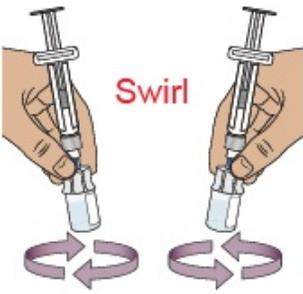
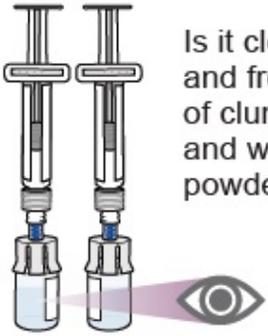
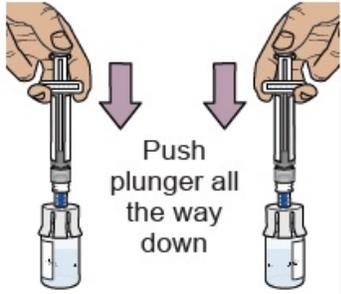


Stop and check to make sure you have both vial 1 and vial 2 prepared.

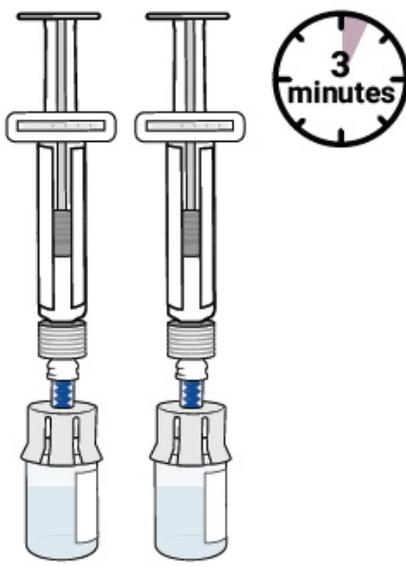


You need both vials prepared before you can continue to Step 11.

11 Swirl to mix medicine

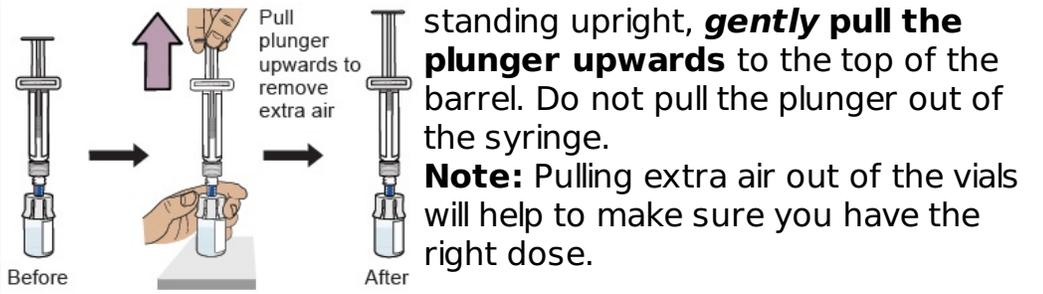
 <p>⚠ Do not shake the vials.</p> <p>Hold 1 prefilled syringe in each hand and gently swirl both vials in a circular motion until the powder is fully dissolved. This may take up to 2 minutes.</p>	 <p>Is it clear and free of clumps and white powder?</p> <p>When the medicine is mixed well, it should be clear. If not, repeat this step until it is clear.</p>	 <p>Push plunger all the way down</p> <p>For each vial, press the plunger down to make sure all the liquid is in the vial (the plunger will move up; this is normal).</p>
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12 Wait for large bubbles in the vials to go away

 <p>It is okay to have slight foam around edges of the vials</p>	<p>Set both vials aside for large bubbles to go away. This may take up to 3 minutes.</p> <p>⚠ Before you continue, make sure the medicine in the vials:</p> <ul style="list-style-type: none"> ✓ Is clear to slightly pearly and colorless to slightly brownish-yellow ✓ Does not have clumps or powder ✓ Does not have large bubbles
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13 Prepare vials by removing extra air

Start with either vial. With the vial



standing upright, **gently pull the plunger upwards** to the top of the barrel. Do not pull the plunger out of the syringe.

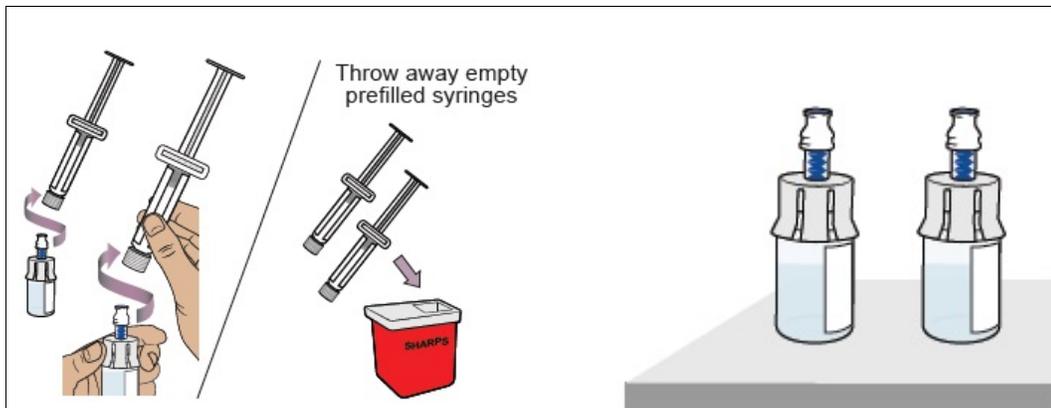
Note: Pulling extra air out of the vials will help to make sure you have the right dose.

⚠ Important:

Perform this step for **both** vials. Stop and check you have **both** vials prepared before continuing.



14 Remove prefilled syringes from vials



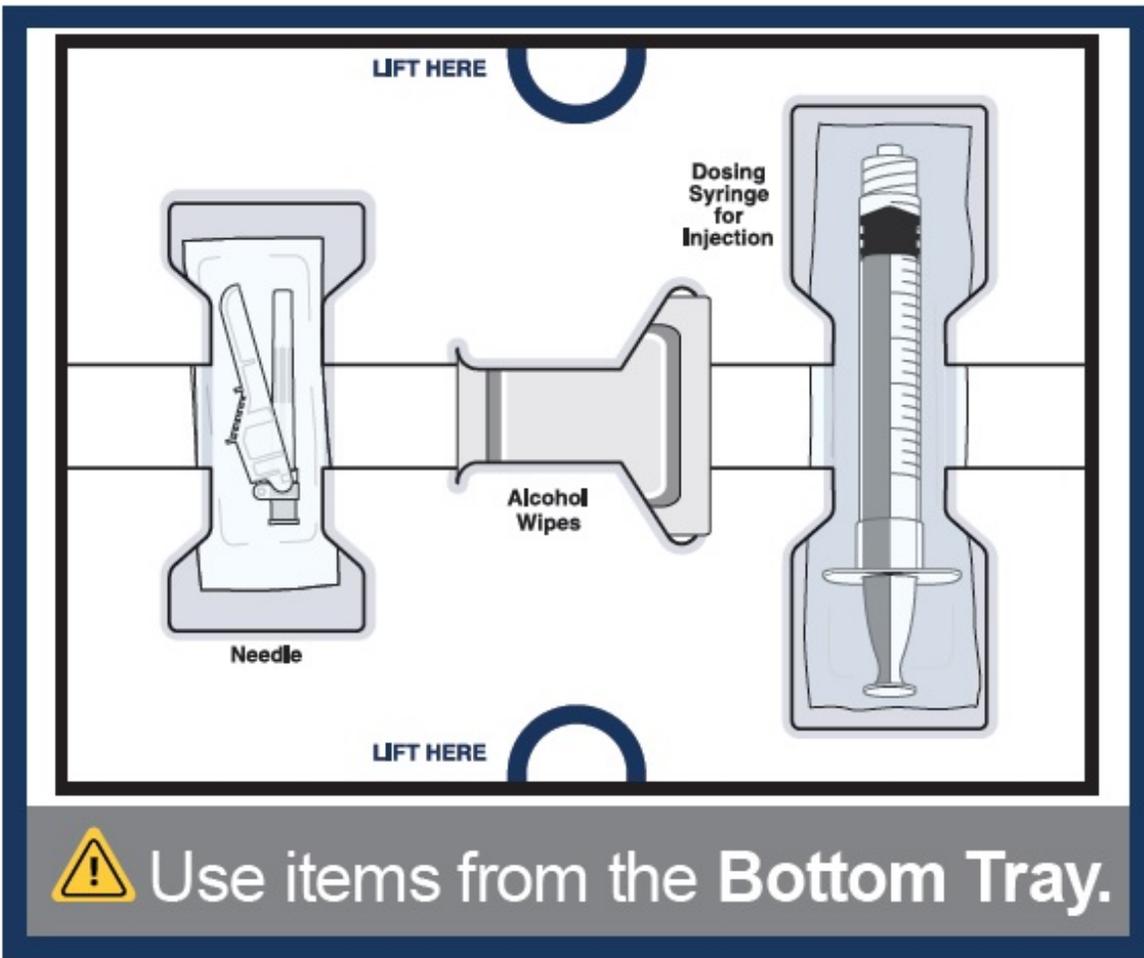
Hold the vial adapters and unscrew **both** prefilled syringes from the vials. Throw away **both** prefilled syringes in the sharps container.

You should have 2 vials of medicine prepared and ready to be combined in the next steps.

Combine medicine from both vials

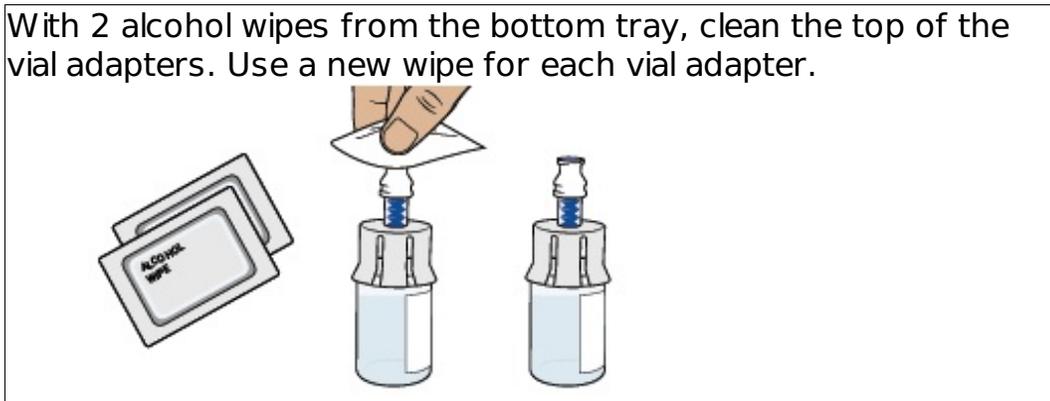
For the next steps, you will need:

- Mixed vial 1 and vial 2
- Items from the Bottom Tray (needle, dosing syringe for injection, and alcohol wipes).



15 Clean top of both vial adapters

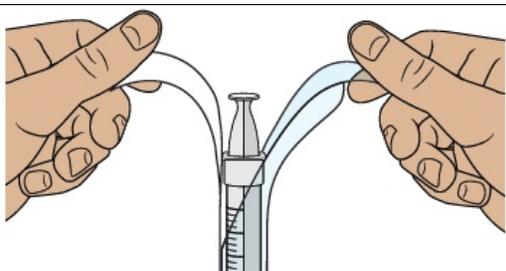
With 2 alcohol wipes from the bottom tray, clean the top of the vial adapters. Use a new wipe for each vial adapter.



16 Remove empty dosing syringe from its package

Find the empty dosing syringe in the bottom tray and remove it from its package.

You will use this dosing syringe to combine medicine from both vials and measure out the



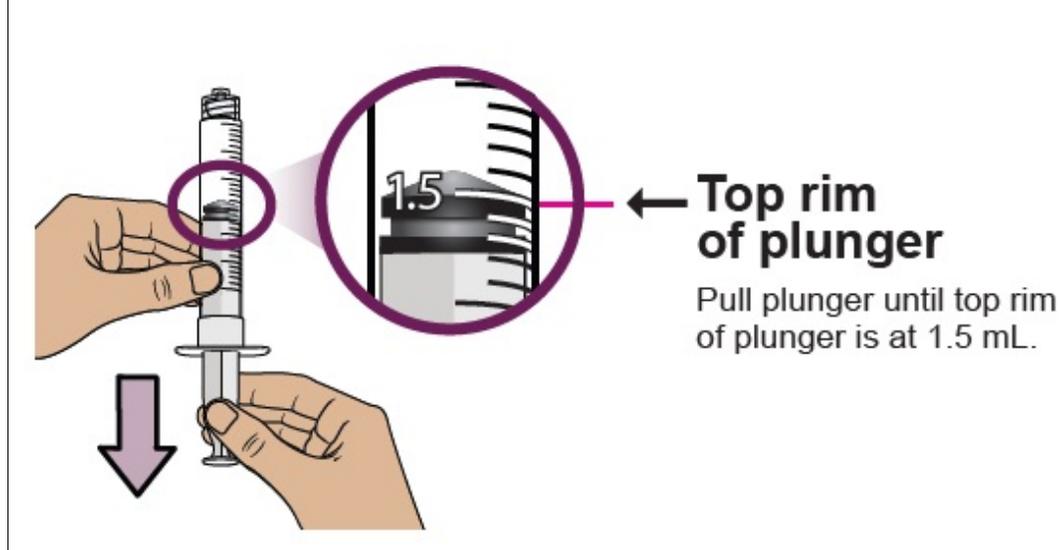
medicine you need (based on your prescribed dose).



17 Pull air into the dosing syringe

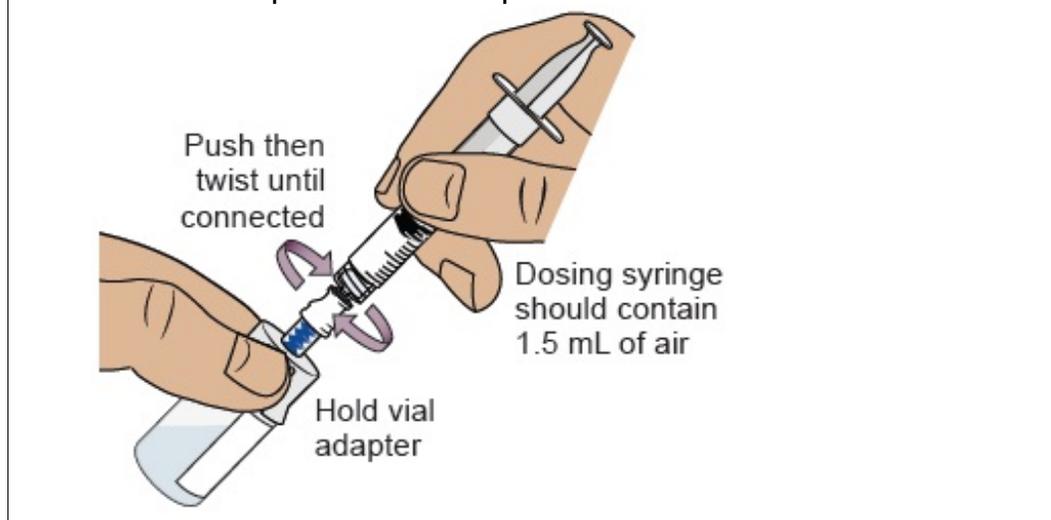
⚠ You must do this to make sure pressure in the vial is even and you get an accurate dose.

Hold the dosing syringe upright and pull down the plunger to **draw in 1.5 mL of air.**



18 Connect dosing syringe to one of the vials

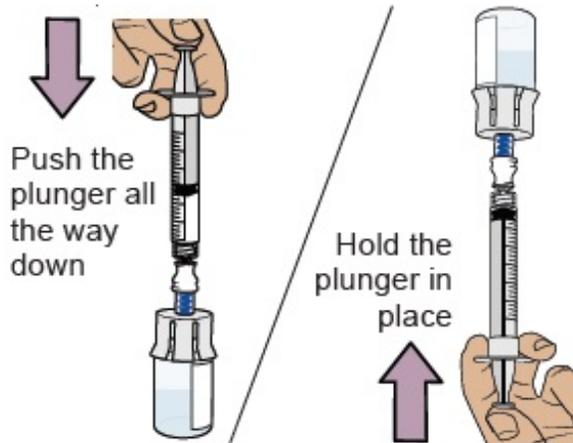
While holding the vial adapter, push then twist the dosing syringe onto the vial adapter until it stops.



19 Push air into vial, then flip upside down

Push the plunger all the way down to transfer all the air into the vial.

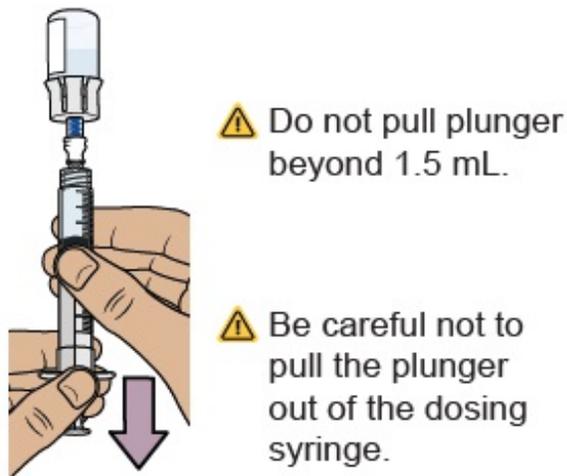
Then **hold the plunger in place** with your thumb and flip the vial upside down.



20 Withdraw all the medicine from 1st vial

With the vial and dosing syringe upside down, slowly pull the plunger back. **Stop at 1.5 mL.**

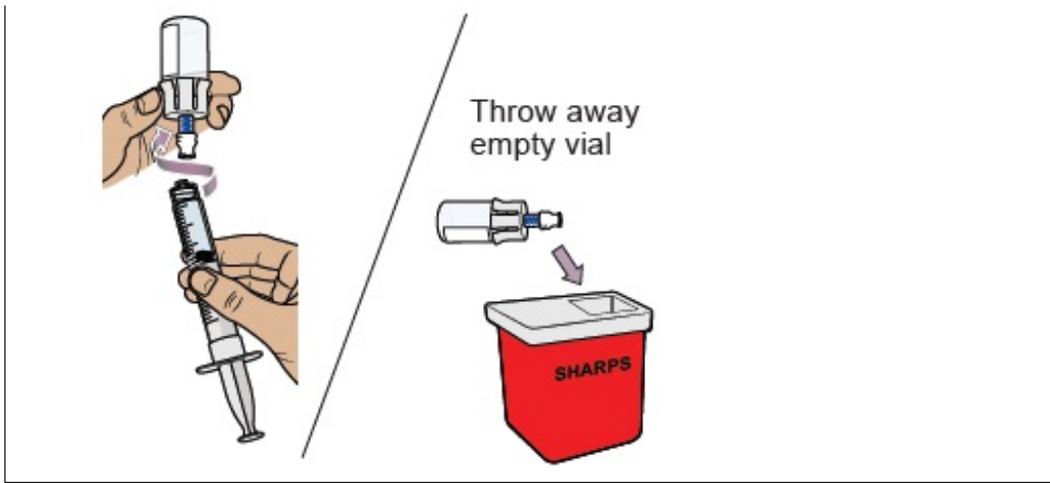
This will make sure all the medicine from the vial is in the dosing syringe.



21 Remove 1st vial from the dosing syringe

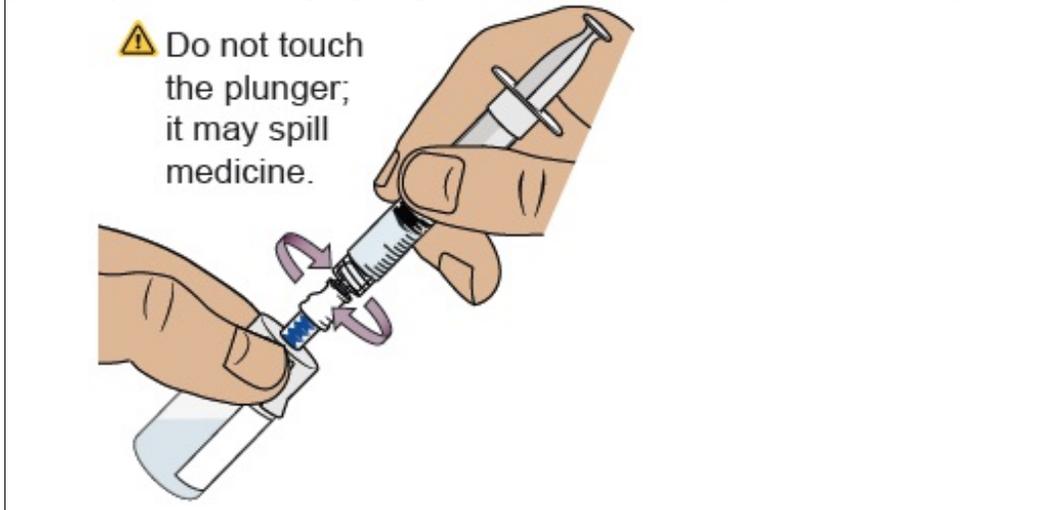
Hold the vial adapter and unscrew the filled dosing syringe from the vial.

Throw away the **empty** vial into the sharps container.



22 Connect the dosing syringe to the 2nd vial

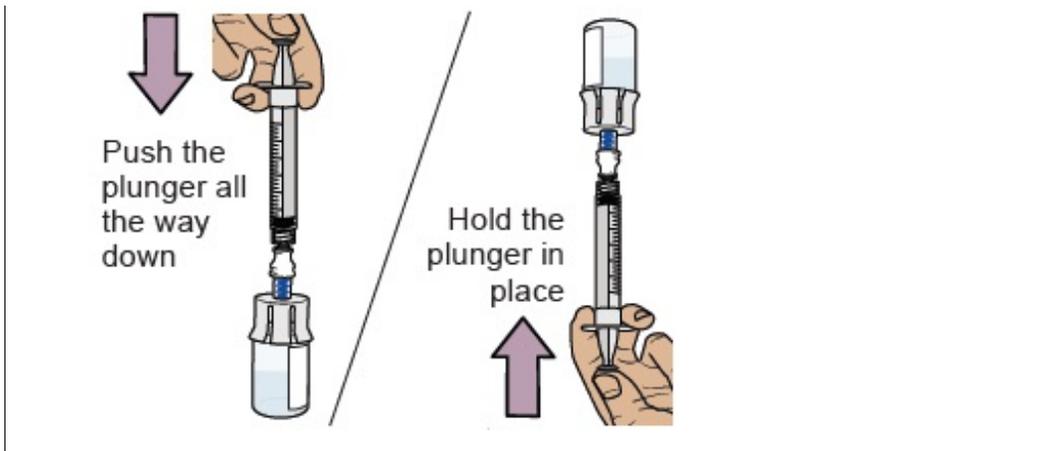
While holding the vial adapter of the 2nd vial, push then twist the partially filled dosing syringe onto the vial adapter until it stops.



23 Push all the medicine into the 2nd vial, then flip vial upside down

Slowly push the plunger all the way down to transfer **all of the medicine into the vial**. This combines the medicine from both vials.

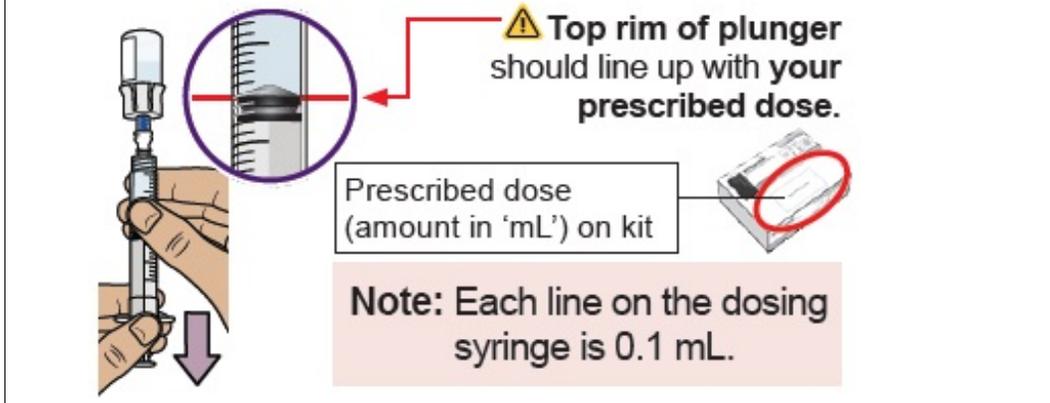
Hold the plunger in place with your thumb and flip the vial upside down.



Withdraw your prescribed dose

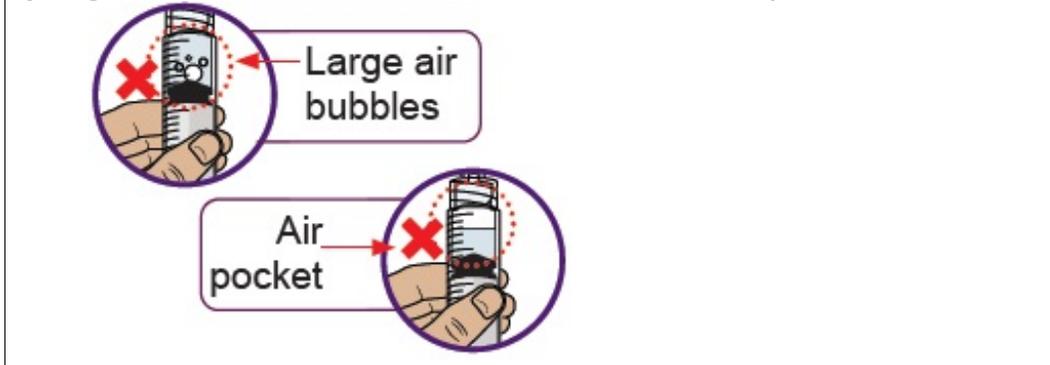
24 Pull plunger back to withdraw your prescribed dose

With the vial and dosing syringe upside down, **slowly pull the plunger back**.
Stop when you get to the **amount in 'mL' that matches your prescribed dose**.



25 Check for large bubbles and air pockets

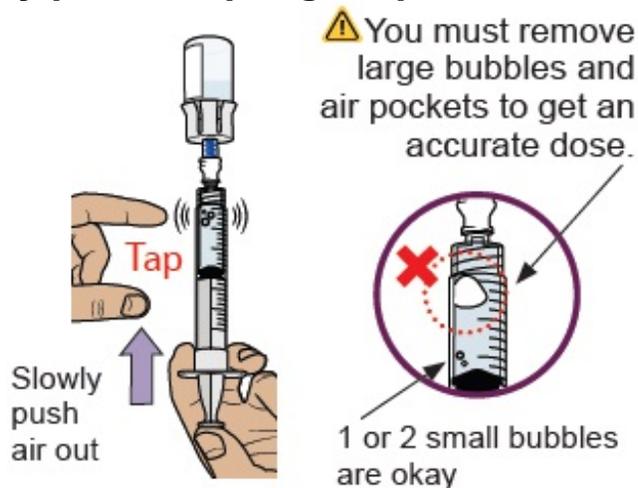
Check to see if there are large air bubbles or an air pocket in the syringe. You will remove extra air in the next steps.



26 Remove air bubbles and air pockets from syringe

If you see large air bubbles or an air pocket, tap the side of the dosing syringe to move the air to the top.

Slowly push the plunger up to remove extra air.



27 Compare amount to prescribed dose

After removing all extra air, **compare the amount in your syringe to your prescribed dose.**

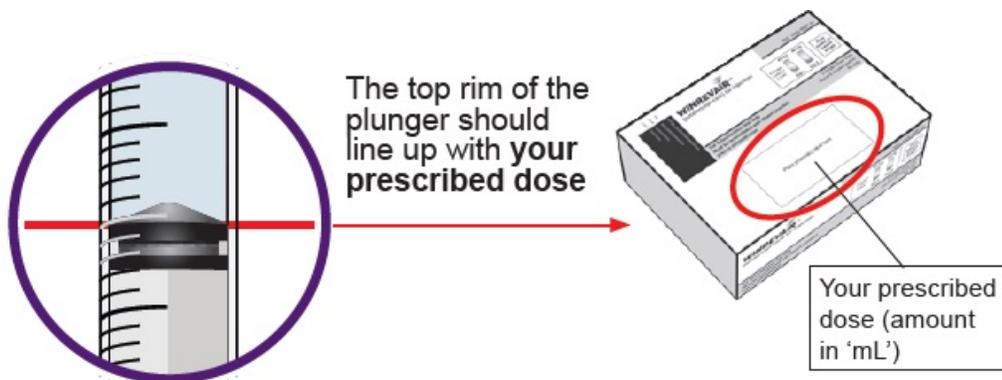
If you do not have your prescribed dose in your syringe, slowly pull the plunger back again to withdraw more medicine from the vial.

Repeat Steps 24 to 26 until you get **your prescribed dose** and no large bubbles or air pockets are visible.



28 Confirm your prescribed dose

Before you continue, check to make sure you have the prescribed dose in the dosing syringe.



⚠ **If the amount does not match your prescribed dose,**

repeat Steps 24 to 27.

29 Remove the dosing syringe from the vial and set dosing syringe aside



Hold the plunger in place



Throw away used vial

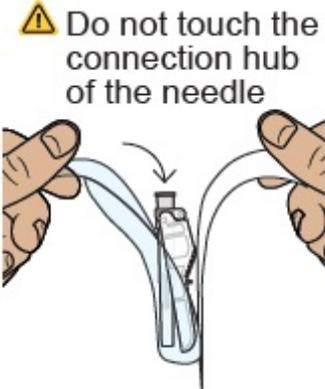
Hold the plunger in place with one hand. With the other hand, hold the vial adapter and unscrew the filled dosing syringe from the vial. Throw away the used vial into the sharps container.



Place the filled dosing syringe on a clean, flat surface.

 Do not touch the dosing syringe tip or let it to touch any surfaces.

30 Attach injection needle to dosing syringe



 Do not touch the connection hub of the needle

Find the needle in the bottom tray and open its package.



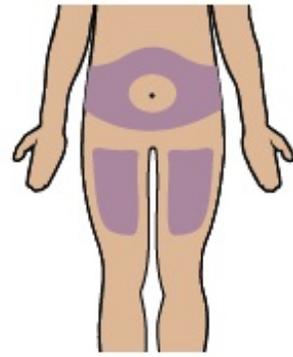
With the needle still in the package, **grip the base of the needle** and **twist on the dosing syringe** until it stops. Remove the needle package.



Move the safety shield away from the needle and toward the syringe. Place the dosing syringe on a clean, flat surface.

 Do not uncap the needle.

31 Choose and clean your injection site



Select an injection site on your stomach (abdomen) or your upper thigh. If injecting on your stomach, avoid a 2-inch area around your belly button. Choose a different site every time you inject.

 Do not inject into skin that is damaged, sore, tender, bruised, or has red patches.

 Do not inject through clothes.



Clean the injection site with a new alcohol wipe.

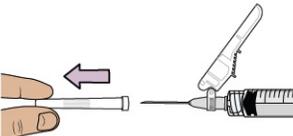
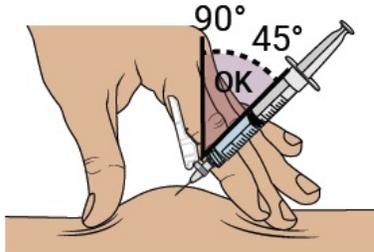
 Do not touch the cleaned injection site

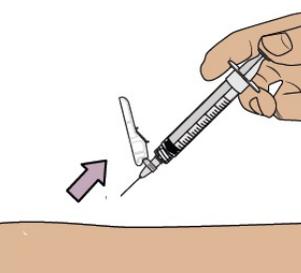
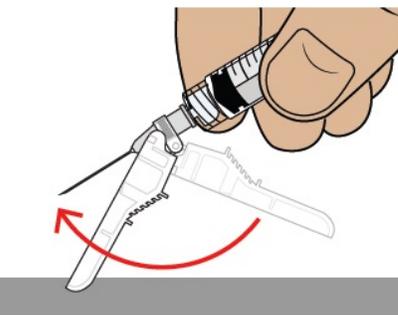
again.

Now, you are ready to inject the medicine.

Inject your medicine

32 Inject your medicine

 <p>Pull the cap straight off the needle. Throw away the cap.</p> <p>⚠ Do not touch the plunger until ready to inject so you do not lose any medicine.</p>	 <p>Gently pinch and hold a fold of skin where you will inject. Insert the needle with a dartlike motion at a 45° to 90° angle.</p>	 <p>Push the plunger with slow, steady pressure all the way down until the dosing syringe is empty. Confirm all the medicine has been injected. You can let go of the skin fold now.</p>
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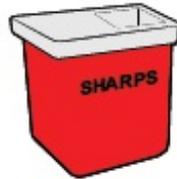
<p>⚠ Keep your fingers away from the needle at all times.</p>		
 <p>While keeping the plunger pushed in, remove the needle from your skin at the same angle you inserted it.</p>	 <p>To reapply the safety shield, push the shield against a flat surface until you hear a “click” and see that the needle is covered.</p>	 <p>Throw away the dosing syringe and used items in a sharps container.</p> <p>⚠ Do not remove the needle from the dosing syringe.</p>

How to throw away WINREVAIR

- Throw away any used vials (including any

remaining WINREVAIR liquid), needle, vial and needle caps, and used syringes in an FDA-cleared sharps disposal container.

- Do not throw away the WINREVAIR vials, syringes, or needle in your household trash.
- **Do not reuse any of the supplies.** WINREVAIR is for one time use only.
- **Important:** Always keep the sharps disposal container out of reach of children and pets.



If you do not have an FDA-cleared sharps disposal container, you may use a household container that is:

- made of a heavy-duty plastic,
- can be closed with a tight-fitting, puncture-resistant lid, without sharp objects being able to come out,
- upright and stable during use,
- leak resistant, and
- properly labeled to warn of hazardous waste inside the container.

When your sharps disposal container is almost full, you will need to follow your community guidelines for the right way to throw away your sharps container. There may be state or local laws about how you should throw away used needles, dosing syringes, and prefilled syringes.

For more information about safe sharps disposal, and for specific information about sharps disposal in the state that you live in, go to the FDA's website at:

<http://www.fda.gov/safesharpsdisposal>.

 Do not recycle your used sharps container.

Frequently asked questions

What should I do if I am bleeding at the injection site?

Place a cotton ball or bandage on your skin right away and apply a small amount of pressure. If the bleeding does not stop, call your healthcare provider right away.

Where can I find my prescribed dose?

Your prescribed dose in 'mL' is on your prescription label on your kit. Contact your healthcare provider if you cannot find your prescribed dose.

What should I do if I accidentally get some medicine on my skin or my work surface?

Wash the area thoroughly with soap and water right away.

What should I do if I am not sure I administered my prescribed dose correctly?

Call your pharmacy or healthcare provider.

What should I do if the plunger of my dosing syringe moves automatically when I try to withdraw medicine from the vial?

Do not worry if your plunger moves slightly on its own when you are filling your dosing syringe with medicine.

With one hand, **hold the plunger in place to stop the plunger from moving.** With the other hand, unscrew the vial from the dosing syringe.

After unscrewed, it is safe to let go of the plunger.

You can avoid this automatic plunger movement by pushing air into the vial before filling your dosing syringe with medicine. Refer to Steps 17 to 28 for detailed instructions.

What should I do if my kit parts are damaged or are discolored, cloudy, or have particles?

Do not use your kit. Call your pharmacy to get a new kit.

What should I do if my medicine does not turn clear after mixing and swirling?

Do not use the medicine if you have swirled the medicine vial for about 2 minutes and let it stand for another 3 minutes but your medicine vial remains cloudy or has clumps, powder, or foreign particles. Call your pharmacy to get a new kit.

What should I do if the sterile water will not come out of the prefilled syringe?

Check that the vial adapter is attached to the vial securely. If not, hold the vial and press the vial adapter down firmly to make sure the vial adapter punctures the vial rubber stopper.

What should I do if I dropped my kit components?

Do not use if any items are damaged. If you are unsure, call your pharmacy to get a new kit.

Can I use my kit that has been left out of the refrigerator?

Store the injection kit in the refrigerator until ready to use it. If the unused injection kit has been out of the refrigerator up to 77°F (25°C) for more than 24 hours, please contact your pharmacy or healthcare provider.

Do I need to use mixed medicine right away?

We recommend you inject the medicine right away after mixing but no later than 4 hours after mixing. If it has been more than 4 hours, throw away unused mixed medicine. If you have questions, please contact your pharmacy or healthcare provider.

How can I get help preparing and giving my injection?

If you have questions about how to give WINREVAIR the correct way or need more information, you can call your pharmacy or healthcare provider.

Call your doctor for advice about side effects.

To report side effects or product quality complaints, please call Merck Sharp & Dohme LLC at 1-800-672-6372. You can also report side effects to FDA at 1-800-FDA-1088.

Manufactured by:
Merck Sharp & Dohme LLC
Rahway, NJ 07065, USA
U.S. License Number 0002

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This Instructions for Use has been approved by the U.S. Food and Drug Administration
Approved: March 2024

PRINCIPAL DISPLAY PANEL - 45 mg Vial Kit Carton

NDC 0006-5090-01

WINREVAIR™
(sotatercept-csrk) for injection

45 mg/vial

Package
contains
1 Vial

**Dose
based on
patient
weight**

For Subcutaneous Use Only
Must be reconstituted with diluent provided
prior to administration

1 Single-Dose Vial
Discard unused portion

Rx only



Place pharmacy label here

Encoding area: Space reserved for 2D serialization Barcode, Serial Number, Expiry and Lot

For Subcutaneous Use Only
Must be reconstituted with diluent provided
1 Single-Dose Vial Discard unused portion
Rx only

Dose based on patient weight
45mg
Vial
Package contains 1 Vial

45 mg/vial
WINREVAIR™
(sotatercept-csrk) for injection

STN 00300065090016
SN
EXP
LOT

NDC 0006-5090-01

Storage: Store refrigerated at 2°C to 8°C (36°F to 46°F) in original carton to protect from light. **Do NOT freeze.**

Do not open the package or mix the medicine until you are ready to use it.

No Preservative.
Keep out of the reach of children and pets.

Dosage: See Prescribing Information. After reconstitution with 1 mL Sterile Water for Injection, the resulting concentration is 50 mg/mL of sotatercept-csrk and the nominal deliverable volume is 0.9 mL.

Each 45 mg single-dose vial contains: 45 mg of sotatercept-csrk, citric acid monohydrate (0.40 mg), polysorbate 80 (0.18 mg), sodium citrate (1.84 mg), and sucrose (72 mg) at pH 5.8.

Each carton contains:

- One WINREVAIR™ single-dose vial
 - One single-dose prefilled syringe of Diluent (Sterile Water for Injection), 1 mL
 - One 3-mL plastic syringe for injection
 - One 27-gauge needle for injection
 - One vial adapter
 - Four alcohol wipes
 - One Instructions for Use
 - One Package Insert for Full Prescribing Information
- Read accompanying directions carefully for the preparation of WINREVAIR™ for Injection.

WINREVAIR™
(sotatercept-csrk) for injection

45 mg/vial

Package contains 1 Vial

45 mg
Vial

Dose based on patient weight

For Subcutaneous Use Only
Must be reconstituted with diluent provided prior to administration

1 Single-Dose Vial
Discard unused portion
Rx only

NDC 0006-5090-01

Manufactured by: Merck Sharp & Dohme LLC
Rahway, NJ 07065, USA
U.S. License No. 0002

At: Patheon Italia, S.p.A.
20600 Monza (MB), Italy

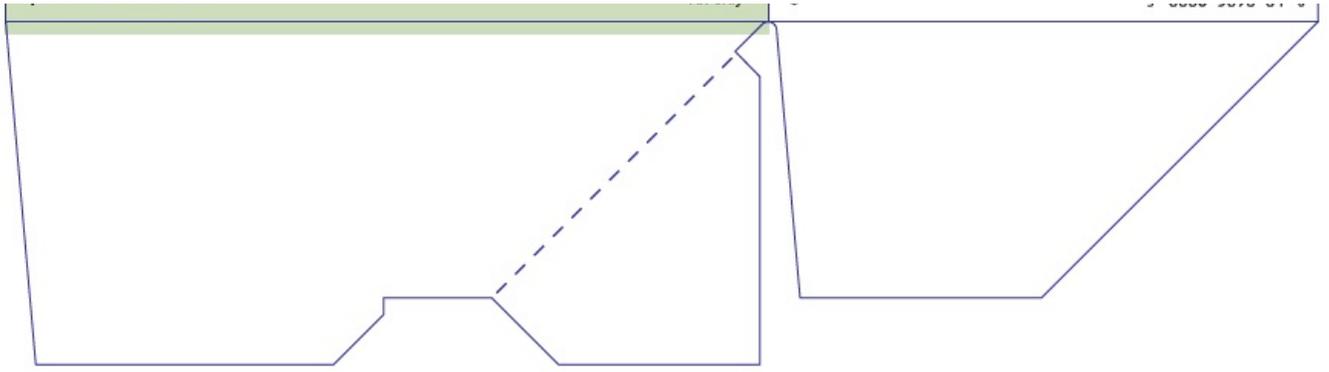
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N 0006-5090-01 4



PRINCIPAL DISPLAY PANEL - 60 mg Vial Kit Carton

NDC 0006-5091-01

WINREVAIR™
(sotatercept-csrk) for injection

60 mg/vial

Package
contains
1 Vial

**Dose
based on
patient
weight**

For Subcutaneous Use Only
**Must be reconstituted with diluent provided
prior to administration**

1 Single-Dose Vial
Discard unused portion

Rx only



Place pharmacy label here

Encoding area: Space reserved for 2D serialization Barcode, Serial Number, Expiry and Lot

For Subcutaneous Use Only
Must be reconstituted with diluent provided
prior to administration
Discard unused portion
Rx only

1 Single-Dose Vial

Dose based on patient weight

Vial

Package contains 1 Vial

60 mg

WINREVAIR™
(sotatercept-csrk) for injection

60 mg/vial

CTIN 00300065091013

SH
EXP
LOT

NDC 0006-5091-01

Storage: Store refrigerated at 2°C to 8°C (36°F to 46°F) in original carton to protect from light. **Do NOT freeze.**

Do not open the package or mix the medicine until you are ready to use it.

No Preservative.

Keep out of the reach of children and pets.

Dosage: See Prescribing Information. After reconstitution with 1.3 mL Sterile Water for Injection, the resulting concentration is 50 mg/mL of sotatercept-csrk and the nominal deliverable volume is 1.2 mL.

Each 60 mg single-dose vial contains: 60 mg of sotatercept-csrk, citric acid monohydrate (0.53 mg), polysorbate 80 (0.24 mg), sodium citrate (2.45 mg), and sucrose (96 mg) at pH 5.8.

Each carton contains:

- One WINREVAIR™ single-dose vial
 - One single-dose prefilled syringe of Diluent (Sterile Water for Injection), 1.3 mL
 - One 3-mL plastic syringe for injection
 - One 27-gauge needle for injection
 - One vial adapter
 - Four alcohol wipes
 - One Instructions for Use
 - One Package Insert for Full Prescribing Information
- Read accompanying directions carefully for the preparation of WINREVAIR™ for Injection.

WINREVAIR™
(sotatercept-csrk) for injection

60 mg/vial

Package contains 1 Vial

60 mg



Vial

Dose based on patient weight

1 Single-Dose Vial
Discard unused portion
Rx only

NDC 0006-5091-01

Manufactured by: Merck Sharp & Dohme LLC
Rahway, NJ 07065, USA
U.S. License No. 0002

At: Patheon Italia, S.p.A.
20800 Monza (MB), Italy

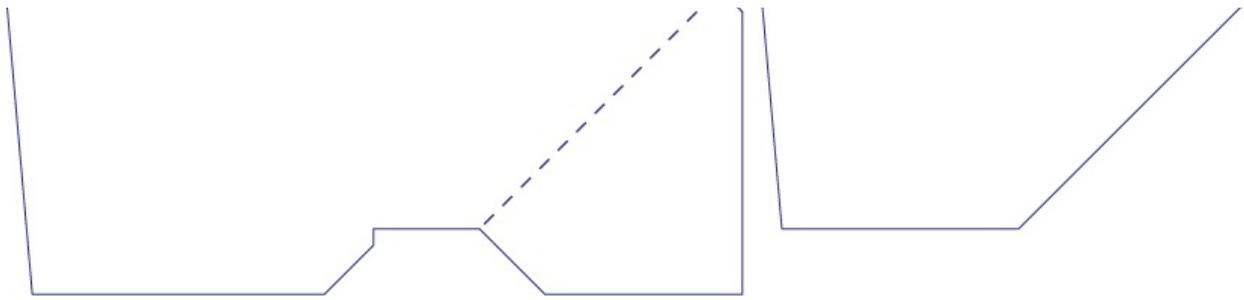
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7010272800



3 0006-5091-01 3



PRINCIPAL DISPLAY PANEL - 45 mg 2 Vials Kit Carton

NDC 0006-5087-01

**WINREVAIR™
(sotatercept-csrk) for injection**

45 mg/vial

Package
contains
2 Vials

**Dose
based on
patient
weight**

**For Subcutaneous Use Only
Must be reconstituted with diluent provided
prior to administration**

2 Single-Dose Vials
Discard unused portion

Rx only

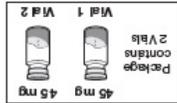


Place pharmacy label here

Encoding area: Space reserved for 2D Serialization Barcode, Serial Number, Expiry and Lot

For Subcutaneous Use Only
Must be reconstituted with diluent provided
prior to administration
2 Single-Dose Vials
Discard unused portion
Rx only

Dose based on patient weight



45 mg/vial

WINREVAIR™
(sotatercept-csrk) for injection

STIN 00300065087016
LOT
EXP
SN

NDC 0006-5087-01

Storage: Store refrigerated at 2°C to 8°C (36°F to 46°F) in original carton to protect from light. **Do NOT freeze.**

Do not open the package or mix the medicine until you are ready to use it.

No Preservative.

Keep out of the reach of children and pets.

Dosage: See Prescribing Information. After reconstitution with 1 mL Sterile Water for Injection, the resulting concentration is 50 mg/mL of sotatercept-csrk and the nominal deliverable volume is 0.9 mL.

Each 45 mg single-dose vial contains: 45 mg of sotatercept-csrk, citric acid monohydrate (0.40 mg), polysorbate 80 (0.18 mg), sodium citrate (1.84 mg), and sucrose (72 mg) at pH 5.8.

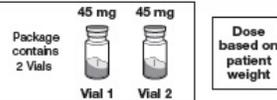
Each carton contains:

- Two WINREVAIR™ single-dose vials
 - Two single-dose pre-filled syringes of Diluent (Sterile Water for Injection), 1 mL
 - One 3-mL plastic syringe for injection
 - One 27-gauge needle for injection
 - Two vial adapters
 - Eight alcohol wipes
 - One Instructions for Use
 - One Package Insert for Full Prescribing Information
- Read accompanying directions carefully for the preparation of WINREVAIR™ for Injection.

WINREVAIR™
(sotatercept-csrk) for injection

45 mg/vial

NDC 0006-5087-01



Dose based on patient weight

For Subcutaneous Use Only
Must be reconstituted with diluent provided
prior to administration

2 Single-Dose Vials
Discard unused portion
Rx only

Manufactured by: Merck Sharp & Dohme LLC
Rahway, NJ 07065, USA
U.S. License No. 0002

At: Patheon Italia, S.p.A.
20900 Monza (MB), Italy

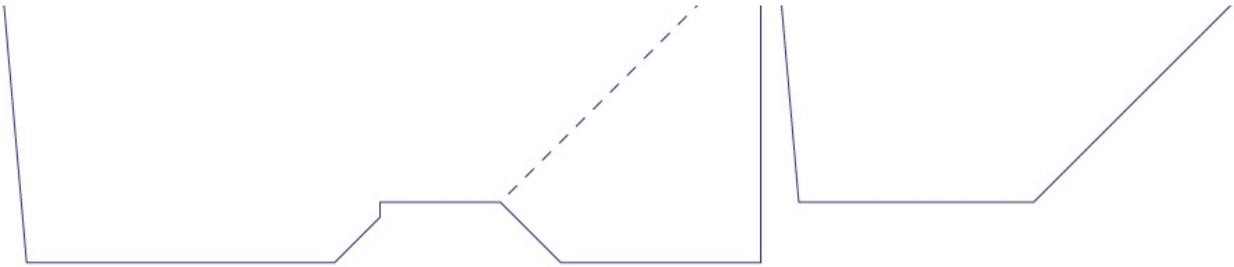
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7010272500



0006-5087-01 6



PRINCIPAL DISPLAY PANEL - 60 mg 2 Vials Kit Carton

NDC 0006-5088-01

WINREVAIR™
(sotatercept-csrk) for injection

60 mg/vial

Package
contains
2 Vials

**Dose
based on
patient
weight**

For Subcutaneous Use Only
Must be reconstituted with diluent provided
prior to administration

2 Single-Dose Vials
Discard unused portion

Rx only



Place pharmacy label here

Encoding area: Space reserved for 2D Serialization Barcode, Serial Number, Expiry and Lot

For Subcutaneous Use Only
Must be reconstituted with diluent provided
prior to administration
Discard unused portion
Rx only

60 mg/vial

Package contains 2 Vials
Vial 1 60 mg
Vial 2 60 mg

Dose based on patient weight

WINREVAIR™
(sotatercept-csrk) for injection

GTIN 00300065088013

SN
EXP
LOT

NDC 0006-5088-01

Storage: Store refrigerated at 2°C to 8°C (36°F to 46°F) in original carton to protect from light. **Do NOT freeze.**

Do not open the package or mix the medicine until you are ready to use it.

No Preservative.
Keep out of the reach of children and pets.

Dosage: See Prescribing Information. After reconstitution with 1.3 mL Sterile Water for Injection, the resulting concentration is 50 mg/mL of sotatercept-csrk and the nominal deliverable volume is 1.2 mL.

Each 60 mg single-dose vial contains: 60 mg of sotatercept-csrk, citric acid monohydrate (0.53 mg), polysorbate 80 (0.24 mg), sodium citrate (2.45 mg), and sucrose (96 mg) at pH 5.8.

Each carton contains:

- Two WINREVAIR™ single-dose vials
- Two single-dose prefilled syringes of Diluent (Sterile Water for Injection), 1.3 mL
- One 3-mL plastic syringe for injection
- One 27-gauge needle for injection
- Two vial adapters
- Eight alcohol wipes
- One Instructions for Use
- One Package Insert for Full Prescribing Information

Read accompanying directions carefully for the preparation of WINREVAIR™ for Injection.

WINREVAIR™
(sotatercept-csrk) for injection

60 mg/vial

Package contains 2 Vials
Vial 1 60 mg
Vial 2 60 mg

Dose based on patient weight

NDC 0006-5088-01

For Subcutaneous Use Only
Must be reconstituted with diluent provided
prior to administration

2 Single-Dose Vials
Discard unused portion
Rx only

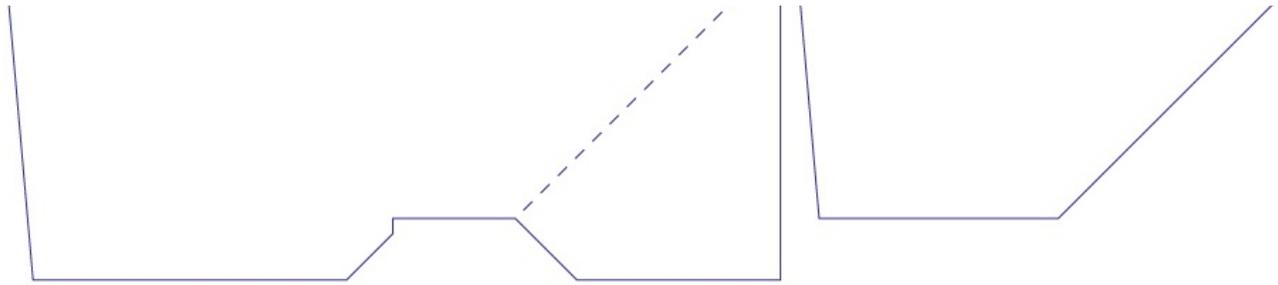
Manufactured by: Merck Sharp & Dohme LLC
Rahway, NJ 07065, USA
U.S. License No. 0002

At: Patheon Italia, S.p.A.
20900 Monza (MI), Italy

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MERCK 701027200

0006-5088-01 3



WINREVAIR

sotatercept-csrk kit

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:0006-5090
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0006-5090-01	1 in 1 CARTON	03/26/2024	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 VIAL, SINGLE-DOSE	1 mL
Part 2	1 SYRINGE	1 mL

Part 1 of 2

WINREVAIR

sotatercept-csrk injection, powder, lyophilized, for solution

Product Information

Item Code (Source)	NDC:0006-5068
Route of Administration	SUBCUTANEOUS

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SOTATERCEPT (UNII: 0QI90BTJ37) (SOTATERCEPT - UNII:0QI90BTJ37)	SOTATERCEPT	45 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
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CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	0.40 mg in 1 mL
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	0.18 mg in 1 mL
SUCROSE (UNII: C151H8M554)	72 mg in 1 mL
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	2.1 mg in 1 mL

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0006-5068-99	1 mL in 1 VIAL, SINGLE-DOSE; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA761363	03/26/2024	

Part 2 of 2

STERILE WATER

water injection, solution

Product Information

Item Code (Source)	NDC:0006-5079
Route of Administration	SUBCUTANEOUS

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)	WATER	1 mL in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0006-5079-89	1 mL in 1 SYRINGE; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA761363	03/26/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA761363	03/26/2024	

WINREVAIR

sotatercept-csrk kit

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:0006-5091
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0006-5091-01	1 in 1 CARTON	03/26/2024	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 VIAL, SINGLE-DOSE	1 mL
Part 2	1 SYRINGE	1.3 mL

Part 1 of 2

WINREVAIR

sotatercept-csrk injection, powder, lyophilized, for solution

Product Information

Item Code (Source)	NDC:0006-5069
Route of Administration	SUBCUTANEOUS

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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SOTATERCEPT (UNII: 0QI90BTJ37) (SOTATERCEPT - UNII:0QI90BTJ37) SOTATERCEPT 60 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	0.53 mg in 1 mL
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	0.24 mg in 1 mL
SUCROSE (UNII: C151H8M554)	96 mg in 1 mL
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	2.8 mg in 1 mL

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0006-5069-99	1 mL in 1 VIAL, SINGLE-DOSE; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA761363	03/26/2024	

Part 2 of 2

STERILE WATER

water injection, solution

Product Information

Item Code (Source)	NDC:0006-5080
Route of Administration	SUBCUTANEOUS

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)	WATER	1.3 mL in 1.3 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0006-5080-89	1.3 mL in 1 SYRINGE; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA761363	03/26/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA761363	03/26/2024	

WINREVAIR

sotatercept-csrk kit

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:0006-5087
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0006-5087-01	1 in 1 CARTON	03/26/2024	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	2 VIAL, SINGLE-DOSE	2 mL
Part 2	2 SYRINGE	2 mL

Part 1 of 2

WINREVAIR

sotatercept-csrk injection, powder, lyophilized, for solution

Product Information

Item Code (Source)	NDC:0006-5068
Route of Administration	SUBCUTANEOUS

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SOTATERCEPT (UNII: 0QI90BTJ37) (SOTATERCEPT - UNII:0QI90BTJ37)	SOTATERCEPT	45 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	0.40 mg in 1 mL
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	0.18 mg in 1 mL
SUCROSE (UNII: C151H8M554)	72 mg in 1 mL
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	2.1 mg in 1 mL

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0006-5068-99	1 mL in 1 VIAL, SINGLE-DOSE; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA761363	03/26/2024	

Part 2 of 2

STERILE WATER

water injection, solution

Product Information

Item Code (Source)	NDC:0006-5079
Route of Administration	SUBCUTANEOUS

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)	WATER	1 mL in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0006-5079-89	1 mL in 1 SYRINGE; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA761363	03/26/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA761363	03/26/2024	

WINREVAIR

sotatercept-csrk kit

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:0006-5088
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0006-5088-01	1 in 1 CARTON	03/26/2024	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	2 VIAL, SINGLE-DOSE	2 mL
Part 2	2 SYRINGE	2.6 mL

Part 1 of 2**WINREVAIR**

sotatercept-csrk injection, powder, lyophilized, for solution

Product Information

Item Code (Source)	NDC:0006-5069
Route of Administration	SUBCUTANEOUS

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SOTATERCEPT (UNII: 0QI90BTJ37) (SOTATERCEPT - UNII:0QI90BTJ37)	SOTATERCEPT	60 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	0.53 mg in 1 mL
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	0.24 mg in 1 mL
SUCROSE (UNII: C151H8M554)	96 mg in 1 mL
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	2.8 mg in 1 mL

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0006-5069-99	1 mL in 1 VIAL, SINGLE-DOSE; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA761363	03/26/2024	

Part 2 of 2

STERILE WATER

water injection, solution

Product Information

Item Code (Source)	NDC:0006-5080
Route of Administration	SUBCUTANEOUS

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)	WATER	1.3 mL in 1.3 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0006-5080-89	1.3 mL in 1 SYRINGE; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA761363	03/26/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
BLA	BLA761363	03/26/2024	

Labeler - Merck Sharp & Dohme LLC (118446553)