

Neos Products Savings Offer

All eligible patients with commercial (non-government) insurance

Cotempla XR-ODT[®]
Extended-Release Orally Disintegrating Tablets
(methylphenidate) 8.6 mg, 17.3 mg, 25.9 mg

Adzenys ER[™]
Extended-Release
(amphetamine) 1.25 mg/mL

Adzenys XR-ODT[®]
Extended-Release Orally Disintegrating Tablets
(amphetamine) 3.1 mg, 6.3 mg, 9.4 mg, 12.5 mg, 15.7 mg, 18.8 mg

Commercial (non-government) insured patients pay*

BIN# 600426
PCN# 54
GRP# ECNEOSRTL
ID# 09115497340

OR

- \$15 Per prescription for insurance "Covered"
- \$50 Per prescription for insurance "Not Covered" or high-deductible health plan

The ACE Customer Excellence Team

If you experience any pharmacy-related issues, call our dedicated staff at 1-877-675-6590.

Support available Monday through Friday, 8:30 am to 8:00 pm ET (except holidays).

*Eligible patients with commercial insurance valid only for 30 day supply. Terms and conditions may apply.

Please see Important Safety Information and the Medication Guide within the accompanying Full Prescribing Information for [Cotempla XR-ODT](#), [Adzenys ER](#) and [Adzenys XR-ODT](#).

Pharmacist Instructions for a Patient with an Eligible Third Party Payer: When you redeem this offer, you certify that you have not submitted and will not submit a claim for reimbursement under any federal, state, or other government programs for this prescription. Submit the claim to the primary Third Party Payer first, then submit the balance due to CHANGE HEALTHCARE, as a Secondary Payer, as a copay only billing using a valid Other Coverage Code (eg, 8). For prescription fill(s) of up to a 30 day supply, eligible commercially insured patients who are "covered" will incur an out-of-pocket expense of \$15. If coverage is rejected due to Prior Authorization, step-edit or NDC block, patients are still considered eligible, "not covered", and pharmacists can submit Other Coverage Code of 03 (secondary claim). The out-of-pocket expense for "not covered" patients will be \$50 per prescription fill for up to a 30 day supply. For commercially insured patients with a deductible, please follow instructions for covered patients using a valid Other Coverage Code (eg, 8). The out-of-pocket expense for deductible patients will be \$50 per prescription fill for up to a 30 day supply.

Pharmacist Instructions for a Cash-Paying Patient: Cash-paying patients may receive up to \$100 off each prescription fill for 30 day supply. Submit this claim to CHANGE HEALTHCARE using a valid Other Coverage Code (eg, 1). After receiving a savings of \$100, patients pay remaining balance; out-of-pocket costs may vary. Reimbursement will be received from CHANGE HEALTHCARE.

For any questions regarding CHANGE HEALTHCARE online processing, please call the Help Desk at 1-800-433-4893.

Restrictions: This offer is valid only in the United States and may not be available in all states. Program is applicable only for commercially insured patients. Cash discount cards are not commercial payers and are not eligible to be used for this program. Offer not valid for prescriptions reimbursed under Medicaid, a Medicare drug benefit plan (including Medicare Advantage and Parts A, B and D plans), Medigap, VA, DOD, CHAMPUS, or TRICARE or other federal or state health programs (such as medical assistance programs). If the patient is eligible for benefits under any such program, the patient cannot use this offer. By using this offer, the patient certifies that he or she will comply with any terms of his or her health insurance contract requiring notification to his or her payer of the existence and/or value of this offer. It is illegal to (or offer to) sell, purchase, or trade this offer. This offer is not transferable and has no cash value. Cannot be combined with any other rebate/coupon, free trial, or similar offer for the specified prescription(s). Not valid if reproduced. Void where prohibited by law. Program managed by ConnectiveRx, on behalf of Neos Therapeutics. The parties reserve the right to rescind, revoke, amend or terminate this offer without written notice at any time. Patient age or insurance restrictions may apply.

Please see Important Safety Information on next page and Medication Guide within the Full Prescribing Information for [Cotempla XR-ODT](#), [Adzenys ER](#) and [Adzenys XR-ODT](#).

IMPORTANT SAFETY INFORMATION FOR COTEMPLA XR-ODT

Cotempla XR-ODT is a federally controlled substance (CII) because it can be abused or lead to dependence. Keep Cotempla XR-ODT in a safe place to protect it from theft. Selling or giving away your Cotempla XR-ODT may cause death or harm to others and is against the law.

Who should not take Cotempla XR-ODT?

Do not give Cotempla XR-ODT to your child if they are:

- allergic to methylphenidate or any ingredients in Cotempla XR-ODT.
- taking or has taken an anti-depression medicine called monoamine oxidase inhibitor (MAOI) within the past 14 days.

What is the most important information I should know about Cotempla XR-ODT?

Cotempla XR-ODT can cause serious side effects. Tell your healthcare provider about health conditions, including if your child:

- has ever abused or been dependent on alcohol, prescription medicines, or street drugs. Cotempla XR-ODT has a high chance for abuse and can cause physical and psychological dependence.
- has any heart problems, heart defects, high blood pressure, or a family history of these problems. This is important because sudden death has occurred in people with heart problems or defects. Increased blood pressure and heart rate have been reported. Your healthcare provider should check for heart problems prior to prescribing Cotempla XR-ODT and will check your child's blood pressure and heart rate regularly during treatment. **Call the healthcare provider or go to the nearest hospital emergency room right away if your child has any signs of heart problems such as chest pain, shortness of breath, or fainting while during treatment.**
- has mental problems, or a family history of suicide, bipolar illness, or depression. This is important because the following could occur: new or worse behavior and thought problems, new or worse bipolar illness, new psychotic symptoms (hearing voices, or seeing or believing things that are not real) or new manic symptoms. **Call your healthcare provider right away if there are any new or worsening mental symptoms or problems during treatment.**
- develops painful and prolonged erections (priapism). Priapism has happened in males who take products that contain methylphenidate. Get medical help right away if your child develops priapism.
- has circulation problems in fingers and toes (peripheral vasculopathy, including Raynaud's phenomenon). Fingers or toes may feel numb, cool, painful, and/or change color from pale, to blue, to red. Tell your healthcare provider if your child has numbness, pain, skin color change, or sensitivity to temperature in their fingers or toes. **Call the healthcare provider right away if any signs of unexplained wounds appear on fingers or toes while taking Cotempla XR-ODT.**
- is having slowing of growth (height and weight). Your child should have his or her height and weight checked often while taking Cotempla XR-ODT. Treatment may be stopped if your child is not gaining weight or height.
- is pregnant or plans to become pregnant. It is not known if Cotempla XR-ODT will harm the unborn baby. If your child becomes pregnant during treatment with Cotempla XR-ODT, talk to your healthcare provider about registering with the National Pregnancy Registry for Psychostimulants.
- is breastfeeding, or plans to breastfeed. You and your healthcare provider should decide if your child will take Cotempla XR-ODT or breastfeed.
- takes any medicines, including prescription and over-the-counter medicines (especially for depression, including MAOIs), vitamins, and herbal supplements. Cotempla XR-ODT and some medicines may interact with each other and cause serious side effects, or sometimes the dose of the other medicine will need to be adjusted.

Do not start any new medicine while taking Cotempla XR-ODT without talking to your healthcare provider first.

What should I avoid during treatment with Cotempla XR-ODT?

- You should avoid drinking alcohol during treatment with Cotempla XR-ODT.

Common side effects of Cotempla XR-ODT include:

Decreased appetite, trouble sleeping, nausea, vomiting, indigestion, stomach pain, weight loss, anxiety, dizziness, irritability, mood swings, increased heart rate, and increased blood pressure.

These are not all the possible side effects of Cotempla XR-ODT. Call your healthcare provider for medical advice about side effects.

What is Cotempla XR-ODT?

Cotempla XR-ODT is a central nervous system (CNS) stimulant prescription medicine used for the treatment of Attention Deficit Hyperactivity Disorder (ADHD) in children 6 to 17 years of age. **Cotempla XR-ODT is a federally controlled substance (CII) because it contains methylphenidate that can be a target for people who abuse prescription medicines or street drugs.** Keep Cotempla XR-ODT in a safe place to protect it from theft. Selling or giving away your Cotempla XR-ODT may cause death or harm to others and is against the law.

For additional safety information, click here for [Medication Guide](#) and [Full Prescribing Information](#) and discuss with your healthcare provider.

You are encouraged to report negative side effects of prescription drugs to the FDA. Visit www.fda.gov/medwatch, or call 1-800-FDA-1088.

IMPORTANT SAFETY INFORMATION FOR ADZENYS XR-ODT AND ADZENYS ER

Adzenys XR-ODT and Adzenys ER are federally controlled substances (CII) because they can be abused or lead to dependence. Keep both Adzenys XR-ODT and Adzenys ER in a safe place to prevent misuse and abuse. Selling or giving away Adzenys XR-ODT or Adzenys ER may harm others and is against the law.

Tell your doctor if you or your child has ever abused or been dependent on alcohol, prescription medicines, or street drugs.

Who should not take Adzenys XR-ODT or Adzenys ER?

Do not take Adzenys XR-ODT or Adzenys ER if you or your child is:

- allergic to amphetamine or any ingredients in Adzenys XR-ODT or Adzenys ER.
- taking or has taken an anti-depression medicine called monoamine oxidase inhibitor (MAOI) within the past 14 days

Adzenys XR-ODT and Adzenys ER are stimulant medicines. Tell your doctor about health conditions, including if:

- you or your child has any heart problems, heart defects, high blood pressure, or a family history of these problems. This is important because sudden death has occurred in people with heart problems or defects, and sudden death, stroke and heart attack have happened in adults. Your doctor should check for heart problems prior to prescribing either Adzenys XR-ODT or Adzenys ER and will check you or your child's blood pressure and heart rate during treatment. **Call the doctor right away if you or your child has any signs of heart problems such as chest pain, shortness of breath, or fainting while taking either Adzenys XR-ODT or Adzenys ER.**
- you or your child has mental problems, or a family history of suicide, bipolar illness, or depression. This is important because the following could occur: new or worse behavior and thought problems, new or worse bipolar illness, new psychotic symptoms (hearing voices, believing things that are not true, are suspicious) or new manic symptoms. **Call the doctor right away if there are any new or worsening mental symptoms during treatment.**
- you or your child has circulation problems in fingers and toes (peripheral vasculopathy, including Raynaud's phenomenon). Fingers or toes may feel numb, cool, painful, sensitive to temperature and/or change color from pale, to blue, to red. **Call the doctor right away if any signs of unexplained wounds appear on fingers or toes while taking either Adzenys XR-ODT or Adzenys ER.**
- you or your child is having symptoms of serotonin syndrome which may happen when Adzenys XR-ODT or Adzenys ER is taken with certain other medicines and may be life threatening. **Call your doctor or go to the nearest hospital emergency room if you have any of the following symptoms:** agitation, hallucinations, coma or other changes in mental status, problems controlling your movements or muscle twitching, fast heartbeat, high or low blood pressure, sweating or fever, muscle stiffness or tightness, nausea, vomiting or diarrhea.
- your child is having slowing of growth (height and weight). Your child should have his or her height and weight checked often while taking either Adzenys XR-ODT or Adzenys ER. The doctor may stop treatment if a problem is found during these check-ups.
- you or your child has kidney problems. Your doctor may lower the dose.
- you or your child is, or plans to become pregnant. If you or your child becomes pregnant during treatment with either Adzenys XR-ODT or Adzenys ER, talk to your doctor about registering with the National Pregnancy Registry for Psychostimulants.
- you or your child is breastfeeding, or plans to breastfeed. You or your child should not breastfeed while taking either Adzenys XR-ODT or Adzenys ER.
- you or your child takes any medicines, including prescription and over-the-counter medicines, vitamins, and herbal supplements. Adzenys XR-ODT and Adzenys ER and some medicines may interact with each other and cause serious side effects.

Do not start any new medicine while taking either Adzenys XR-ODT or Adzenys ER without talking to your doctor first.

What should I avoid while taking Adzenys XR-ODT or Adzenys ER?

- drinking alcohol.

Common side effects for Adzenys XR-ODT and Adzenys ER include:

- **Children 6–12 Years:** Decreased appetite, problems sleeping, stomach pain, extreme mood change, vomiting, nervousness, nausea, and fever.
- **Children 13–17 Years:** Decreased appetite, problems sleeping, stomach pain and weight loss, and nervousness.
- **Adults:** Dry mouth, decreased appetite, problems sleeping, headache, weight loss, nausea, anxiety, restlessness, dizziness, fast heart beat, diarrhea, weakness, and urinary tract infections.

These are not all the possible side effects of Adzenys XR-ODT and Adzenys ER. Call your doctor for medical advice about side effects.

What are Adzenys XR-ODT and Adzenys ER?

Adzenys XR-ODT and Adzenys ER are central nervous system (CNS) stimulant prescription medicines used for the treatment of Attention Deficit Hyperactivity Disorder (ADHD) in people 6 years of age and older. **Adzenys XR-ODT and Adzenys ER are federally controlled substances (CII) because they contain amphetamine that can be a target for people who abuse prescription medicines or street drugs.** Keep both Adzenys XR-ODT or Adzenys ER in a safe place to protect it from theft. Selling or giving away your Adzenys XR-ODT or Adzenys ER may cause death or harm to others and is against the law.

For additional safety information for Adzenys XR-ODT, click here for the [Medication Guide](#) and [Full Prescribing Information](#) and discuss with your healthcare provider.

For additional safety information for Adzenys ER, click here for the [Medication Guide](#) and [Full Prescribing Information](#) and discuss with your healthcare provider.

You are encouraged to report negative side effects of prescription drugs to the FDA. Visit www.fda.gov/medwatch, or call 1-800-FDA-1088.



Adzenys ER™ is a trademark of Neos Therapeutics.

Adzenys XR-ODT® and Cotempla XR-ODT® are registered trademarks of Neos Therapeutics.

©2019 Neos Therapeutics Inc. All rights reserved. Printed in USA

N00497 01/19