Aytu BIOPHARMA | Aytu Product Savings Offer

Present this to your pharmacist

100% of eligible patients with commercial non-governmental insurance may pay as low as*

$15 Per prescription if **"covered"** on insurance plan and where deductible has been met

OR

$50 Per prescription if **"not covered"** on insurance plan or if the deductible has not been met. No prior authorization required

*Terms and conditions apply.

Pharmacist Instructions

1. Visit www.aytucoupon.com
2. Click "I'm a pharmacy"
3. Input necessary information and press submit
4. Utilize the offer to assist patient

ACE Customer Excellence Team

If you experience any pharmacy-related issues, call our dedicated staff at **1-888-AYTURxC (298-8792)**. Support available Monday through Friday, 9:00 am to 7:30 pm ET (except holidays).

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

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

Karbinal® ER (carbinoxamine maleate) extended-release oral suspension / 4mg/5mL

ZolpiMIST® (zolpidem tartrate) ORAL SPRAY C

Retail Copay Offer Terms and Conditions: This program provides non-government copay assistance with out-of-pocket costs for eligible patients. By using this offer, patient and pharmacist understand and agree to comply with these terms and conditions. Offer may only be used by eligible residents of the U.S. at participating pharmacies and may not be redeemed at government-subsidized clinics. Offer user must be at least 18 years of age to use for themselves or on behalf of a minor. Patient age or insurance restrictions may apply.

Offer limited to one per person and is not transferrable. No substitutions are permitted. Offer eligible only with valid prescription, has no cash value, and cannot be combined with any free trial, discount, prescription savings card or other offer. This offer is not insurance. Valid only for patients with commercial insurance and NOT valid for prescriptions eligible to be reimbursed in whole or in part by Medicaid, Medicare (including Medicare Advantage and Part D plans), or any other federally or state funded healthcare benefit program, or by commercial plans or other health or pharmacy benefit programs that reimburse for entire cost of the prescription drug or prohibit offer's use. Medicare Part D enrollees who are in the prescription drug coverage gap are not eligible for offer. Void where prohibited by law, taxed or restricted. It is illegal to sell, purchase, trade, or counterfeit, or offer to sell, purchase, trade or counterfeit the offer.

Patient, pharmacist and prescriber agree not to seek reimbursement for all or any part of the benefit received by patient through the offer. Certain information pertaining to use of the offer will be shared with Aytu BioPharma, the sponsor of the offer, and its affiliates. The information disclosed will include the patient copay ID, pharmacy demographics, prescriber information, and details relating to the claim, such as copay amount, insurance details, and therapy received. For more information, please see the Aytu BioPharma privacy policy.

Aytu BioPharma reserves the right to rescind, revoke or amend the offer at any time without notice.

Please see Important Safety Information on next page and Medication Guide within the Full Prescribing Information for Adzenys XR-ODT, Cotempla XR-ODT, Karbinal ER, and ZolpiMist.

IMPORTANT SAFETY INFORMATION FOR ADZENYS XR-ODT

Adzenys XR-ODT is a federally controlled substance (CII) because it can be abused or lead to dependence. Keep Adzenys XR-ODT in a safe place to prevent misuse and abuse. Selling or giving away Adzenys XR-ODT 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?

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

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

Adzenys XR-ODT is a stimulant medicine. 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 Adzenys XR-ODT 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 Adzenys XR-ODT.**
- 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 Adzenys XR-ODT.**
- your child is having slowing of growth (height and weight). Your child should have his or her height and weight checked often while taking Adzenys XR-ODT. 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.
- you or your child is breastfeeding, or plans to breastfeed. You should not breastfeed while taking Adzenys XR-ODT.
- you or your child takes any medicines, including prescription and over-the-counter medicines, vitamins, and herbal supplements. Adzenys XR-ODT and some medicines may interact with each other and cause serious side effects.

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

What should I avoid while taking Adzenys XR-ODT?

- drinking alcohol.

Common side effects of Adzenys XR-ODT include:

- Decreased appetite and problems sleeping.
- **Children 6 — 12 Years also include:** Stomach pain, extreme mood change, vomiting, nervousness, nausea, and fever.
- **Children 13 — 17 Years also include:** Stomach pain and weight loss.
- **Adults also include:** Dry mouth, 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. Call your doctor for medical advice about side effects.

What is Adzenys XR-ODT?

Adzenys XR-ODT is a prescription medicine used for the treatment of Attention Deficit Hyperactivity Disorder (ADHD) in patients 6 years and above.

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 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 KARBINAL ER

What is the most important information I should know about Karbinal ER?

- Karbinal ER is contraindicated in children younger than 2 years of age, and nursing mothers.
- Karbinal ER may produce marked drowsiness and impair the mental or physical abilities required for the performance of potentially hazardous tasks such as driving a car or operating machinery. Advise patients to avoid engaging in hazardous tasks requiring mental alertness and motor coordination after ingestion of Karbinal ER. Avoid concurrent use of Karbinal ER with alcohol or other central nervous system depressants because additional impairment of central nervous system performance may occur.
- Avoid use of Karbinal ER with alcohol and other CNS depressants (hypnotics sedatives, tranquilizers, etc.) due to additive effects.
- Karbinal ER is contraindicated in patients who are taking monoamine oxidase inhibitors (MAOI).

What are the possible side effects of Karbinal ER?

Use of Karbinal ER may result in decreased mental alertness with impaired mental or physical abilities.

The most frequent side effects include:

- Sedation
- Sleepiness
- Dizziness
- Disturbed coordination
- Epigastric distress
- Thickening of bronchial secretions

Ask your doctor or pharmacist for more information.

Call your doctor for medical advice about side effects.

You may report side effects to FDA at 1-800-FDA-1088.

For additional safety information, click here for Full Prescribing Information and discuss with your healthcare provider.

Keep Karbinal ER and all medicines out of the reach of children.

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 ZOLPIMIST

Zolpimist (zolpidem tartrate) Oral Spray is indicated for the short-term treatment of insomnia characterized by difficulties with sleep initiation. Zolpidem tartrate has been shown to decrease sleep latency for up to 35 days in controlled clinical studies. The clinical trials performed in support of efficacy were 4-5 weeks in duration with the final formal assessments of sleep latency performed at the end of treatment.

WARNING: COMPLEX SLEEP BEHAVIORS!

Complex sleep behaviors including sleep-walking, sleep-driving, and engaging in other activities while not fully awake may occur following use of ZOLPIMIST. Some of these events may result in serious injuries, including death. Discontinue ZOLPIMIST immediately if a patient experiences a complex sleep behavior.

IMPORTANT RISK INFORMATION

What is the most important information I should know about ZOLPIMIST?

Complex sleep behaviors that have caused serious injury and death.

After taking ZOLPIMIST, you may get up out of bed while not being fully awake and do an activity that you do not know you are doing (complex sleep behaviors). The next morning, you may not remember that you did anything during the night. These activities may happen whether or not you drink alcohol or take other medicines that make you sleepy with ZOLPIMIST. Reported activities include:

- driving a car ("sleep-driving")
- making and eating food
- talking on the phone
- having sex
- sleep-walking

Stop taking Zolpimist and call your doctor right away if you find out that you have done any of the above activities after taking ZOLPIMIST.

Important:

1. Take ZOLPIMIST exactly as prescribed

- Do not take more ZOLPIMIST than prescribed.
- Take ZOLPIMIST right before you get in bed, not sooner.

2. Do not take ZOLPIMIST if you:

- have ever experienced a complex sleep behavior (such as driving a car, making and eating food, talking on the phone or having sex while not fully awake) after taking ZOLPIMIST
- drink alcohol
- take other medicines that can make you sleepy. Talk to your doctor about all of your medicines. Your doctor will tell you if you can take ZOLPIMIST with your other medicines.
- cannot get a full night sleep

What is ZOLPIMIST?

ZOLPIMIST is a sedative-hypnotic (sleep) medicine. ZOLPIMIST is used in adults for the short-term treatment of a sleep problem called insomnia. Symptoms of insomnia include:

- trouble falling asleep

ZOLPIMIST is not for children.

ZOLPIMIST is a federally controlled substance (C-IV) because it can be abused and lead to dependence. Keep ZOLPIMIST in a safe place to prevent misuse and abuse. Selling or giving away ZOLPIMIST may harm others and is against the law. Tell your doctor if you have ever abused or have been dependent on alcohol, prescription medicines, or street drugs.

Who should not take ZOLPIMIST?

Do not take ZOLPIMIST if you have had an allergic reaction to zolpidem (Ambien, Ambien CR, ZOLPIMIST). Some signs of allergic reaction may be swelling of the face, a feeling of the throat closing, or difficulty breathing shortly after taking Zolpidem.

See the end of this Medication Guide for a complete list of ingredients in ZOLPIMIST.

ZOLPIMIST may not be right for you. Before starting ZOLPIMIST, tell your doctor about all of your health conditions, including if you:

- have a history of depression, mental illness, or suicidal thoughts
- have a history of drug or alcohol abuse or addiction
- have kidney or liver disease
- have a lung disease or breathing problems
- are pregnant, planning to become pregnant, or breastfeeding

Tell your doctor about all the medicines you take including prescription and nonprescription medicines, vitamins, and herbal supplements. Medicines can interact with each other, sometimes causing serious side effects.

Do not take ZOLPIMIST with other medicines that can make you sleepy.

Know the medicines you take. Keep a list of your medicines with you to show your doctor and pharmacist each time you get a new medicine.

How should I take ZOLPIMIST?

- Take ZOLPIMIST exactly as prescribed. Do not take more ZOLPIMIST than prescribed for you.
- **Take ZOLPIMIST right before you get into bed.**
- **Do not take ZOLPIMIST unless you are able to stay in bed a full night (7-8 hours) before you must be active again.**
- For faster sleep onset, ZOLPIMIST should NOT be taken with or immediately after a meal.
- Call your doctor if your insomnia worsens or is not better within 7 to 10 days. This may mean that there is another condition causing your sleep problem.
- If you take too much ZOLPIMIST or overdose, call your doctor or poison control center right away, or get emergency treatment.

What are the possible side effects of ZOLPIMIST?

Serious side effects of ZOLPIMIST include:

- **getting out of bed while not being fully awake and doing an activity that you do not know you are doing.** (See "What is the most important information I should know about ZOLPIMIST?")
- **abnormal thoughts and behavior.** Symptoms include more outgoing or aggressive behavior than normal, confusion, agitation, hallucinations, worsening of depression, and suicidal thoughts or actions.
- **memory loss**
- **anxiety**
- **severe allergic reactions.** Symptoms include swelling of the tongue or throat, trouble breathing, and nausea and vomiting. Get emergency medical help if you get these symptoms after taking ZOLPIMIST.

Call your doctor right away if you have any of the above side effects or any other side effects that worry you while using ZOLPIMIST.

The most common side effects of ZOLPIMIST are:

- drowsiness
- dizziness
- diarrhea
- "drugged feelings"
- You may still feel drowsy the next day after taking Zolpimist. **Do not drive or do other dangerous activities after taking ZOLPIMIST until you feel fully awake.**

After you stop taking a sleep medicine, you may have symptoms for 1 or 2 days such as: tiredness, trouble sleeping, nausea, flushing, lightheadedness, uncontrolled crying, vomiting, stomach cramps, panic attack, nervousness, and stomach area pain.

These are not all the side effects of ZOLPIMIST. Ask your doctor or pharmacist for more information.

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

ADVERSE EVENT REPORTING

To report suspected adverse events or product complaints, please contact **Aytu BioPharma at 855-AYTU-BIO; FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.**

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