



Illinois Naloxone Standardized Procedure

This updated Naloxone Standardized Procedure outlines how entities may become authorized to obtain, dispense, and administer naloxone hydrochloride for the purpose of reversing an opioid overdose. This Procedure also presents the educational requirements for obtaining the Illinois Naloxone Standing Order and the technique for administering naloxone.

Introduction

In September 2015, Illinois enacted Public Act 99-0480 (Act), expanding access to the opioid antagonist, naloxone. Naloxone may be used to reverse opioid overdoses, including those caused by heroin, fentanyl, and certain prescription pain medications. The law authorizes trained pharmacists and first responders to dispense naloxone as an opioid antagonist intervention.

Pursuant to the Act, the Illinois Department of Financial and Professional Regulation (IDFPR) – in consultation with the Illinois Department of Public Health (IDPH) and Illinois Department of Human Services (IDHS) – has issued a standardized procedure for appropriately trained professionals to obtain, dispense, or administer naloxone.

Naloxone Entity

Naloxone Entities may include pharmacies, pharmacists, or opioid overdose education and naloxone distribution (OEND) programs.

- Participating pharmacies and pharmacists must be licensed under the Illinois Pharmacy Practice Act (225 ILCS 85), complete training approved by IDHS pursuant to Public Act 99-0480, and have knowledge of this document, the Illinois Naloxone Standardized Procedure. Pharmacies/pharmacists should report naloxone dispensing to the Illinois Prescription Monitoring Program at <https://www.ilpmp.org/>.
- Any non-pharmacy OEND program must be registered as a Drug Overdose Prevention Program with the IDHS's Division of Alcoholism and Substance Abuse, at <http://www.dhs.state.il.us/>. This may include law enforcement agencies, drug treatment programs, local health departments, hospitals or urgent care facilities, or other for-profit or not-for-profit community-based organizations.

Educational Requirement

Under this standardized procedure, eligible entities must complete training in opioid overdose reversal, which includes the following:

- Opioid overdose recognition and prevention
- Naloxone administration techniques
- The importance of calling 911 for the care of the overdose victim after naloxone administration

Naloxone Hydrochloride

Naloxone is indicated for the reversal of opioid overdose, induced by natural or synthetic opioids, relative to respiratory depression or unresponsiveness. It should not be given to anyone known to be allergic to naloxone hydrochloride. It may be delivered subcutaneously or intramuscularly using an auto-injector, or needle and syringe, or intranasally.

Signs of Symptoms of Opioid Overdose

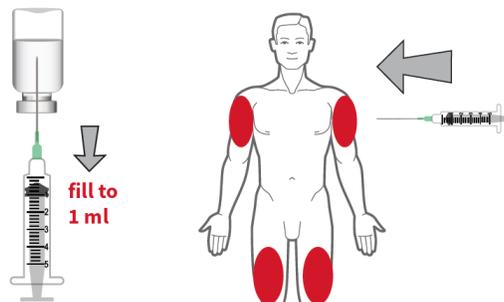
- Slowed, irregular, or no breathing
- Skin, nails turn blue
- Extreme sleepiness
- Unresponsive to sternal rub or when shaken
- Pinpoint pupils

Standardized Procedure for Naloxone Administration

1. Confirm signs and symptoms of potential opioid overdose
2. Call 9-1-1 and administer naloxone as follows (**select dispensed dosage form**):

Intramuscular Naloxone:

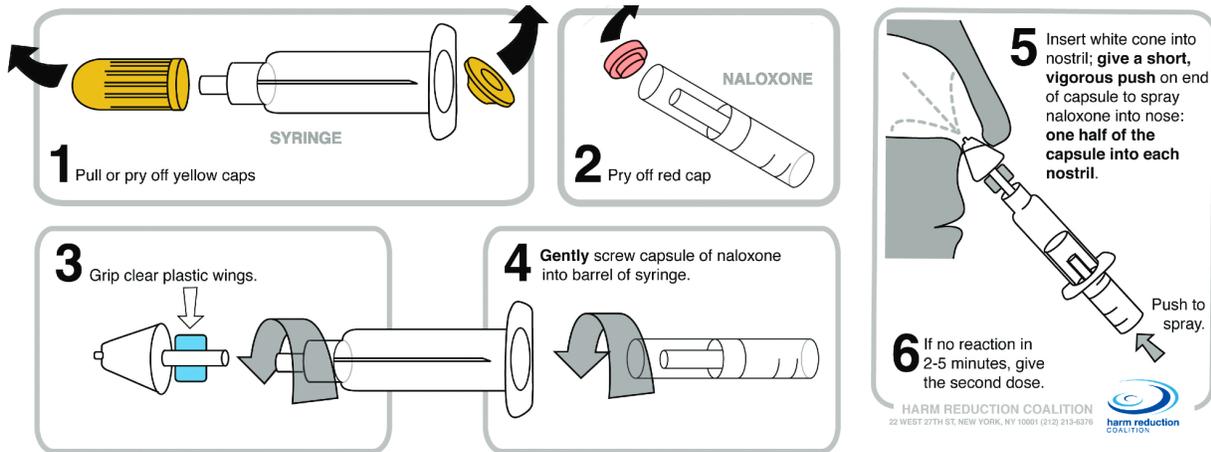
- Uncap the naloxone vial and uncap the muscle needle-syringe
- Insert the muscle needle through the rubber membrane on the naloxone vial, turn the vial upside down, draw up 1 ml of naloxone liquid, and withdraw the needle
- Insert the needle into the muscle of the upper arm or thigh of the victim, through clothing if needed, and push on the plunger to inject the naloxone
- Repeat the injection if there is no response after three minutes



Multi-Step Intranasal Naloxone:

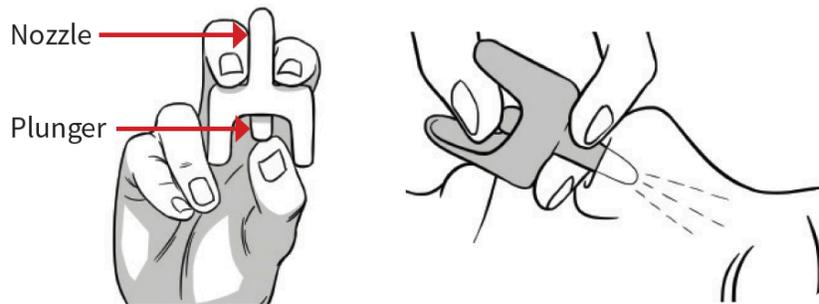
- Pop off two colored caps from the delivery syringe and one from the naloxone vial
- Screw the naloxone vial gently into the delivery syringe
- Screw the mucosal atomizer device onto the top of the syringe

- Spray half (1 mL) of naloxone in one nostril and the other half (1 mL) in the other
- Repeat if there is no response after three minutes



Single-Step Intranasal Naloxone:

- Peel back the package to remove the device
- Hold the device with your thumb on the bottom of the plunger and 2 fingers on the nozzle
- Place and hold the tip of the nozzle in either nostril until your fingers touch the bottom of the patient's nose
- Press the plunger firmly to release the dose into the patient's nose
- Repeat if there is no response after 3 minutes



Auto-injector Naloxone:

- Pull auto-injector from outer case and pull off red safety guard
- Place the black end of the auto-injector against the outer thigh, through clothing if needed, press firmly and hold in place for 5 seconds
- Repeat if there is no response after 3 minutes

3. Provide rescue breathing using a disposable rescue breathing device, chest compressions, or full cardiopulmonary resuscitation (CPR) based on the training and abilities of the responder¹ or follow the instructions of the 911 dispatcher.
4. If person becomes unresponsive again, administer another dose of naloxone. Stay with the person until emergency help arrives.

Contraindications

- Patient is known to be hypersensitive to naloxone hydrochloride

Precautions

- Pre-existing cardiac disease or seizure disorder
- Person is suspected to be physically dependent on opioids including newborns of mothers with opioid dependence (Reversal of opioid effect will precipitate acute abstinence syndrome)
- Use in Pregnancy:
 1. Teratogenic Effects: Pregnancy category C, no adequate or well-controlled studies in pregnant women
 2. Non-teratogenic Effects: Pregnant women known or suspected to have opioid dependence often have associated fetal dependence. Naloxone crosses the placenta and may precipitate fetal withdrawal symptoms
 3. Naloxone should only be used in pregnant women with opioid dependence in situations of life-threatening overdose
- Nursing Mothers: Caution should be exercised when administering to nursing women due to transmission in human milk
- Geriatric Use: Caution should be exercised for potential decreased hepatic, renal and cardiac function, as well as concomitant disease and other pharmacotherapies

Adverse Reactions

- Adverse reactions are related to precipitating opioid withdrawal. They include fever, hypertension, tachycardia, agitation, restlessness, diarrhea, nausea/vomiting, myalgias, diaphoresis, abdominal cramping, yawning, and sneezing.
- These symptoms may appear within minutes of naloxone administration and subside in approximately 2 hours.
- The severity and duration of the withdrawal syndrome is related to the dose of naloxone and the degree of opioid dependence.
- Adverse effects beyond opioid withdrawal are rare

¹As there is insufficient data to recommend one resuscitation method over another, naloxone entities will need to determine whether rescue breathing, chest compressions, both, or neither, is most appropriate for inclusion in their training curricula. (New York State Technical Working Group on Resuscitation Training in Naloxone Provision Programs. 2016)