

Healthcare Professional Specific Adverse Reaction Follow Up Form

This questionnaire is intended to follow-up on specific adverse reactions regarding opioid-related harms for your patient.

HEALTHCARE PROFESSIONAL SPECIFIC ADVERSE REACTION FOLLOW UP FORM	
MINT CONTACT INFORMATION: Telephone: +1 877-398-9696 Fax: +1 647-723-7696 Email: drugsafety@mintpharmaceuticals.com	FOR MINT USE ONLY: Reference case no: Mint Received Date: _____ (YYYY-MM-DD)
Reporter Information	
1. Reporter Name	
2. Reporter ☐ Qualification Physician ☐ Pharmacist ☐ Other health professional: _____	
3. Contact Information Email: _____ Phone: _____ Address: _____	
4. Type of Report ☐ Initial: _____ (YYYY-MM-DD) ☐ Follow-up: _____ (YYYY-MM-DD)	
5. Date of this Report _____ (YYYY-MM-DD)	
Patient Information	
1. Initials or Unique Identifier (if available)	
2. Age	
3. Sex ☐ Female ☐ Male	

4. Relevant Medical History/Risk Factors (e.g., pregnancy, smoking and alcohol use, hepatic/renal dysfunction, etc.)				
5. Relevant Tests/Laboratory data				
Lab Test:	Lab Test Date: (YYYY-MM-DD)	Results:	Normal Range:	Comments:
MINT-TRAMADOL/ACETAMINOPHEN Therapy Information				
1. Tramadol/Acetaminophen dosage				
2. Tramadol/Acetaminophen route of administration				
3. Indication				
4. Accidental Exposure ☐ Yes OR ☐ No				
5. Duration of Treatment				
Start date:		End date:		
Ongoing? ☐				
a) Long term exposure (greater than 12 weeks): ☐ Yes OR ☐ No				
6. Concomitant medications/treatments/supplements				
Product Name:	Dosage Regimen	Start Date: (YYYY-MM-DD)	Stop Date/Ongoing: (YYYY-MM-DD)	Indication of Use:

Adverse Reaction Information

1. Important identified risks (select all that apply):

☐ Abuse/ Misuse

Does the patient display any of the following signs or symptoms of **abuse/ misuse** (check all that apply):

- ☐ Pinpoint (very small) pupils
- ☐ Changes in appetite
- ☐ Nausea
- ☐ Vomiting
- ☐ Drowsiness
- ☐ Slurred speech
- ☐ Headaches
- ☐ Impaired coordination
- ☐ Death
- ☐ Other (please specify): _____

☐ Accidental Exposure

Does the patient display any of the following signs or symptoms of **accidental exposure** (check all that apply)

- ☐ Feelings of euphoria and relaxation
- ☐ False sense of well-being
- ☐ Confusion
- ☐ Sedation
- ☐ Drowsiness
- ☐ Dizziness/ lightheadedness
- ☐ Nausea and vomiting
- ☐ Constipation
- ☐ Respiratory depression or arrest
- ☐ Death
- ☐ Other (please specify): _____

☐ Death

☐ Interaction with alcohol, benzodiazepines, and other CNS depressants

Is the patient taking any of the following medications

- ☐ Alcohol
- ☐ Benzodiazepines
- ☐ Other central nervous system (CNS) depressants
- ☐ Other (please specify): _____

☐ Life-threatening respiratory depression

Has the patient experienced **life-threatening respiratory depression** (symptoms can include slowed breathing, long pauses between breaths, or shortness of breath)?

- ☐ Yes
- ☐ No

☐ Neonatal Opioid Withdrawal Syndrome (NOWS)

Has the infant experienced **Neonatal Opioid Withdrawal Syndrome** (NOWS)?

- ☐ Yes
- ☐ No
- ☐ Not applicable

If yes, what following signs or symptoms of **Neonatal Opioid Withdrawal Syndrome** did the patient display (check all that apply):

- ☐ Body shakes (tremors), seizures (convulsions), overactive reflexes (twitching) and tight muscle tone
- ☐ Fussiness, excessive crying or having a high-pitched cry
- ☐ Poor feeding or sucking or slow weight gain
- ☐ Breathing problems, including breathing really fast
- ☐ Fever, sweating or blotchy skin
- ☐ Trouble sleeping and lots of yawning
- ☐ Diarrhea or throwing up
- ☐ Stuffy nose or sneezing
- ☐ Other (please specify): _____

☐ Opioid Use Disorder (Addiction)

Has the patient shown any signs or symptoms of tramadol **addiction** (check all that apply):

- ☐ No signs of addiction
- ☐ Unable to Control Tramadol Use (The individual may attempt to reduce the dosage or is continue the tramadol, but cannot)
- ☐ Experiencing Substance Cravings
- ☐ Doctor Shopping (When the refills are denied, the individual may seek out other physicians to acquire new prescriptions for the tramadol)
- ☐ Experiencing Withdrawal Symptoms
- ☐ Other (please specify): _____

☐ Overdose

Has the patient shown any symptoms of **overdose** (check all that apply):

- ☐ Decreased size of the pupil (the black circle in the center of the eye)
- ☐ Difficulty breathing
- ☐ Slow or shallowing breathing
- ☐ Extreme drowsiness or sleepiness
- ☐ Unable to respond or wake up
- ☐ Slowed heartbeat
- ☐ Muscle weakness
- ☐ Cold, clammy skin
- ☐ Other (please specify): _____

☐ Physical and psychological dependence/tolerance (Drug withdrawal syndrome)/ Withdrawal

Has the patient shown signs or symptoms of **physical and psychological dependence/tolerance or withdrawal** (check all that apply)?

- ☐ Feeling agitated or anxious
- ☐ Panic attacks
- ☐ Feeling your heartbeat (palpitations)
- ☐ Difficulty sleeping
- ☐ Shaking
- ☐ Sweating
- ☐ Body aches
- ☐ Feeling restless
- ☐ Other (please specify): _____

Comments (if any):

2. Important potential risks (select all that apply): ☐ Medication error ☐ Off-label use (Ex. pediatric/adolescent populations, restless legs syndrome, antidepressant, premature ejaculation), please specify: _____ ☐ Diversion
3. Adverse Reaction Description:
4. Onset Date of Reaction _____ (YYYY-MM-DD)
5. Therapy Stop Date _____ (YYYY-MM-DD)
6. Outcome: ☐ Recovered ☐ Fatal ☐ Not Recovered ☐ Recovered with Sequelae ☐ Recovering ☐ OR Unknown
7. Was it treated?
8. Reporter Causality: ☐ Certain ☐ Probable ☐ Possible ☐ Unlikely ☐ Conditional OR ☐ Unassessable
9. Additional Information or Comments:

Reporter Signature:	Date (YYYY-MM-DD):
FOR MINT USE ONLY: Signature: Print Name:	Date (YYYY-MM-DD):