

Appendix 2

 <div style="display: inline-block; vertical-align: middle;"> <p style="font-size: small; margin: 0;">Winnipeg Regional Health Authority Office régional de la santé de Winnipeg</p> <p style="font-weight: bold; margin-top: 5px;">RIFAPENTINE/INH TREATMENT (Tx) FORM</p> </div>	<p style="font-size: large; color: #ccc; margin: 0;">ADDRESSOGRAPH</p>
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Rifapentine Tx Start Date (mmm/dd/yyyy): _____ Rifapentine Tx End Date (mmm/dd/yyyy): _____

Dose in mg: _____ Number of doses taken: _____

Completed Tx*: Yes No → Please fill in Reason Treatment Stopped/Reported Adverse Events below

* defined as at least 11 weekly doses administered within 16 weeks of Tx start; doses must be separated by > 72 hours to count.

If client did not complete Tx, please select the Reason Treatment Stopped (choose one):

- Client Stops Drug- Other
- Client Stops Drug- Side Effects
- Clinician Stops Drug- Other
- Clinician Stops Drug- Other Contraindication
- Clinician Stops Drug- Poor Compliance
- Clinician Stops Drug- Side Effects
- Lost to follow-up
- Other Please specify: _____

Reported Adverse Events (see Key below):

Adverse Event Code	Date Reported (mmm/dd/yyyy)	Grading of Event Code	Notes

Key

Adverse Event Code:

- A = Hospitalization related to severe adverse event
 - B = Hypotension (systolic BP <90mmHG) related to severe adverse event
 - C = Loss of Consciousness related to severe adverse event
 - D = Anaphylaxis** related to severe adverse event
 - E = Grade 4 Toxicity related to severe adverse event
 - F = Serious adverse event that resulted in death
 - G = Serious adverse event that was life threatening
 - H = Serious adverse event that required hospitalization or prolongation of existing hospitalization
 - I = Serious adverse event that resulted in persistent or significant disability/incapacity
 - J = Serious adverse event that resulted in a congenital anomaly/birth defect
 - K = Serious adverse event → Pregnancy during treatment
 - L = Hepatotoxicity with LFTS greater than 5x upper limit of normal
- ** defined as at least one major dermatological reaction and at least one cardiovascular and/or respiratory criteria

Grading of Adverse Events:

- 3 = Severe inability to work or perform normal daily activity
- 4 = Life Threatening or disabling event that represented an immediate threat to life
- 5 = Death related to the adverse event