

AA-Clozapine Patient Care Network - PATIENT REGISTRATION

Phone: 1-877-276-2569 / Fax: 1-866-836-6778 / Website: www.aaspire.ca

1165 Creditstone Rd., Unit 1, Vaughan, Ontario L4K 4N7

19-AA054_AAC0350E1

check one:	for registering the Patient in the AA-Clozapine Patient Care Network. Please ne Patient Restart Modify currently registered AA-Clozapine Patient Discontinuation
1 PATIENT REGISTRATION (MASS TRANSFER SEE ATTACHED)	2 PATIENT'S TREATMENT RESOURCE TEAM - PHARMACIST REGISTRATION
Initials: Date of Birth: / / DD MMM YYYY Sex: M F Other: Ethnicity: Caucasian Asian Black Other (specify): No interruption in treatment Health Card #: Monitoring Frequency: Weekly Biweekly Every Four Weeks	Pharmacist Name: Pharmacist License #: APA Designation: Y N Pharmacy Name: Address: Prov: Postal Code:
3 PATIENT'S TREATMENT RESOURCE TEAM - PHYSICIAN REGISTRATION	City: Tel: Fax: Fax:
Physician Name: Physician License #: Facility Name:	Email: I confirm that all dispensing pharmacists at this location will only dispense AA-Clozapine at the specified frequency upon confirmation that the patient has had his/her blood drawn for a Complete Blood Count and differentials for the current period. If applicable, I also confirm responsibility for all actions undertaken by the website login. Date: / / / Pharmacist's Signature:
Address:Prov: Postal Code: City:	4 PATIENT'S TREATMENT RESOURCE TEAM - LABORATORY AND COORDINATOR REGISTRATION
Tel: Email: Statement by Treating Physician I, the treating physician, authorized nurse practitioner or authorized pharmacist* will ensure that blood testing (white blood cell count and differential) for these patients (identified as per the list attached) as required by the AA-Clozapine Product Monograph is performed at the specified frequency. I understand that no pharmacy will dispense any brand other than AA-clozapine to my patients	Laboratory Name: Address/City: Tel: Fax: Coordinator Name:
without my prior knowledge and permission regarding which brand is being dispensed. In this way I will be able to inform the laboratory to send my patients results to the appropriate manufacturer's clozapine database (AA-Clozapine Risk Management Program). I will not prescribe AA-Clozapine until the non-rechallengeable status of these patients has been verified. I have informed the patients and they have not objected to the release of relevant safety information held within a clozapine database to any other clozapine database of an approved manufacturer of clozapine in Canada, if needed for the safe utilization of this medication and/or for the continuous monitoring of these patients. The information which may be released, includes the non-rechallengeable hematological status of the patient, white blood cell counts and absolute neutrophil counts, dates and other information as may be relevant to the safe treatment of the patient with clozapine. *In selected provinces, according to the College of Nurse, College of Pharmacy guidelines/regulations for applicable provinces — By selecting this box, I authorize the laboratory to release to AA-Pharma (1-866-836-6778) all hematological CBC and	Address/City: Tel: Fax: Ext:
Date: / / differential lab results for this patient. Physician's Signature:	PLEASE NOTE: Please ensure AA Pharma is placed on the Standing Order/Requisition form so that we receive copies of the CBC results for the patient TO COMPLETELY PROCESS NEW PATIENTS AND RESTARTS, CBC RESULTS WITH DIFFERENTIALS WITHIN THE LAST 30 DAYS ARE REQUIRED
DISCLOSURE I have reviewed and understand the AA-CLOZAPINE product monograph. (d) 1, the Physician, will only prescribe AA-CLOZ	

- b) I understand that death can occur as a result of agranulocytosis with the use of AA-CLOZAPINE, and that all patients on AA-CLOZAPINE must be enrolled in the AA-CLOZAPINE Patient Care Network to help reduce the risk of a nonrechallengeable patient reusing AA-CLOZAPINE. Iunderstand that patients placed on the nonrechallengeable list have had previous unacceptable WBC counts, and/or ANC values, and/or Clozapine induced myocarditis as defined in the AA-CLOZAPINE product monograph.
- c) I understand that the patient's rechallengeable status will be verified prior to the initiation of treatment for all patients that are new to treatment, or for patients with an unknown or interrupted history on Clozapine.
- from the AA-CLOZAPINE Patient Care Network.
- (e) I agree to notify the AA-CLOZAPINE Patient Care Network of any discontinued patients or interruptions in AA-CLOZAPINE therapy.
- (f) I, the Physician, will ensure that if the patient is female, she is not pregnant nor breastfeeding.
- (g) I, the Pharmacist, will only dispense AA-CLOZAPINE following the receipt of aPIN number from the AA-CLOZAPINE Patient Care Network.
- (h) I, the Physician agree to ensure that hematological testing is performed at the required frequency (as per the product monograph) and submit copies of all lab reports indicating WBC and ANC results to AA Pharma within 7 days.
- (i) I agree to submit the four required weekly lab reports containing WBC and ANC counts after a patient discontinues Clozapine therapy.

 (j) I understand that the AA-CLOZAPINE Patient Care Network will monitor compliance with
- reporting requirements and will notify the patient's physician and/or pharmacist of any discrepancies or overdue lab reports.
- (k) In the event of agranulocytosis, clozapine induced myocarditis, or any other serious event, including any lack of drug effect and any clozapine related hospitalization I agree to fill out the required Serious Adverse Event form(s) and send this form(s) directly to AA-Clozapine Patient Care Network within 24 hours via fax to 18668366778 or report via phone to 18772762569