

DATA TO BE SUBMITTED RELATED TO THE STANDARD OF PRACTICE FOR THERAPEUTIC PRESCRIBING

1. Prescribing, Dispensing, Compounding and Selling Drugs

Registrants must submit the following data related to prescribing, dispensing, compounding or selling drugs in their individual practices, including all practice locations:

- Number of drugs prescribed to patients,
- Number of drugs dispensed to patients,
- Number of drugs compounded for patients,
- Number of drugs sold to patients,
- Number of Adverse Occurrences when a patient has been prescribed, dispensed, compounded or sold a drug. An adverse occurrence would be any of the following events if they are known:
 - Patient referred to emergency services within 5 days,
 - An emergency drug had to be administered to the patient,
 - Patient subsequently diagnosed with shock or convulsions within 5 days,
 - Patient's condition did not improve or worsened,
 - An unscheduled treatment had to be provided to the patient,
 - Patient had an adverse drug reaction.
- From the following list, which types of adverse occurrences have been encountered in your practice (see list in prior bullet),
- Conditions that emerged requiring an unscheduled treatment (see additional data sets below for details), and
- Types of adverse drug reactions encountered, if applicable (see additional data sets below for details).

2. Administering substances by Injection

Registrants must submit the following data related to administering a substance by injection (non-IVIT), from their individual practices, including all practice locations:

- Number of injections administered to patients,
- Number of patients who experienced an Adverse Occurrence (see above for definition of adverse occurrences),
- Types of Adverse Occurrences encountered (see above for definition of adverse occurrences),
- Type of infection encountered, if applicable (see additional data sets below for details),
- Conditions for which an unscheduled treatment was required, if applicable (see additional data sets below for details), and
- Type of adverse substance reactions encountered (see additional data sets below for details).

3. Administering a substance by Inhalation

Finally, Registrants must submit the following data related to administering a substance by inhalation, from their individual practices, including all practice locations:

- Number of inhalations administered to patients,
- Number of patients who experienced an Adverse Occurrence (see above for definition of adverse occurrences),
- Types of Adverse Occurrences encountered (see above for definition of adverse occurrences),
- Type of infection encountered, if applicable (see additional data sets below for details),
- Conditions for which an unscheduled treatment was required, if applicable (see additional data sets below for details), and
- Type of adverse drug reactions encountered (see additional data sets below for details).

Additional Data Sets

The College data collection will include the following additional data:

- Where an infection has occurred, the type of infection seen from the following pick list, although number of instances will not be required:
 - Bronchitis,
 - Cystitis,
 - Gastroenteritis,
 - Influenza,
 - Pharyngitis,
 - Pneumonia,
 - Sinusitis.
- Where an unscheduled treatment has occurred, the type of condition treated from the following pick list, although number of instances will not be required:
 - Anxiety,
 - Dizziness,
 - Headache,
 - Fatigue,
 - Low back strain,
 - Muscular spasms,
 - Phlebitis,
 - Pneumonia.
- Where an adverse drug/substance reaction has occurred, the type of reaction from the following pick list, although number of instances will not be required:

- Anxiety,
- Diarrhea,
- Headache,
- Hypertension,
- Hypoglycemia.
- Injection site extravasation,
- Maculo-papular rash,
- Nausea,
- Phlebitis,
- Pre-syncope,
- Seizure,
- Shortness of breath,
- Syncope,
- Urticaria,
- Vomiting.