Notice from the Executive Officer: Updates to the Ontario Guidelines for Drug Submission and Evaluation

- Effective July 1st, 2019, the ministry will no longer require drug manufacturers to provide a full printed copy of their submission.
- The Drugs and Devices Division will accept one electronic copy of the submission for all product submissions (including but not limited to Single Source Drug Product submissions, Multiple Source Drug Product submissions, Valved Holding Chamber submissions, Diabetic Testing Agent submissions, and Nutrition Product submissions.)

The Drugs and Devices Division is working to improve efficiencies and to reduce burden on manufacturers for drug submission requirements.

The updated policy will also be available on the ministry's website the week of July 1st, 2019.

Ontario