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Product information

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The product monograph is developed by a drug sponsor according to guidelines published by Health Canada that provide direction on the content and format. The veterinary labelling is developed by the drug sponsor according to the Food and Drug Regulations. While Health Canada reviews the product monograph or the veterinary labelling as part of the drug review process, it remains the responsibility of the drug sponsor to ensure that the product monograph or the veterinary labelling is complete and accurate.

Current status: **Marketed****Current status date:** 2013-06-11**Original market date:** [1](#) 1998-01-05**Product name:** ACET 650

Help on accessing alternative formats, such as Portable Document Format (PDF), Microsoft Word and PowerPoint (PPT) files, can be obtained in the [alternate format help section](#).

DIN: 02230437**Product Monograph/Veterinary** **Date:** 2017-07-19**Labelling:**  [Product monograph/Veterinary Labelling \(PDF version ~ 175K\)](#)**Company:** [PENDOPHARM DIVISION OF PHARMASCIENCE INC](#)

100 6111 Royalmount Ave

Montreal

Quebec

Canada H4P 2T4

Class: Human**Dosage form(s):** Suppository**Route(s) of administration:** Rectal**Number of active ingredient(s):** 1**Schedule(s):** OTC**Anatomical Therapeutic Chemical (ATC):** [4](#) N02BE01 ACETAMINOPHEN (PARACETAMOL)**Active ingredient group (AIG) number:** [5](#) 0102009008

List of active ingredient(s)

Active ingredient(s)	Strength
ACETAMINOPHEN	650 MG

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Footnotes

- 1 The earliest marketed date recorded in the Drug Product Database.
- 4 The purpose of the World Health Organization (WHO) Anatomical Therapeutic Chemical (ATC) classification system is to be used as a tool for drug utilization research in order to improve quality of drug use. Drugs are divided into different groups according to the organ or system on which they act and their chemical, pharmacological and therapeutical properties.
- 5 The AIG number is a 10 digit number that identifies products that have the same active ingredient(s) and ingredient strength(s). The AIG is comprised of three portions:
 - the first portion (2 digits) identifies the number of active ingredients,
 - the second portion (5 digits) identifies the unique groups of active ingredients(s),
 - the last portion (3 digits) identifies the active ingredient group strength. The strength group has a tolerance of -2% to +10%.

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