

Government of Canada

Gouvernement du Canada

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> Reports and Publications - Drugs and Health Products > Compliance and Enforcement

Inspection tracker: Drug manufacturing establishments

As part of Health Canada's ongoing commitment to openness and transparency, the Department is publishing information regarding emerging issues identified through our drug inspection program.

This tracker provides a snapshot of the potential health and safety issues Health Canada is tracking with companies that fabricate, package/label, test, wholesale, distribute or import drugs for sale in Canada. The information in the chart will expand to eventually include details about affected Canadian companies and products.

How the inspection tracker works

- Health Canada responds to potential risks as soon as we hear about them whether from our own inspections, the companies themselves, adverse reaction reports or from our regulatory partners.
- The tracker highlights actions Health Canada is taking such as: requests for voluntary quarantine, stop sales, import restrictions, or product recalls. It also indicates those circumstances where no action has been warranted.
- Even if a company is listed on the tracker, it doesn't necessarily mean there is an immediate risk to the health of Canadians. It means Health

Canada is looking into a potential issue.

- Health Canada will continue to take action to manage risks identified to the health of Canadians using the most appropriate level of intervention, proportional to the risk to health.
- This tracker currently doesn't list specific drugs and health products affected but links to <u>Recalls and Safety Alerts</u> if a risk has been linked to specific products on the Canadian market.
- This tracker is updated regularly.
- Once an item is listed in the <u>closed section</u>, it indicates that there is no further action being taken by Health Canada at this time and there will be no more updates regarding this item.
- Last update: July 30, 2024

Open items

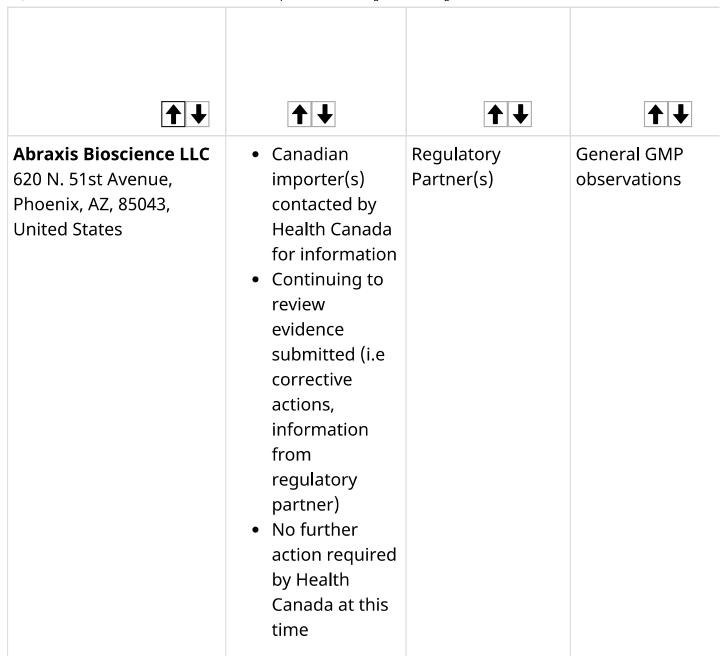
Establishment	Status of issue	Source of information under review	Primary reason for action
Spectrum Chemical Mfg Corp. 777 Jersey Avenue, New Brunswick, NJ, 08901, United States New	 Continuing to review evidence submitted (i.e corrective actions, information from regulatory partner) 	Regulatory Partner(s)	General GMP observations

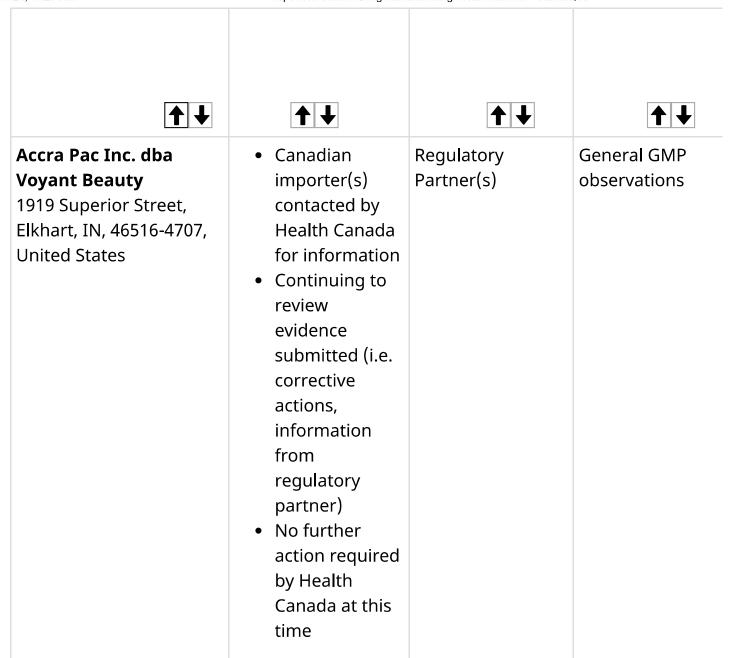
Establishment	Status of issue	Source of information under review	Primary reason for action
Sri Krishna Pharmaceuticals Ltd. Unit II, A-34 & A-35, IDA, Nacharam, Medchal- Malkajgiri District, Hyderabad, Telangana, India, 500076	 Canadian importer(s) contacted by Health Canada for information Continuing to review evidence submitted (i.e corrective actions, information from regulatory partner 	Regulatory Partner(s)	General GMP observations
Sun Pharmaceutical Industries Limited Survey No. 1012, Dadra, Union Territory of Dadra & Nagar Haveli and Daman & Diu, 396 193, India	 Canadian importer(s) contacted by Health Canada for information Continuing to review evidence submitted (i.e corrective actions, information from regulatory partner) 	Regulatory Partner(s)	General GMP observations

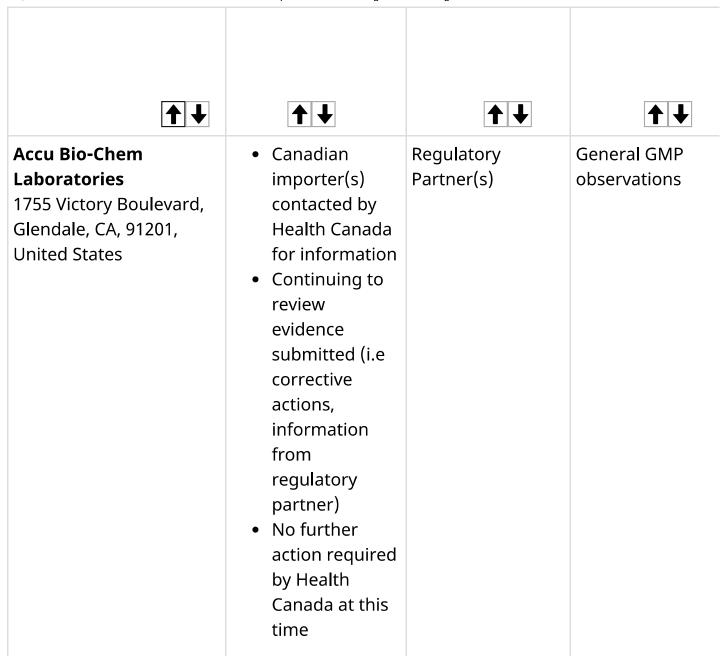
Filter items	Showing 1 to 10 of 234 entries			
	 Show	10	~	entries

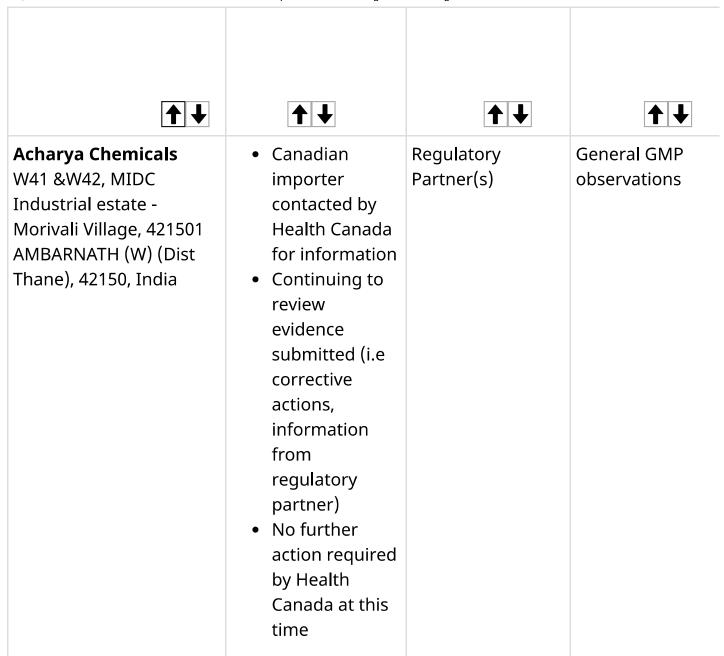
Closed items

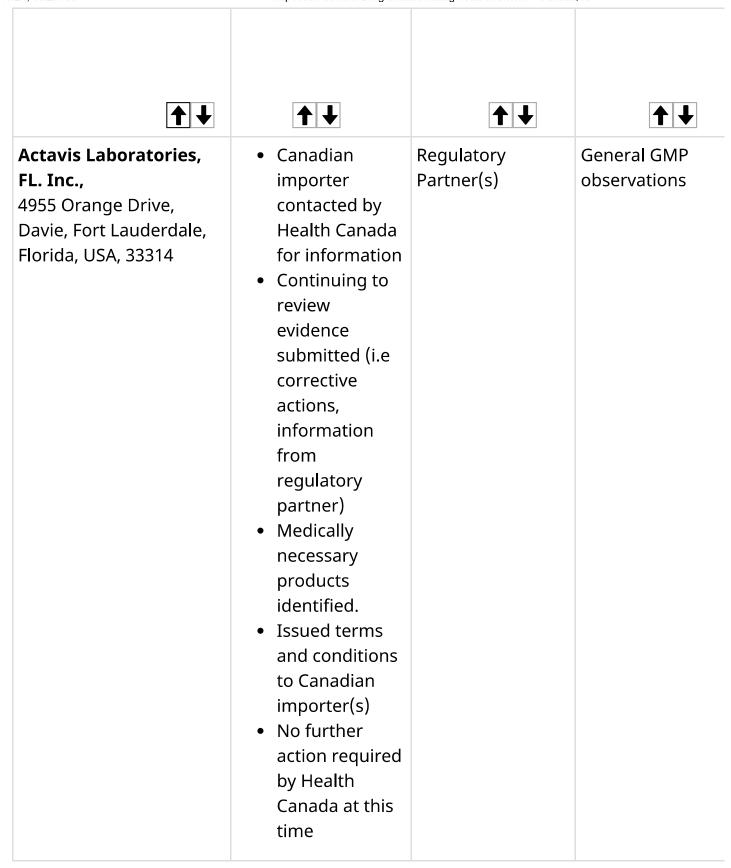
		Source of information	Primary
	Status of	under	reason for
Establishment 🚹 🖶	issue ↑↓	review ↑ ↓	action ↑ ↓

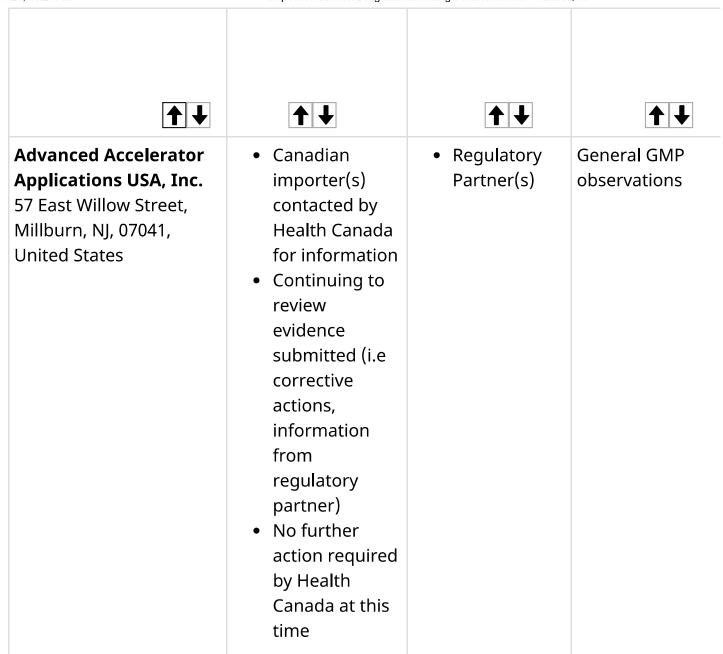


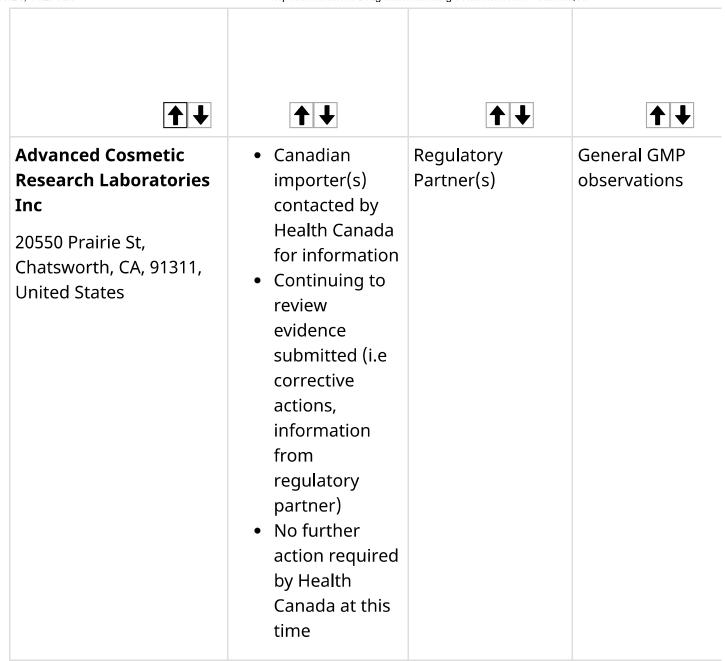


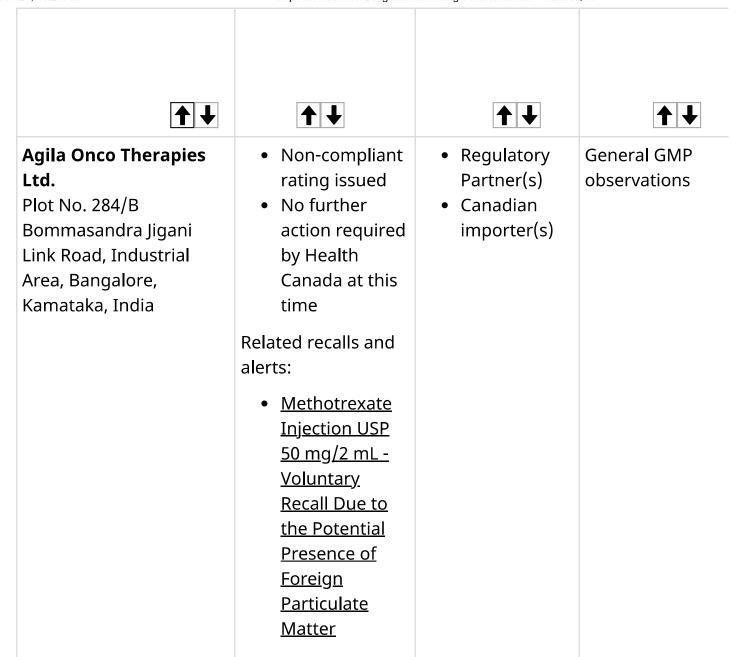












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AGILA SPECIALTIES PRIVATE LTD. Opp II M, Bilekahalli, Bannerghatta Road, Bangalore, Karnataka, India - 560 076	 Canadian importer(s) contacted by Health Canada for information No critical risks identified to date No further action required by Health Canada at this time 	Regulatory Partner(s)	General GMP observations









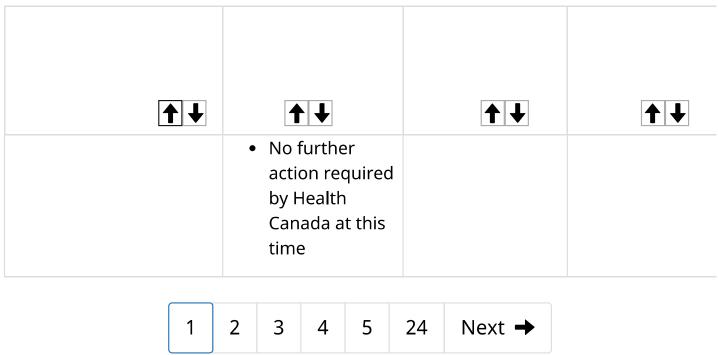
AGILA SPECIALTIES PRIVATE LTD., SPECIALTY FORMULATION FACILITY (SFF)

19A, Plot No. 284-B/1 Bommasandra Jigani Link Road, Anekal Taluk, Bangalore India - 560 105

- Canadian importer(s) contacted by Health Canada for information
- Continuing to review evidence submitted (i.e corrective actions, information from regulatory partner)
- Medically necessary products identified
- Requested voluntary quarantine of non-medically necessary products
- Voluntary quarantine in place
- Issued terms and conditions to Canadian importer(s)

Regulatory Partner(s)

General GMP observations



Date modified:

2024-07-30