

## Pharma Medica Research Inc. Successfully completes another US FDA Inspection!

**Toronto, Canada (July 14, 2017)** – Pharma Medica Research Inc. (PMRI) is extremely pleased to announce that the US FDA has concluded a 4-day intense inspection of the clinical unit in St. Charles, Missouri, USA with no 483 Form issued. The Inspector performed an extensive review of documentation from two recent studies in addition to SOPs and training & equipment records. The Inspector indicated that the records are in very good order, there are strong documentation practices and summarized the audit as being very successful!

It's a very exciting time at PMRI as they celebrate their 20<sup>th</sup> anniversary. Over the years, their reputation has been built upon principles of delivering uncompromised quality.

This follows the two most recent US FDA audits of both Canadian facilities which also concluded with no 483 findings! These inspections, which took place over a 3 week period, included full inspections of the clinical and bioanalytical operations in addition to the Quality Systems.

*"We are very proud of this outcome which is a true testament to our continued commitment to exceptional quality and is a direct result of the dedication of our staff and strong leadership & direction from our Senior Management Team"* stated Latifa Yamlahi, President & CEO.

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### **About Pharma Medica Research Inc.**

Pharma Medica Research Inc. (PMRI) is a privately owned, full-service contract research organization that has been successfully audited by many regulatory agencies including FDA, EMA, MHRA, AGES, Health Canada and ANVISA. Over the past 20 years, PMRI has specialized in conducting complex early-phase clinical trials in healthy volunteers, special and patient populations. To date, the company has completed more than 4,000 clinical trials intended for multiple regulatory bodies around the world. These include studies across various therapeutic areas for New Chemical Entities (NCEs), hybrid and generic submissions. With sites in Canada and the USA, all study needs can be fulfilled in-house including clinical conduct, bioanalysis, clinical data management and full scientific affairs services.

To learn more or contact PMRI, please visit [www.pharmamedica.com](http://www.pharmamedica.com).