



NAME: \_\_\_\_\_

HSN: \_\_\_\_\_

D.O.B.: \_\_\_\_\_

- RUH     SCH     SPH
- PH     RGH     WRC     Other \_\_\_\_\_

**HEALTH INFORMATION MANAGEMENT SERVICES  
CLINICAL TRIAL HEALTH RECORD RETENTION**

This individual participated in a pharmaceutical research study. Health Canada Food and Drug Regulations – Division 5, C.05.012 require all records be retained for 15 years following completion of the study.

**SECTION BELOW TO BE COMPLETED BY RESEARCHER:**

Study title: \_\_\_\_\_  
\_\_\_\_\_

Study protocol #: \_\_\_\_\_

REB #: \_\_\_\_\_

Date trial started at local site: \_\_\_\_\_  
*(dd/mmm/yyyy)*

Date trial ended at local site: \_\_\_\_\_  
*(dd/mmm/yyyy)*

Saskatchewan Health Authority operational approval to conduct this research was granted on:

\_\_\_\_\_  
*(dd/mmm/yyyy)*

**Contact information (please print)**

Investigator: \_\_\_\_\_

Department: \_\_\_\_\_

Phone #: \_\_\_\_\_

Signature: \_\_\_\_\_  
*(Investigator)*

Date: \_\_\_\_\_  
*(dd/mmm/yyyy)*

**FOR COMPLETION BY HEALTH INFORMATION MANAGEMENT SERVICES (MANAGER AND/OR HIM ANALYST)**

Date to destroy patient chart visit(s): \_\_\_\_\_  
*(dd/mmm/yyyy)*

**KEEP ON TOP RIGHT-HAND SIDE OF THE CHART  
PLACE ON ALL VOLUMES INCLUDED IN THE STUDY PERIOD**