

PREGNANCY INFORMATION FAX FACSIMILE TRANSMISSION		Study #: SAE FAX NO: (301) 230-0159 ALTERNATE FAX NO: (301) 897-7404
Ticket Number: _____		
Initial Report Date: DD - MMM - YY	Follow-up Report Date: DD - MMM - YY	
Principal Investigator:	Reporter:	
Reporter Telephone #:	Reporter FAX #:	
Investigator Number <small>Complete all of the investigator and subject number boxes provided. Use leading zeros, when necessary, to complete all expected boxes.</small> Example: Investigator #407 would be filled in as: 0 0 4 0 7	Subject Number	Subject Initials <small>Record the first letter of the subject's first, middle and last name, in that sequence. If the subject has no middle name, enter a dash.</small> Example: A - C
Subject's Sex: <input type="checkbox"/> Female <input type="checkbox"/> Male	Subject's Weight: _____ kg	Subject's Date of Birth: DD - MMM - YYYY
Subject's Ethnicity (check one only): <input type="checkbox"/> Hispanic or Latino <input type="checkbox"/> Not Hispanic or Latino <input type="checkbox"/> Not Available		
Subject's Race (check all that apply): <input type="checkbox"/> American Indian or Alaska Native <input type="checkbox"/> Asian <input type="checkbox"/> Black or African American <input type="checkbox"/> Native Hawaiian or Other Pacific Islander <input type="checkbox"/> White <input type="checkbox"/> Not Available		
Study Drug:	Study Drug Start Date: DD - MMM - YY	Study Drug Stop Date: DD - MMM - YY OR <input type="checkbox"/> Study Drug Continuing
Dose:	Route: ORAL	Frequency: QD Kit #:
First Day of Last Menstrual Period: DD - MMM - YY	Estimated Date of Delivery: DD - MMM - YY	
Method of Contraception (check all that apply): <input type="checkbox"/> Oral Contraceptive Pills <input type="checkbox"/> Condoms <input type="checkbox"/> Periodic Abstinence <input type="checkbox"/> Progestin Injection or Implants <input type="checkbox"/> Spermicide <input type="checkbox"/> Diaphragm <input type="checkbox"/> Intrauterine Device (IUD) <input type="checkbox"/> Tubal Ligation <input type="checkbox"/> Other, specify: _____		
Reproductive History: <input type="checkbox"/> Gravida _____ <input type="checkbox"/> Para _____		
Tests performed during pregnancy: <input type="checkbox"/> None <input type="checkbox"/> Unknown <input type="checkbox"/> CVS Results: <input type="checkbox"/> Normal <input type="checkbox"/> Abnormal <input type="checkbox"/> Amniocentesis Results: <input type="checkbox"/> Normal <input type="checkbox"/> Abnormal <input type="checkbox"/> Ultrasound Results: <input type="checkbox"/> Normal <input type="checkbox"/> Abnormal		
Pregnancy Outcome Was pregnancy interrupted? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, specify: <input type="checkbox"/> Elective Termination <input type="checkbox"/> Spontaneous Abortion <input type="checkbox"/> Ectopic Date of Termination: DD - MMM - YY If pregnancy was not terminated, specify pregnancy outcome (and provide infant outcome information) <input type="checkbox"/> Vaginal Birth: <input type="checkbox"/> Premature <input type="checkbox"/> Term OR <input type="checkbox"/> C-Section: <input type="checkbox"/> Scheduled <input type="checkbox"/> Emergency Date of Delivery: DD - MMM - YY Infant outcome information: <input type="checkbox"/> Normal <input type="checkbox"/> Abnormal		
Additional Case Details (if needed):		
NOTE: For an initial reporting fax both the Pregnancy Report CRF and Additional Pregnancy Information Fax Page. For follow-up reporting, fax only the Additional Pregnancy Information Fax Page.		

NOTE: The patient should have appropriate follow-up as deemed necessary by their physician.
If the the baby is born with a birth defect or anomaly, then a second AdEERS report is required.