

Site Number: _____

Screening ID: _____ - ____

Participant Letters: _____

Complete this form upon confirmation that a study participant is pregnant, regardless of assigned treatment group. No further study medication should be given.

Additional form(s) that need to be completed:

- Adverse Event Report Form (RIT13)

- Pregnancy Outcome Report Form (RIT14R)*

* When pregnancy has ended

A. REPORT INFORMATION

Pregnancy Identification Number: _____

1. Report Date:

____ / ____ / ____
DAY MONTH YEAR

2. Last attended study visit prior to the confirmed pregnancy:

- | | | | | |
|-------------------------------------|-------------------------------------|--------------------------------------|--------------------------------------|-------------------------------------|
| <input type="checkbox"/> 2 Baseline | <input type="checkbox"/> 10 Week 10 | <input type="checkbox"/> 18 Month 12 | <input type="checkbox"/> 29 Month 24 | <input type="checkbox"/> 99 Other |
| <input type="checkbox"/> 3 Week 1 | <input type="checkbox"/> 11 Month 3 | <input type="checkbox"/> 21 Month 13 | <input type="checkbox"/> 30 Month 30 | <input type="checkbox"/> 98 PhiX174 |
| <input type="checkbox"/> 4 Week 2 | <input type="checkbox"/> 15 Month 5 | <input type="checkbox"/> 26 Month 15 | <input type="checkbox"/> 31 Month 36 | Visit ONLY |
| <input type="checkbox"/> 5 Week 3 | <input type="checkbox"/> 16 Month 6 | <input type="checkbox"/> 27 Month 18 | <input type="checkbox"/> 32 Month 42 | |
| <input type="checkbox"/> 6 Week 5 | <input type="checkbox"/> 17 Month 9 | <input type="checkbox"/> 28 Month 21 | <input type="checkbox"/> 33 Month 48 | |

a. If OTHER, specify date of visit:

____ / ____ / ____
DAY MONTH YEAR

b. If PhiX174 Visit ONLY (i.e. Weeks 6, 7, 8, 13, 14, 16, 53, 54, 58, 59, 60, or 62),
Record week number: _____

B. PREGNANCY INFORMATION

1. Date of positive pregnancy test:

____ / ____ / ____
DAY MONTH YEAR

2. Date of last menstrual cycle:

____ / ____ / ____
DAY MONTH YEAR

3. Estimated date of delivery:

____ / ____ / ____
DAY MONTH YEAR

4. Is the participant planning on carrying the pregnancy to term?

Y N

5. Is the participant willing to continue with future follow-up visits?

Y N

6. Has the participant's obstetric care provider been informed of her participation in this study?

Y N

C. PREGNANCY HISTORY

1. Record total number of prior pregnancies (not including this one):

2. Has the participant ever had a pregnancy complication?

Y N

If YES,

a. Has the participant ever had a miscarriage?

Y N

b. Has the participant ever had a pregnancy that resulted in a stillbirth?

Y N

c. Has the participant ever had a pregnancy result in neonatal death?

Y N

d. Has the participant ever had a pre-term delivery (< 37 gestational weeks)?

Y N

e. Has the participant ever had a post-term delivery (> 42 gestational weeks)?

Y N

Initials (first, middle, last) of person completing this form:

F M L

Date form completed:

____ / ____ / ____
DAY MONTH YEAR

On all questions write "?" if the desired information is currently unavailable, but is being checked and will be known in future updates. Write "" if the desired information is permanently unavailable (i.e. will not be known in any future updates).*



Anti-CD20 Study
PREGNANCY CONFIRMATION FORM

Form RIT14

15 MARCH 2006

Version 1.0

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