

Anti-CD20 Study PREGNANCY OUTCOME REPORT FORM

Form RIT14R

15 MARCH 2006 Version 1.0 Page 1 of 2

			Page 1 of 2
Site Number:	Screening ID:	 Participant Letters:	

Complete this form when the outcome of an active pregnancy becomes known. Complete this form for all participants that become pregnant during the course of the trial.

A. PREGNANCY OUTCOME INFORMATION					
1. Record the Pregnancy Identification Number:					
The Pregnancy Identification Number is located on the subject's initial Pregnancy	Confirmation For	n (RI '	Γ14)		
2. Is the outcome of the pregnancy unknown due to loss of participant to follow-up	?	Y	N		
If YES, STOP HERE					
3. Date pregnancy ended:	DAY MONTH	/— <u></u>	 EAR		
4. Was the pregnancy terminated as a result of an induced abortion?	DAT MONTH	Y	N		
If YES,					
a. Was the reason for the abortion medically indicated?		Y	N		
If YES, Complete Adverse Event Report Form (RIT13)					
1) Specify reason:					
5. Did the pregnancy result in a miscarriage? Complete Adverse Event Report Fo	orm (RIT13)	Y	N		
6. Did the pregnancy result in a live birth or multiple live births?					
7. Did the pregnancy result in a stillbirth?					
If YES, Complete Adverse Event Report Form (RIT13)					
a. Did the stillbirth have any congenital malformations?					
If YES,					
1) Specify:					
b. Did the stillbirth have any other complications?					
If YES,					
1) Specify:					
8. Record number of infants (both living and deceased) the birth resulted in:		_	_		
9. Were there any complications during the delivery?		Y	N		
10. Was an HbA1c measured at any time during the pregnancy?					
If YES,					
a. Record HbA1c:			%		
b. Date measured:	/	/— 			
11. Is the participant currently breastfeeding?	DAY MONTH	Y	EAR N		
11. 15 the participant currently breastreeding:		1	14		

Complete Section B to record the details of any live birth(s).

B. INFANT INFORMATION

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).

Diabetes TrialNet

Anti-CD20 Study PREGNANCY OUTCOME REPORT FORM

Form RIT14R

15 MARCH 2006 Version 1.0 Page 2 of 2

Site Number:	Screening ID:	Particip	ant Letters:				
1. Indicate the Pregnancy Identification Number:							
2. Birth Order:	01	0 2	03†				
3. Sex (M/F):	M F	M F	M F				
4. Gestational age:	wks	wks	wks				
5. Birth weight:	————gm OR	——— gm OR	gm OR				
	lbs oz	lbs oz	lbs oz				
6. 1 minute APGAR score:							
7. 5 minute APGAR score:							
8. Was the infant born with any congenital malformations?	Y N	Y N	Y N				
a. If YES*, specify:							
9. Was the infant born with other complications?	Y N	Y N	YN				
a. If YES*, specify:							
10. Was the infant admitted to the Neonatal Intensive Care Unit (NICU) at any time*?	Y N	Y N	Y N				
11. Was the infant discharged from the hospital alive? If YES,	Y N	Y N	Y N				
a. Date discharged:	DAY MONTH YEAR	DAY MONTH YEAR	DAY MONTH YEAR				
If NO*,							
b. Date of death:	DAY MONTH YEAR	DAY MONTH YEAR	DAY MONTH YEAR				
c. Specify cause of death:							
* Requires completion of an Adverse Event Report Form (RIT13)							
If more space is needed, attach additional copies of the second page of this form							
Initials (first, middle, last) of person completing this form: ${F} {M} {L}$							

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).

Date form completed:

DAY MONTH