TrialNet		CD20 Study Y EVENT FORM		m RIT13M 5 MARCH 2006 Version 1.0 Page 1 of 2
Site Number:	Screening ID:		Participant Letters:	e

Complete this form if a participant dies during the study, <u>regardless</u> of whether the death was <u>related</u> to the study medication.

- This form should be sent to the Coordinating Center within 24 hours of notification of the death.
- Once a death certificate has been obtained, a copy MUST be sent to the Coordinating Center.

Additional form(s) that need to be completed:			Documentation that needs to be obtained:								
- Adverse Event Report Form (RIT13)			Death Certificate (when available)Autopsy report (when available)								
						pore (ma					
A. REPORT IN	NFORMATION										
1. Date of re	port:					- $ -$	MONTH /	YEA	 R		
2. Date of de	eath:					- DAY $/$ -	/ MONTH				
3. Type of report:						Initial		Follow			
B. GENERAL	EVENT CLASSIFICATION										
1. Where did	the death occur? (check one)										
	\Box_1 Hospital			\square_4	Long-term care institution						
					Unknown						
	School/Work			99	Other						
If OTHE a. Specif											
-	-										
\square_1	2. The death was (<i>check one</i>): \Box_1 Sudden, explained					Followi	ng illness				
\square_2	Sudden, unexplained				_ 5						
3. At the tim	e of onset of the terminal event,	the p	artici	pant was	(check o	ne):					
					rate phy	vsical ac	tivity				
	1				D ₅	Engaged in heavy physical activity					
	Engaged in light physical activ	ity			9	Unknov	vn				
4. Was the participant receiving study medication at the time of the death event?						Y	Ν				
5. Will an autopsy report be available?							Y	Ν			
6. Has a death certificate been obtained?							Y	Ν			
If NO,											
	ne been requested?								Y	Ν	
7. Record the	e sources of information that we	re use	ed to o	complete	this form	n:					
a. Death	certificate?	Y	Ν	d. Iı	nterview	of attend	ling physi	cian?	Y	Ν	
b. Autopsy report?		Y	Ν	e. Ir	nterview	w of family member?		Y	Ν		
c. Hospi	tal report on fatal illness?	Y	Ν	f. 0	ther?				Y	Ν	
If OTHE											
1) Speci	fy:										

C. SPECIFIC EVENT INFORMATION

1. Describe the immediate cause of death:

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).

TrialNet	Anti-CD20 Study MORTALITY EVENT FOR	RM Form RIT13N Version 1.
Site Number:	Screening ID:	Page 2 of Participant Letters:
2. Describe the underlyin	g cause of death:	
3. Describe any contribu	tory causes of death:	
	mmediate, underlying and/or contributory of	causes of death were present at
randomization:		
	Initials (first, middle, last) o	of person completing this form: $\frac{1}{F} \frac{1}{M}$
	Date for	m completed: $\frac{1}{DAV} = \frac{1}{MONTH} = \frac{1}{VEAP}$
		DAY MONTH YEAR
Signature of Principal	nvestigator	

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).