

SERIOUS ADVERSE EVENT REPORTING FORM

Initial Reporting: Once you have become aware of a SAE or SUSAR, please email this form to the BAMI Trial Office (BTO) [e-mail: bami@qmul.ac.uk] within 24 hours of learning of the event. "SAE report" must be in the email's subject line. The form should be signed by the Pl/delegated doctor, scanned and e-mailed to the BTO. If that could not be achieved, a signed copy of the same form (without editing and additional information) must be sent to the BTO at the earliest opportunity by fax [+44 20 3465 6491]. Once this form is received by the BTO an acknowledgment email will be sent to your site during office hours. If you do not receive an acknowledgement on the following working day, please contact the BTO [Tel: +44 20 3765 8740].

Full title of the study	The effect of intracoronary reinfusion of bone marrow-derived mononuclear cells (MN-MNC) on all cause-mortality in acute myocardial infarction		
Study Acronym	BAMI		
Name of sponsor	Queen Mary University of London		
Sponsor R&D Number	EudraCT Number: 2012-001495-11		
MREC Number	13/LO/0315		
Chief Investigator	Name:	Prof Anthony Mathur	
	Phone No:	(+44) 20 3765 8715	
	Email address:	a.mathur@qmul.ac.uk	
Is this a double blind study?	No		
Name of IMP	T2c001 (Bone marrow derived mononuclear cells (BM-MNCs))		

The following sections should be completed by the SITE.

The following sections should be completed by the off E.				
Report type:				
Subject identification code:	Patient/initials (first, last):			
	john, hawking			
Date of Birth: (Day/Month/Year)	Patient's Age:	Sex:		
23 /11 /1987	31	⊠M □ F		
Principal Investigator				
Name:	Phone No:	Email address:		
Aalap	7895432176			
Trial Co-ordinator at local site:				
Name:	Phone No:	Email address:		
Mark				
Reporting host institution				
Trust/ Institution name:		Site number:		
Walt Blood testing institute		49		
Date of site become aware of event:	Onset date of SAE:	Resolution date of SAE:		
30 /12 /2006	30 /12 /2006	01 /02 /2007		
Event description (eg. body site, symptoms): Please use separate form for each event patient suffering from sleepind disorder which is of type Narcolepsy.				





Type of SAE		Severity		
□ F	Results in Death	Mild		
□ L	_ife threatening	☐ Moderate		
	Hospitalisation or prolongation of nospitalisation	Severe Severe		
Ш i	Persistent or significant disability or ncapacity			
\boxtimes	Congenital anomaly or birth defect			
	Other" important medical event			
ľ	f "Other", please describe:			
Bone	marrow derived mononuclear cells re-info	usion		
	IMP (t2c001) infused?	Re-infusion Date:	Re -Infusion Time:	
□ Y	es 🛛 No	/ /	:	
			24 hour clock	
Causal relationship to IMP		Causal relationship to intervention		
Is the SAE likely to be related to the IMP t2c001?		Does the SAE have a causal relationship to the intervention?		
⊠ F	Related Unrelated	⊠ Bone Marrow Aspiration	on	
_		☐ Cardiac Catheterisation		
∐Pa	atient in control arm	Not related to invasive procedure		
	0.5	No intervention taken		
Is the SAE expected? (Expected reactions will be found in the Investigator Brochure and protocol)		Is the SAE due to the progression of an underlying illness?		
⊠ E	Expected Unexpected	⊠ Yes □ No		
Is the event classified as a SUSAR? (i.e. RELATED to the IMP and UNEXPECTED)		Is the SAE related to the tri	ial CONDUCT?	
□ Y	∕es ⊠ No			
	s, please give IMP batch number			
	01 Batch Number:			
	garded as SUSAR, please also fill in the IS form attached to this form			
Name	es of non IMPs concomitant medicines:	Names of concomitant dise	eases:	
PAN-50		Narcolepsy		
If SAE during index admission, action taken with study treatment		Did the PI withdraw the p	atient from the study?	
	nfused			
	Not Infused	□ No		
⊠ I	nfusion stopped			
	nfusion interrupted			
	Not applicable, please Give details:			



Outcome of SAE						
□ Resolved	☐ Fatal					
Resolved with sequelae	*Date of death:	/ /				
*specify sequelae:	*Copy of post-morte	m available? Yes				
☐ Improved		□ No				
□ Persisting						
Worsened	│					
	UIKIOWII					
Additional information requested by the Cl's team for this project:						
Relevant Medical History :	Relevant Lab Results / Imaging Results (e.g. ECG, blood test):					
	blood test: AIDS					
	l					
Delegated person completing the form if not the PI						
Print name:	Phone No.:	Email address:				
john hawking	8765432189					
Signature:	Job title:	Date:				
	software engineer					
Principle Investigator / Delegated person	,					
Print name:	Phone No.:	Email address:				
george						
Signature:	Date:					
For BAMI Trial Office use only						
Date form RECEIVED by BAMI Trial Office from	Reviewed by:					
Investigator Site:						
31 /12 /2006 (This data will be DAY 1 for SUSARs)	Date reviewed: 22 /5 /2007					
(This date will be DAY 1 for SUSARs)						
Chief investigator reviewed	Fully Agree with Di's seems	-amant				
Date: / /	Fully Agree with PI's assessment					
	∑ Yes □No					
Cl's signature:	If No, please specify					
For R&D Office use only						
Date form RECEIVED by R&D team:	For SUSAR only					
/ /	Date Reported to MHRA:					
Reviewed by:	, ,					
Date reviewed: / /						

