

## 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](mailto: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 PI/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:

Report type: <input checked="" type="checkbox"/> Initial <input type="checkbox"/> Follow-up		
Subject identification code:	Patient/initials (first, last): john, hawking	
Date of Birth: (Day/Month/Year) 23 /11 /1987	Patient's Age: 31	Sex: <input checked="" type="checkbox"/> M <input type="checkbox"/> F

<b>Principal Investigator</b>		
Name: Aalap	Phone No: 7895432176	Email address:
<b>Trial Co-ordinator at local site:</b>		
Name: Mark	Phone No:	Email address:
<b>Reporting host institution</b>		
Trust/ Institution name: Walt Blood testing institute	Site number: 49	
Date of site become aware of event: 30 /12 /2006	Onset date of SAE: 30 /12 /2006	Resolution date of SAE: 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
<input type="checkbox"/> Results in Death <input type="checkbox"/> Life threatening <input checked="" type="checkbox"/> Hospitalisation or prolongation of hospitalisation <input type="checkbox"/> Persistent or significant disability or incapacity <input checked="" type="checkbox"/> Congenital anomaly or birth defect "Other" important medical event If "Other", please describe:	<input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input checked="" type="checkbox"/> Severe

Bone marrow derived mononuclear cells re-infusion		
Was IMP (t2c001) infused?	Re-infusion Date:	Re -Infusion Time:
<input type="checkbox"/> Yes <input checked="" type="checkbox"/> No	/    /	: 24 hour clock

<b>Causal relationship to IMP</b> Is the SAE likely to be related to the IMP t2c001? <input checked="" type="checkbox"/> Related <input type="checkbox"/> Unrelated  <input type="checkbox"/> Patient in control arm	<b>Causal relationship to intervention</b> Does the SAE have a causal relationship to the intervention? <input checked="" type="checkbox"/> Bone Marrow Aspiration <input type="checkbox"/> Cardiac Catheterisation <input checked="" type="checkbox"/> Not related to invasive procedure <input type="checkbox"/> No intervention taken
<b>Is the SAE expected?</b> (Expected reactions will be found in the Investigator Brochure and protocol) <input checked="" type="checkbox"/> Expected <input type="checkbox"/> Unexpected	Is the SAE due to the progression of an underlying illness? <input checked="" type="checkbox"/> Yes <input type="checkbox"/> No
<b>Is the event classified as a SUSAR?</b> (i.e. RELATED to the IMP and UNEXPECTED) <input type="checkbox"/> Yes <input checked="" type="checkbox"/> No If Yes, please give IMP batch number T2c001 Batch Number: *If regarded as SUSAR, please also fill in the CIOMS form attached to this form	Is the SAE related to the trial CONDUCT? <input checked="" type="checkbox"/> Yes <input type="checkbox"/> No
Names of non IMPs concomitant medicines: PAN-50	Names of concomitant diseases: Narcolepsy

If SAE during index admission, action taken with study treatment	Did the PI withdraw the patient from the study?
<input type="checkbox"/> Infused <input type="checkbox"/> Not Infused <input checked="" type="checkbox"/> Infusion stopped <input type="checkbox"/> Infusion interrupted <input type="checkbox"/> Not applicable, please Give details:	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No



