



EuPreCHO Appendix 06 - Case Report Form

EuPreCHO: European study on perioperative management and outcome following Preoperative Transthoracic Echocardiography in noncardiac surgery patients.

	Inclusion criteria		
Elective, inhospit following criteria	tal, intermediate or high-risk noncardiac surgery procedures AND any of the 3	<u>Yes</u>	No
- aged ≥65 y	ears OR	Yes	No
family histo	AND ≥2 of the following: hypertension, smoking, dyslipidaemia, diabetes, or ory of cardiovascular disease (coronary artery disease, cerebral vascular pripheral arterial disease, heart failure) OR	<u>Yes</u>	No
	AND known cardiovascular disease (coronary artery disease, cerebral vascular ripheral arterial disease, heart failure, valvular disease)	<u>Yes</u>	No
	Exclusion criteria		
- under 18 y	years of age	No	Yes
- day surge	ry (outpatient surgery)	No	Yes
- urgent/en	nergency surgery	No	Yes
- current IC (day 0),	U patient (i.e. in ICU day prior of surgery or the day of the index surgery	<u>No</u>	Yes
- cardiac su	rgery within the last month prior to the index noncardiac surgery	<u>No</u>	Yes
- unwilling	or unable to provide informed consent	<u>No</u>	Yes
- unable to	complete the WHODAS questionnaire (literacy or language barrier)	<u>No</u>	Yes
- Previous e	enrollment in EuPreCHO (in case of repeated surgery).	<u>No</u>	Yes

Only IF the first inclusion criterion AND any of the 3 subsequent eligibility criteria are answered with YES and all exclusion criteria are answered with NO, THEN, is the patient eligible for recruitment





	PATIENT BASELINE CHARACTERISTICS					
1	Study Subject ID:	_ _ - - - Enter Study Subject ID in this format xxx-xxx-xxx 3 digit code for the country, 3 digit code for the hospital and 3 digit individual patient number				
		Did patient receive TTE within 6 months of surgery?				
2	Exposed / Non-Exposed	(or within 8 months if index surgery has been postponed)				
_		Yes No				
3	Written informed consent was obtained from patient	Date of informed consent (before or equal to date of surgery) _ - - - - - - - - - - in this format dd-Mmm-YYYY (Month in English starting with capital letter)				
4	Age (on day of surgery):	_ * under 18 is exclusion criteria				
5	Biological sex:	☐ Male ☐ Female				
		in case of gender-affirming hormone therapy use the therapy-induced sex				
6	Weight:	_ kg [40-e]				
7	Height:	_ cm [140-210]				
8	Preoperative functional status	☐ fully independent ☐ partially dependent ☐ fully dependent				
9	ASA Physical Status:					
10	Active cancer	☐ Yes ☐ No undergoing surgery for cancer OR known metastatic disease OR patient				
		has received active treatment for their cancer (e.g., chemotherapy, radiation or surgery) within the last 6 months				
11	Camily history of					
11	Family history of cardiovascular disease	☐ Yes ☐ No yes, if reported in any preoperative clinical documentation or if history provided by patient				
12	History of dyslipidemia	☐ Yes ☐ No yes, if diagnosis reported in any preoperative clinical documentation or if history of correspondent treatment				
13	Hypertension	☐Yes ☐ No				
		yes, if diagnosis reported in any preoperative clinical documentation or if history of correspondent treatment				
14	Current smoker	☐Yes ☐ No				
		< 1 yr prior intervention, excluding pipes, cigars, chewing tobacco				
15	Diabetes mellitus with treatment	☐Yes ☐ No				
	treatment	yes, if diagnosis reported in any preoperative clinical documentation or if history of correspondent treatment				
		15.1 If yes, treated with Diet Classical Oral antidiabetics GLP-1 Receptor Agonist/ SGLT2 inhibitors Insulin				
16	Haemoglobin Preop	16.1 g/L[60-180]				
	(closest to operation date and	g/dL[6-18]				
	≤3 months)	mg/mL[60-180]				
		16.2 .				





17	Preoperative hs-Troponin	17.1 _ ng/L Please be aware of unit					
		47.2 Associates must be Transmin					
		17.2 Assay information Troponin 99th percentile of local assay _ _ _ _ _ ng /L Please be aware of unit					
		99th percentile of local assay ng /L Please be aware of unit					
		17.3 Check which applies (single choice)					
		high-sensitivity Troponin I					
		high-sensitivity Troponin T					
18	Preoperative B-type Natriuretic Peptide	18.1 _ ng/L Please be aware of unit					
		18.2 Assay information					
		99th percentile of local assay _ _ ng /L Please be aware of unit					
		18.3 Check which applies (single choice)					
		□ NTproBNP					
		BNP					
19	Creatinine Preop:	19.1 μmol/L [1.0-3000.0]					
	(closest to surgery date and ≤3	mg/dL [0.1-30.0]					
	months)	19.2 _ . _					
20	Renal disease	☐Yes ☐ No					
		including chronic kidney disease, dialysis-dependent, hypertensive renal/heart disease with renal					
		failure, kidney-transplantation, acute renal failure					
		20.4 If was does notice to good dishuis Ves No					
		20.1 If yes ,does patient need dialysis Yes No					
		1					





	CARDIOPULMONARY COMORBIDITIES					
21	Chronic heart failure or cardiomyopathy	 Yes ☐ No yes, if diagnosis reported in any preoperative clinical documentation or if history of correspondent treatment 21.1 If yes, history of congestive heart failure? Yes ☐ No 				
22	Intake of heart failure medication (>30 days before surgery)	22.1 ACE-Inhibitors;				
23	History of Coronary Artery Disease	Yes No yes, if diagnosis reported in any preoperative clinical documentation or if history of myocardial infarction, correspondent interventions or medical treatment		23.1 If yes, check all that apply: previous myocardial infarction previous coronary intervention previous coronary artery bypass		
24	History of peripheral vascular disease	Yes No yes, if diagnosis reported in any preoperative clinical documentation or if history of correspondent interventions (e.g. PTA, vascular surgery)				
25	History of Stroke or TIA	Yes No yes, if diagnosis reported in any preoperative clinical documentation or if history of correspondent interventions				
26	History of severe valvular stenosis	26.1 If yes, check all that apply : Yes No Mitral valve Aortic valve Tricuspid valve				
27	History of severe valvular regurgitation	Yes No	27.1 If yes, check all that annly:			





28	History of cardiac arrhythmia or heart blocks History of pulmonary embolism	Yes No 28.1 If yes, check all that apply: atrial fibrillation, atrial flutter Heart blocks History of cardiac arrest Yes No yes, if diagnosis reported in any preoperative clinical documentation or if history of correspondent				
30	Chronic obstructive pulmonary disease	treatment Yes No yes, if diagnosis reported in any preoperative clinical documentation or if history of correspondent treatment 30.1 If yes History of severe COPD? Yes No Functional disability, hospitalization in the past for treatment of COPD, O2 therapy, Gold III-IV				
31	History of restrictive pulmonary disease	Yes No	ical documentation or if history of correspondent			
32	History of OSAS		ical documentation or if history of correspondent			
		SYMPTOMS AND CLINICAL SIGN	s			
33	Presence of symptoms	☐ Yes ☐ No	33.1 If yes, check all that apply: dyspnoea chest discomfort orthopnea history of syncope history of arrhythmia			
34	Presence of clinical signs	☐ Yes ☐ No	34.1 If yes , check all that apply : murmur crackles jugular vein distension peripheral oedema ascites pleural effusion arrhythmia			
35	Was ECG recorded?	Yes No 35.1 If yes, was any of the following detected? Check all that apply Atrial fibrillation Other non-sinus rhythms RBBB LBBB ischemic changes (ST depression, T inversion, ST elevation) non-acute/nonspecific ischemic changes(e.g. pathological Q waves, poor R wave progression, T-flattening) none of the above 35.2 If yes and patient had TTE, when was ECG recorded? Before TTE After TTE Not Applicable, Patient did not receive TTE				



	SELF-RE	PORTED FUN	CTIONAL (CAPACITY											
36	having t	o stop to rest	-	inuously climb without		<1	1	2	3	4	. 🗀 :	>4			
37	Single choice Choose one activity category that best describes your usual pattern of daily physical activities, (including activities related to house and family care, transportation, occupation, exercise and wellness and leisure or recreational purposes) Single choice					 □ Inactive or little activity other than usual daily activities □ Regularly (≥ d/wk) participate in physical activities requiring low levels of exertion that result in slight increases in breathing and heart rate for at least 10 minutes at a time □ Brisk walking, jogging or running, cycling, swimming, or vigorous sports at a comfortable pace or other activities requiring similar levels of exertion for 20 to 60 minutes per week □ Brisk walking, jogging or running, cycling, swimming, or vigorous sports at a comfortable pace or other activities requiring similar levels of exertion for 1 to 3 hours per week □ Brisk walking, jogging or running, cycling, swimming, or vigorous sports at a comfortable pace or other activities requiring similar levels of exertion for over 3 hours per week. 									
38	CLINICA	L FRAILTY SC	ALE		□ 1	2	<u></u> 3 [4	<u></u>		5 🗆 :	7 🔲	8 [] 9	
		CLIN 1 2 3 4 5 5	VERY FIT MANAGING WELL LIVING WITH VERY MILD FRAILTY LIVING WITH MILD FRAILTY	People who are robust, active, energetic and motivated. They tend to exercise regularly and are among the fittest for their age. People who have no active disease symptoms but are less fit than category 1. Often, they exercise or are very active occasionally, e.g., seasonally. People whose medical problems are well controlled, even if occasionally symptomatic, but often are not regularly active beyond routine walking. Previously "vulnerable," this category marks early transition from complete independence. While not dependent on others for daily help, often symptoms limit activities. A common complaint is being "slowed up" and/or being tired during the day. People who often have more evident slowing, and need help with high order instrumental activities of daily living (finances, transportation, heavy housework). Typically, mild frailty progressively impairs shopping and walking outside alone, meal preparation, medications and begins to restrict light housework.	The de corres; demen mild d the de still re repeat and so	gree of fra conds to to tia. Commementia in tails of a memberin mg the sa cial withd	LIVING WITH MODERATE FRAILTY LIVING WITH SEVERE FRAILTY LIVING WITH VERY SEVERE FRAILTY TERMINALLY ILL FRAILTY II INTERMINALLY ILL LIVING WITH VERY SEVERE FRAILTY ILL LIVING WITH VERY SEVERE FRAILTY ILL LIVING WITH VERY SEVERE FRAILTY ILL LIVING WITH SEVERE FRAILTY LIVING WITH SEVERE	activ Insidio	ities ance, e, they o as and need of the	I with ke Iten have Iten h	at for persuause (physey seems so dying (wi tor persuause) the persuause (physey seems so dying (wi tor persuause) the persuause to fife. This ople can be to death the seems to die persuause the persuause to die persuause to di	sse, s with ng and (culing, sonal sical or table thin -6 onal care cically, n a life e not allty, still h.) TIA mory is emingly s well, mpting, o often sood, eremission: re of fitness.			



BASELINE WHODAS 2.0							
	Difficulties?	None	Mild	Moderate	Severe	Extreme or Cannot do	
In the past 30 days, how much difficulty did you have in	39 Standing for long periods such as 30 minutes	□ 0	1	□ 2	□з	<u>4</u>	
nave in	40 Taking care of your household responsibilities	□ 0	1	□ 2	<u></u> 3	<u>4</u>	
	41 Learning a new task, for example, learning how to get to a new place	□ 0	1	2	□ 3	<u></u> 4	
	42 How much of a problem did you have joining in community activities (for example festivities, religious or other activities) in the same way as everyone else can?	□ 0	□ 1	☐ 2	□ 3	□ 4	
	43 How much have you been emotionally affected by your health problems?	□ 0	1	☐ 2	□ 3	_ 4	
	44 Concentrating on doing something for ten minutes	□ 0	1	2	З	<u></u> 4	
	45 Walking a long distance such as a kilometre	□ 0	□ 1	□ 2	□ 3	<u>4</u>	
	46 Washing your whole body	□ 0	□ 1	□ 2	□ 3	<u>4</u>	
	47 Getting dressed	□ 0	□ 1	2	<u></u> 3	□ 4	
	48 Dealing with people you do not know	□ 0	□ 1	□ 2	□ 3	□ 4	
	49 Maintaining a friendship	□ 0	□ 1	2	<u></u> 3	□ 4	
	50 Your day-to-day work/school	□ 0	□ 1	□ 2	□ 3	□ 4	



	PREOPERATIVE TRANSTHORACIC ECHOCARDIOGRAPHY				
51	Was preoperative transthoracic	☐ Yes ☐ No			
	echocardiography conducted within 6 months of the surgery	55.1 If yes Date: _ - _ - _ _			
52	The TTE was requested	specifically due to the surgery independent of the surgery			
53	Why was TTE requested?	new symptoms/ signs routine follow-up (prior disease)			
54	Type of TTE	 extensive TTE examination (cardiological TTE lab) focused TTE 54.1 If focused TTE, when was TTE conducted? at the time of preoperative assessment 			
		immediately preoperatively			
55	How was the estimated Ejection Fraction (EF)?	☐ EF in percentage ☐ EF qualitative 55.1 If EF reported in percentage, estimated EF % ☐ ☐ ☐ ☐ / %			
		55.2 If qualitative report only, check what applies (single choice): Normal EF mildly reduced EF moderately reduced EF severely reduced EF			
56	Regional wall motion abnormality	Yes No 56.1 If yes, is regional wall motion abnormality new/not previously described?			
57	Significant (Grade II or more) LV diastolic dysfunction with evidence of increased LV filling pressures	☐ Yes ☐ No ☐ Yes ☐ No			
58	Significant RV dysfunction	☐ Yes ☐ No			
59	Clinically relevant mitral valve stenosis	☐ Yes ☐ No			
60	Severe mitral regurgitation	☐ Yes ☐ No			
61	Severe aortic stenosis	☐ Yes ☐ No			
62	Severe aortic regurgitation	☐ Yes ☐ No			



	PERIOPERATIVE MANAGEMENT					
63	Was surgery postponed?	 Yes				
64	Was Surgery Cancelled?	Yes No If yes: 64.1 □ cancellation (based on doctor/board decision) Date _ _ - _ _ _ _ (dd/Mmm/YYYY) NOTE: even if the patient was cancelled complete CRF including follow-up 30 days after cancellation 64.2 □ cancellation after shared decision with patient Date _ _ - _ _ _ _ (dd/Mmm/YYYY) NOTE: even if the patient was cancelled complete CRF including follow-up 30 days after cancellation				
65	Discussion in preoperative multidisciplinary board	G5.1 If yes: decision derived from board discussion (check all that apply)				
66	Cardiological work-up	Yes No 66.1 If yes check all that apply □ cardiac MRI 66.1.1 If yes and patient had TTE, when was cardiac MRI conducted? □ Before TTE After TTE Not Applicable, Patient did not receive TTE □ cardiac CTscan 66.1.2 If yes and patient had TTE, when was cardiac CTscan conducted? □ Before TTE After TTE Not Applicable, Patient did not receive TTE				





			stress imaging				
			66.1.3 If yes and patient had TTE, when was stress imaging conducted?				
			☐ Before TTE ☐ After TTE ☐ Not Applicable, Patient did not receive TTE				
			coronary angiography				
			66.1.4 If yes and patient had TTE, when was coronary angiography conducted?				
			☐ Before TTE ☐ After TTE ☐ Not Applicable, Patient did not receive TTE				
			☐ PCI or CABG				
			66.1.5 If yes and patient had TTE, when was PCI or CABG conducted ?				
			☐ Before TTE ☐ After TTE ☐ Not Applicable, Patient did not receive TTE				
			valvuloplasty or TAVI				
			66.1 6 If yes and patient had TTE, when was valvuloplasty or TAVI conducted?				
			☐ Before TTE ☐ After TTE ☐ Not Applicable, Patient did not receive TTE				
67	Changes in	Yes	No				
	medication						
		67.1 If yes, exis	sting drug or introduction of new drug				
			tation of dosage of existing drug				
		intro	duction of new drug class				
			1.1 If new drug, check all that apply				
			Aspirin				
			ADP-inhibitors e.g.clopidogrel, prasugrel, ticagrelor, ticlodipine, cangrelor				
			☐ Vit K-antagonits ☐ LMWH				
			NOACs e.g. rivoroxaban, apixaban, dabigatran, edoxaban				
			B-Blockers				
			Inhibitors of renin angiotensin system (ACE-inhibitors, ADII Antagonists, Renin inhibitors,				
			Angiotensin receptor neprilysin inhibitors) Calcium channel-blockers				
			Statins				
			Diuretics				
			Mineralocorticoid/aldosterone receptor antagonists, e.g. spironolactone, eplerenone,				
			finorenone PDE-Inhibitors (e.g. Avanafil, Sildenafil, Tadalafil, Vardenafil) or Endothelin-Inhibitors (e.g.				
			Bosentan, Macitentan, Darusentan, Ambrisentan)				
			SGLT-2 inhibitors (e.g. but not limited to : canagliflozin, dapagliflozin, empagliflozin				
			ertugliflozin)				
			GLP-1 receptor agonists (e.g. but not limited to liraglutide, semaglutide, tirzepatide, exenatide, lixisenatide, albiglutide, dulaglutide, exenatide				
68	Planned ICU/IMC a	dmission or	☐ Yes ☐ No				
	planned extended I						
			68.1 If yes: Is ICU/IMC admission/ extended PACU stay standard of care for the conducted				
			surgery?				
			(always done in your institution for this specific surgery)				
			Yes No				
69	Systematic periope	rative	Yes No				
	Troponin screening						
			69.1 If yes: Is systematic perioperative Troponin screening standard of care for the conducted				
			surgery?				
1			(always done in your institution for this specific surgery)				
l			☐ Yes ☐ No				





	INTRAOPERATIVE DATA							
70	Date of surgery	Date _ - _ _	_ - _ _	_l (dd/Mmm/YYYY)				
71	Type of surgery	☐ Moderate risk ☐	☐ Moderate risk ☐ High risk					
72	Intrathoracic, intra-abdominal, or suprainguinal vascular surgery?	Yes No						
73	Surgery site (as per NSQIP MICA)	Anorectal Aortic Bariatric Brain Breast ENT (not thyroid/pa Foregut/Hepatopan Gallbladder, append spleen Hernia (ventral, inge	ncreatobiliary dix, adrenal and	Neck (thyroid/parathyroid) Obstetric/Gynaecologic Orthopaedic and non-vascular extremity Other abdominal Peripheral vascular Skin Spine Non-oesophageal thoracic Vein Urology				
74	Anaesthesia technique	General General combined with regional neuraxial peripheral Regional-peripheral Regional-neuraxial Regional-combined peripheral and neuraxial						
75	Invasive blood pressure measurement	Yes No 75.1 If yes: Is invasive blood pressure measurement local standar care for the conducted surgery? (always done in your institution for this specific surgery) Yes No						
76	Central venous line	Yes No	conducted surge	our institution for this specific surgery)				
77	Pulmonary arterial catheter	Yes No	77.1 If yes: Is pulmonary arterial catheter local standard of care for the conducted surgery? (always done in your institution for this specific surgery) Yes No					
78	Intraoperative TransoEsophageal Echocardiogram - TEE	Yes No	78.1 If yes: Is intraoperative TEE local standard of care for the conducted surgery? (always done in your institution for this specific surgery) Yes No					
79	Pulse contour cardiac output (PiCCO) or devices for cardiac output estimation	Yes No	79.1 If yes: are pulse contour cardiac output or devices for cardiac output estimation local standard care for the conducted procedure? (always done in your institution for this specific surgery) Yes No					
80	Goal-directed haemodynamic management as per locally implemented protocol	Yes No	care for the con	our institution for this specific surgery)				





INHOSPITAL I	INHOSPITAL POSTOPERATIVE EVOLUTION						
81 ICU Admission?	☐ Yes ☐ No						
If Yes, First ICU admission 81.1	☐ Planned, i.e. decision of ICU admission prior to anesthesia induction ☐ Unplanned, i.e. decision of ICU admission during/after anaesthesia induction or during hospital stay						
81.2 Length of first stay in ICU (days)	_ _ _						
81.3 Was patient readmitted to ICU (second stay or more in ICU) ?	☐ Yes ☐ No	81.3.1 If yes: Total Length of <u>additional</u> stay in ICU (days) _ if several readmissions please report the cumulative length of stay in ICU					
82 Grade of most severe complication in hospital according to Clavien-Dindo class (single choice)							
None I III IIIa IIIb IVa IVb	□v						
GRADE I: Any deviation from the normal postoperative course without the need for pharmacological treatment or surgical, endoscopic and radiological interventions (Allowed therapeutic regimens are: drugs as antiemetics, antipyretics, analgetics, diuretics and electrolytes and physiotherapy. This grade also includes wound infections opened at the bedside) GRADE II: Requiring pharmacological treatment with drugs other than such allowed for grade I complications. Blood transfusions and total parenteral nutrition are also included. GRADE III: Requiring surgical, endoscopic or radiological intervention. - IIIa: Intervention not under general anaesthesia - IIIb: Intervention under general anaesthesia GRADE IV: Life-threatening complication (including CNS complications) * requiring IC/ICU-management - IVa: single organ dysfunction (including dialysis) - IVb: multiorgan dysfunction GRADE V: Death of a patient							
82.1 If any, organ system of most severe compl							
Cardiovascular Respiratory Neurological Gastrointestinal Renal Surgical site infection Other Not Applicable / no complication for this patient	mignest Clavien-Dindo	class, if several with the same class check all that apply					
83 Date of Hospital Discharge	Patient has been d 83.1 If discharged , da - _ [>=Date of surgery]	oital at 30-day follow up lischarged (or died in hospital) te of discharge: - (dd/Mmm/YYYY) of hospital discharge = date of death					





30-DAY OUTCOMES									
84	When was follow-up performed?	_ - _ - (dd/Mmm/YYYY) [>=30 days Date of surgery]							
85	Did the patient die?	Yes No 85.1 If yes, did patient die from cardiac death? Yes No (Please refer to appendix 2 for cardiac death definition) 85.2 If patient died, please enter date of death _ _ - _ _ _ _ _ (dd/Mmm/YYYY) 85.3 If exact date unknown, confirm that death occurred within 30 days of surgery Confirmed Not confirmed							
86	Did the patient suffer a Myocardial Infarction?	86.1 If yes , please enter date of MI _ _ - _ _ - _ _ (dd/Mmm/YYYY) 86.2 If exact date unknown, confirm that MI occurred within 30 days of surgery Confirmed Not confirmed							
87	Did the patient suffer a non-fatal cardiac arrest?	Yes No 87.1 If yes , please enter date of non-fatal cardiac arrest _ - _ - _ (dd/Mmm/YYYY) 87.2 If exact date unknown, confirm that non-fatal cardiac arrest occurred within 30 days of surgery Confirmed Not confirmed							
88	Did the patient suffer an acute heart failure or decompensation of chronic heart failure?	Yes No 88.1 If yes , date of acute heart failure or decompensation of chronic heart failure _ - _ - _ _ (dd/Mmm/YYYY) 88.2 If exact date unknown, confirm that acute heart failure or decompensation of chronic heart failure occurred within 30 days of surgery Confirmed Not confirmed							
89	Was the patient submitted to coronary revascularisation	Yes No 89.1 If yes , date of coronary revascularisation _ - _ - _ _ (dd/Mmm/YYYY) 89.2 If exact date unknown, confirm that coronary revascularisation occurred within 30 days of surgery Confirmed Not confirmed							
90	Did patient suffer any recurring event? (check all that apply)	 No Myocardial Infarction Non-fatal cardiac arrest Acute heart failure or decompensation of chronic heart failure Coronary revascularisation 							





30-DAY WHODAS 2.0										
91	Was 30-Day WHODAS Completed		es No,patient died							
			Difficulties?	None	Mild	Moderate	Severe	Extreme or Cannot do		
	In the past 30 days, how much difficulty did you have in	92	Standing for long periods such as 30 minutes	□ 0	□ 1	□ 2	<u></u> 3	□4		
		93	Taking care of your household responsibilities	□ 0	<u> </u>	□ 2	□ 3	□ 4		
		94	Learning a new task, for examplelearning how to get to a new place	□ 0	□ 1	□ 2	□3	□4		
		95	How much of a problem did you have joining in community activities (for example festivities, religious or other activities) in the same way as everyone else can?	□ 0	□ 1	□ 2	<u> </u>	□ 4		
		96	How much have you been emotionally affected by your health problems?	□ 0	1	□ 2	<u></u> 3	□4		
		97	Concentrating on doing something for ten minutes	□ 0	□ 1	□ 2	□ 3	□4		
		98	Walking a long distance such as a kilometre	□ 0	□ 1	□ 2	□ 3	□ 4		
		99	Washing your whole body	□ 0	□ 1	□ 2	□ 3	4		
		100	Getting dressed	□ 0	□ 1	□ 2	□ 3	<u></u> 4		
		101	Dealing with people you do not know	□ 0	□ 1	□ 2	□ 3	□4		
		102	Maintaining a friendship	□ 0	□ 1	□ 2	□ 3	<u></u> 4		
		103	Your day-to-day work/school	□ 0	□ 1	□ 2	□ 3	□4		
DAYS ALIVE AND OUT OF HOSPITAL AT 30 DAYS										
Was the patient admitted to an acute medical care facility after discharge from the initial institution? (Please note also early inter-institutional transfer should be reported as admission) Yes No 104.1 If yes, length of stay in health care facility (days) _ 104.2 If yes, was readmission for cardiovascular reason? Yes No 104.3 Was the patient readmitted more than once? (Please note also early inter-institutional transfer should be as admission) Yes No 104.3.1 If yes: how many additional times? 104.3.2 If yes: cumulative length of stay in health care facility(ies) (days) _								ıld be reported		

104.3.3 If yes, was any readmission for cardiovascular reason? Yes No