	failure (HF) in primary and secondary care
	,
Study ID (surname of first author ar	nd year first full report of study was published e.g. Smith 2001)
Report IDs of other reports of th	is study (e.g. duplicate publications, follow-up studies)
1. General Information	
1. General mornida	
Date form completed	
(dd/mm/yyyy)	
Name of person extracting data	
Report title	
(title of paper/ abstract/ report	
that data are extracted from)	
Report ID	
(ID for this paper/ abstract/ report)	
Reference details	
Report author contact details	
Publication type	
(e.a. full report, abstract, letter)	

Review title or ID

Study funding sources (including role of funders) Possible conflicts of interest

(for study authors)

Notes:

## 2. Study Eligibility

Study	Eligibility criteria				Location in text
Characteristics	(Insert eligibility criteria for each characteristic as				(pg & ¶/fig/table)
	defined in the Protocol)	Yes	No	Unclear	
Type of study	Randomised Controlled Trial				
	Controlled Clinical Trial (quasi-randomised trial)				
Participants	Patients >18 years who are being treated for HF in primary or secondary care.				
Intervention	Treatment guided by serial BNP measurements (BNP-guided therapy).  Treatment guided by clinical assessment (standard care).				
Any outcome measures reported	All-cause mortality Death related to HF Cardiovascular death All-cause hospital admission Hospital admission for HF Adverse events Quality of life				
	INCLUDE EXCLUDE				
If excluded, provide reason					
Notes:					

DO NOT PROCEED IF STUDY EXCLUDED FROM REVIEW

# 3. Participants and Setting

	Description	Location in text
Number of participants		(pg & ¶/fig/table)
randomised, incl. no. in each		
group		
Population description		
(from which study		
participants are drawn)		
Setting		
(e.g. Primary or secondary		
care, in-hospital etc)		
Inclusion criteria		
Exclusion criteria		
Exclusion criteria		
Method/s of recruitment of		
participants		
Informed consent obtained		
	Yes No Unclear	
Any baseline imbalances	les No Officieal	
identified?		
Withdrawals and exclusions		
(if not provided below by		
outcome)		
Age		
Sex		
Race/Ethnicity		
Severity of illness		
(e.g. LV function, NYHA class)		
Co-morbidities		
Other treatment received		
(additional to study		
intervention)		
Other relevant		
sociodemographics		
Subgroups measured		
Subgroups reported		
Notes:		

### 4. Methods

	Descriptions as stated in report/paper	Location in text (pg & ¶/fig/table)
Aim of study		
<b>Design</b> (e.g. parallel, crossover, cluster)		
Unit of allocation (by individuals, cluster/ groups or body parts)		
Start date		
End date		
Total study duration		
Ethical approval needed/ obtained for study	Yes No Unclear	
Notes:		

### 5. Intervention Details

**Intervention Group (BNP guided therapy)** 

	Description as stated in report/paper	Location in text (pg & ¶/fig/table)
BNP/NTproBNP target		(pg & <sub>II/J</sub> ig/table)
Description of intervention (include sufficient detail for replication, e.g. content, dose, components)		
Treatment algorithm used		
Duration of treatment period		
<b>Timing</b> (e.g. frequency, duration)		
<b>Delivery of intervention</b> (how was the intervention delivered)		
Intervention providers (who delivered the intervention)		
Co-interventions		
Notes:		

## **Control Group (Symptom-Guided Therapy)**

	Description as stated in	report/paper	Location in text
			(pg & ¶/fig/table)
Clinical target			
Description of intervention			
(include sufficient detail for			
replication, e.g. content,			
dose, components)			
Treatment algorithm used			
Duration of treatment period			
Timing (e.g. frequency,			
duration)			
Delivery of intervention (how			
was the intervention			
delivered)			
Intervention providers			
(who delivered the			
intervention)			
Co-interventions			
Notes:			
Notes.			
6. Outcomes Reported	l		
Outcome reported	Yes No Unclear	Comments	Location in text
All-cause mortality			(pg & ¶/fig/table)
Death related to HF			
Cardiovascular death			
All-cause hospital admission			
Hospital admission for HF			
Adverse events			
Quality of life			

### 7. Risk of Bias Assessment

Domain	Judgement			Description	Location in text
	Low risk	High risk	Unclear		(pg & ¶/fig/table)
Random sequence generation (selection bias)					
Allocation concealment (selection bias)					
Blinding of participants and personnel (performance bias)					
All-cause mortality					
Cause specific mortality (death from HF / CVD)					
Adverse events/hospital admission					
QoL					
Blinding of outcome assessment (detection bias)					
All-cause mortality					
Cause specific mortality (death from HF / CVD)					
Adverse events/hospital admission					
QoL					
Incomplete outcome data (attrition bias)					
All-cause mortality					
Cause specific mortality (death from HF / CVD)					
Adverse events/hospital admission					
QoL					
Selective outcome reporting? (reporting bias)					
Other bias					

#### 8. Outcome data

Trial name	Number of patients	Point estimates (Hazard ratio or other information reported)	Lower confidence interval	Upper confidence interval	Number of patients in BNP group	Number of patients in control group	Number of events in BNP group	Number of events in control group	Comments