A more granular view on pulmonary embolism
Mos, I.C.M.

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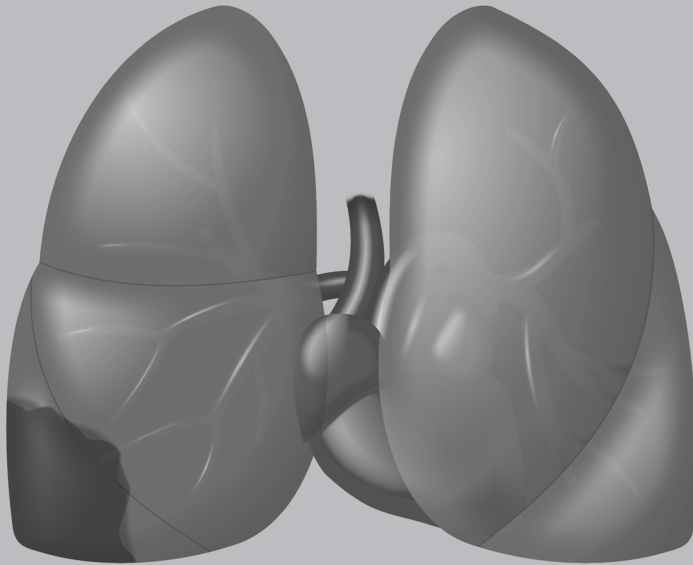
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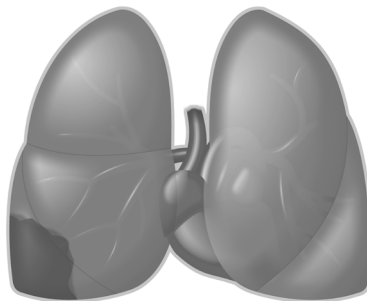
Brain-type natriuretic peptide levels in the prediction of adverse outcome in patients with pulmonary embolism: a systematic review and meta-analysis.

I.C.M. Mos*, F.A. Klok* and M.V. Huisman

*Equally contributed

ABSTRACT

Rationale

The potential role of elevated brain-type natriuretic peptides in the differentiation of patients suffering from acute pulmonary embolism at risk for adverse clinical outcome has not been fully established.

Objectives

We evaluated the relation between elevated (NT-pro-)BNP levels and clinical outcome in patients with pulmonary embolism.

Methods

Articles reporting on studies that evaluated the risk of adverse outcome in patients with pulmonary embolism and elevated (NT-pro-)BNP levels were abstracted from Medline and EMBASE. Information on study design, patient- and assay characteristics and clinical outcome were extracted. Primary endpoints were overall mortality and pre-defined composite outcome of adverse clinical events.

Measurements and Main Results

Data from 13 studies were included. In 51% (576/1132) of the patients (NT-pro-)BNP levels were increased. The different analyses were performed in subpopulations. Elevated levels of (NT-pro-)BNP were significantly associated with right ventricular dysfunction (p<0.001). Patients with high (NT-pro-)BNP concentration were at higher risk of complicated in-hospital course (OR 6.8, 95%CI 4.4-10) and 30-day mortality (OR 7.6, 95%CI 3.4-17).Patients with a high (NT-pro-)BNP had a 10% risk of dying (68/671, 95%CI 8.0-13) while 23% (209/909, 95%CI 20-26) had an adverse clinical outcome.

Conclusions

High concentrations of BNP distinguish patients with pulmonary embolism at higher risk of complicated in-hospital course and death from those with low BNP levels. Increased (NT-pro-)BNP concentrations alone however do not justify more invasive treatment regimens.

INTRODUCTION

Right ventricular dysfunction on echocardiography is a common clinical finding in patients with acute pulmonary embolism (PE)[1-3] and predicts poor outcome in these patients. Prognostic stratification in acute PE patients may have consequences on management decisions. Patients identified with a low risk of complicated outcome may be eligible for outpatient management and high risk patients may benefit of more aggressive treatment.[1-2]

Several cardiac biomarkers have emerged as indicator of right ventricular dysfunction and predictor of clinical outcome in patients with acute PE. A recent meta-analysis demonstrated that elevated troponin levels identify patients with PE at high risk of short term death and adverse outcome.[4] Also, Brain-type natriuretic peptide (BNP) is a marker of ventricular dysfunction. This hormone is released in response to myocyte stretch. It is synthesized as an inactive prohormone (pro-BNP) that is split into the active hormone BNP and the inactive N-terminal fragment (NT-pro-BNP).[5] Several prospective studies have been performed to identify to potential role of either BNP or NT-pro-BNP in the risk stratification of patients with PE.[6-18] However, reported studies have limited patient numbers, used different cutoff points and involved different clinical endpoints. Therefore, we performed a meta-analysis of studies in patients with acute PE to evaluate the relation between elevated levels of BNP or NT-pro-BNP and clinical outcome.

METHODS

Data sources

A literature search was performed to identify all published prospective studies on BNP or NT-pro-BNP levels and clinical outcome in patients with PE. Medline and EMBASE were searched using pre-defined search terms, between January 1980 and October 2007. Search criteria included "Pulmonary Embolism" and "pro-brain natriuretic peptide" or "Brain Natriuretic Peptide" or "natriuretic peptide". Also, by searching the reference lists of all established studies, the researchers aimed to identify additional relevant papers. Papers were not limited to the English language. Only full papers were applicable for this analysis.

Study outcome

Objectively adjudicated short term adverse clinical events were used as primary outcome of this meta-analysis. These included mortality or an adverse clinical outcome defined as the occurrence of any of the following: death, cardiopulmonary resuscitation, mechanical ventilation, use of vasopressors, thrombolysis, thrombosuction, open surgi-

cal embolectomy or admission to the ICU. Right ventricular dysfunction was used as secondary endpoint.

Study selection and data extraction

Two independent researchers (F.K. and I.M., both MD) performed study selection. In case of disagreements a third researcher (M.H., MD, PhD) was consulted. Criteria for selection were a prospective design, consecutive inclusion, pre-defined endpoints, clear description of in- and exclusion criteria, objective criteria for diagnosis of PE, standardized treatment and the possibility of creating a 2 by 2 table based on BNP or NT-pro-BNP levels and clinical endpoints. Study sample size was not an eligibility criterion. Objective criteria for PE were positive CT findings, high probability VQ scan, positive pulmonary angiography or clinical suspicion of PE in combination with an ultrasonography proven deep vein thrombosis. Le Gal et al recently described that a positive compression ultrasonography of the lower limb veins is highly predictive of PE on computed tomography in suspected patients.[19] Data regarding patient characteristics, exclusion criteria, diagnostic criteria for PE, severity of PE (inclusion of hemodynamic instable patients and use of thrombolytic therapy), completeness of follow-up, immunoassay, timing of sampling, cutoff level, follow-up period and endpoints were abstracted.

Statistical analysis

Data were entered in Review Manager (Version 4.2 for Windows. Copenhagen: The Nordic Cochrane Centre, the Cochrane Collaboration, 2003). Individual and pooled odds ratios were calculated to assess the relation between elevated BNP or NT-pro-BNP levels and clinical outcome. Mantel-Haenszel Methods for Combining Trials were used for weighting the studies. Cochran's chi-square test and the I^2 test for heterogeneity were used to assess inter study heterogeneity. The chi-square test assesses whether observed differences in results are compatible with chance alone. I^2 describes the percentage of the variability in effect estimates that is due to heterogeneity rather than sampling error. Statistically significant heterogeneity was considered present at chi-square $p<0.10$ and $I^2 >50\%$.

RESULTS

Study selection

As a result of the literature search, 124 studies were revealed. Articles were excluded by review of title and abstract in case of review articles (n=48), animal studies (n=2), case reports (n=5), editorials, letters or author replies (n=13), studies not including the clinical course of PE (n=6) and if it concerned studies on other diseases than PE (n=17, Figure 1).
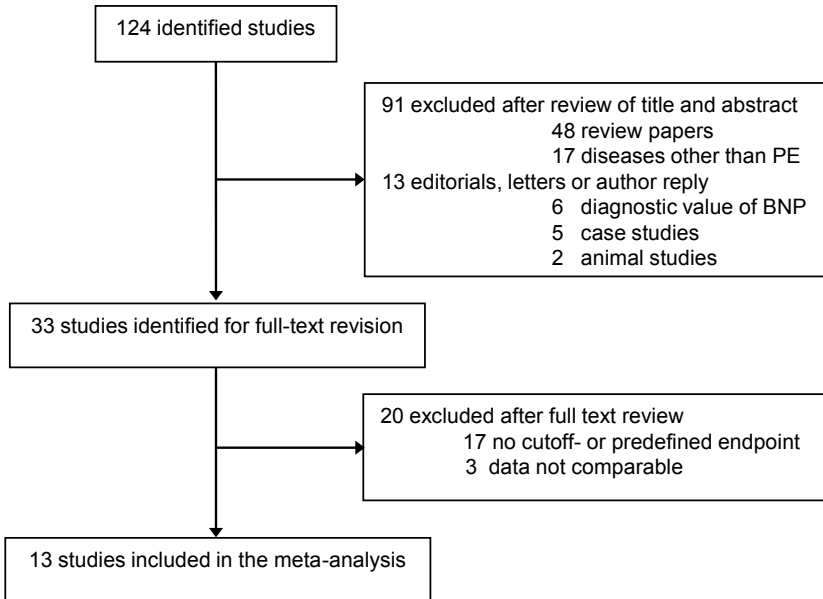
Figure 1. Flow diagram of study selection. PE: pulmonary embolism.

After full review, 20 more studies were excluded because our predefined endpoints were not reported (17) or no cutoff points were mentioned (3). We identified 13 studies that met our criteria.[6-18]

Characteristics of included studies

Demographic characteristics of the patients were comparable between all included studies (Table 1). Mean age of the patients varied between 53 and 75 years, the proportion of females ranged from 36-74%. In most patients, the diagnosis of PE was confirmed by CT-scan, high probability V/Q scan or pulmonary angiography. In three studies, hemodynamically unstable patients were excluded.[7,11,17] Noticeably, in two of these latter studies, some patients received thrombolytic therapy during their hospital stay.[7,11] Two included studies reported on partially overlapping patient cohorts.[16,18] Because one of them used BNP[16] and the other NT-pro-BNP[18] levels as outcome parameter, both studies could be incorporated into subgroup analyses based on type of BNP testing.

Assays and cutoff points

As shown in Table 1, all studies reporting NT-pro-BNP levels used a Roche analyzer (2 types), with 3 different cutoff levels, varying from 500 till 1000 pg/ml. In the BNP studies, two assays with 4 different cutoff levels varying between 75 and 100 pg/ml were used. In all included studies, the timing of sampling is comparable. Cutoff levels were not predefined in most studies. In these 10 papers, receiver operating characteristics

Table 1. Characteristics of included studies.

Marker	Ref	n	Female (%)	Age*	Assay†	Timing of sampling	Cut-off	Follow-up	PE diagnosis	Hemodynamic instability¥	Thrombolysis (n, %)
NT-pro-BNP	6	60	60	72 ±15	Roche, Elecsys 2010 analyzer	Admission	1000pg/ml§	In hospital	PA, V/Q, ultrasonography[a]	Yes	1 (1.7)
	8	107	63	61 ±6	Roche, Elecsys 2010 analyzer	Admission, 4h, 8h, 24h	1000pg/ml	30 days	PA, V/Q, ultrasonography[a]	Yes	–[¤]
	12	124	60	60 ±18	Roche, Elecsys 2010 analyzer	Admission, 4h, 8h, 24h	1000pg/ml	In hospital	PA, V/Q, ultrasonography[a]	Yes	12 (11)
	13	100	65	63 ±18	Roche, ECLIA	Admission	600pg/ml	40 days	PA, V/Q	Yes	7 (7.0)
	15	79	63	63 ±17	Roche, Elecsys 2010 analyzer	Admission	600pg/ml	In hospital	PA, V/Q	Yes	8 (10)
	18	73	41	61 ±18	Roche, Elecsys 2010 analyzer	Admission	500pg/ml	In hospital	PA, V/Q	Yes	10 (14)
BNP	7	67	41	64 ±17	Biosite Diagnostics, Triage	Admission	100 pg/ml	NA‡	CT, V/Q	No	6 (9.0)
	9	181	58	53 ±17	Biosite Diagnostics, Triage	Admission	90pg/ml§	6 months	PA, V/Q	Yes	13 (22)
	10	51	65	79 ±9	Biosite Diagnostics, Triage	Admission	100 pg/ml	In hospital	PA, V/Q, ultrasonography[a]	Yes	0 (0)
	11	61	74	75 ±14	Biosite Diagnostics, Triage	Admission	89 pg/ml	In hospital	PA, pulmonary angiography	No	7 (11)
	14	46	36	57 ±19	Biosite Diagnostics, Triage	Admission	90 pg/ml	In hospital	PA, V/Q, echocardiography[Δ]	Yes	22 (48)
	16	73	41	61 ±18	Biosite Diagnostics, Triage	Within 4 hours	90pg/ml§	In hospital	PA, V/Q, embolectomy	Yes	6 (8.2)
	17	110	–¶	58 ±18	Immuno radiometric assay, Shionoria	Admission	75 pg/ml	3 months	PA, V/Q, ultrasonography	No	0 (0)

*Mean ±SD; †manufacturer and kind of assay (all were quantitative assays); ‡not applicable: endpoint was right ventricular dysfunction at time of diagnosis; §information was not provided; §predefined cut-off point; [a]typical clinical presentation and positive ultrasonography of lower limbs; [Δ]typical presentation and suggestive echocardiography; [¤]specific information was not provided; ¥patients with hemodynamic instability were eligible for the study. PA: computed tomography or conventional pulmonary angiography; VQ: ventilation perfusion scintigraphy; n: number.

(ROC) analyses were performed to retrospectively determine optimal cutoff values with regard to complicated PE. Normal levels are defined as levels beneath or equal to the cutoff point.

Clinical outcome

Overall, in 51% (576/1132) of the patients the assays showed elevated plasma concentrations of (NT-pro-) BNP. Data on overall mortality were reported in 4 studies using BNP[10,11,14,17] and 4 studies using NT-pro-BNP.[8,12,13,15] In the BNP cohort, 17 of 123 patients (14%, 95%CI 8.3-21) with elevated BNP levels died compared to 3 of 138 (2.2%, 95%CI 0.45-6.2) with normal BNP levels. This resulted in an overall OR for death of 6.5 (95%CI 2.0-21, Figure 2). One study had a follow up of 3 months[17], as compared to the other 3 which had in-hospital follow-up. If this single study was left out of the analysis, overall OR decreased to 3.3 (95%CI 0.6-18). In the NT-pro-BNP cohort, 46 of 250 patients (18%, 95%CI 14-24) with elevated NT-pro-BNP levels died in comparison with 2 of 160 (1.3%, 95%CI 0.15-4.4) with normal NT-pro-BNP levels, OR for death was 8.7 (95%CI 2.8-27, Figure 2).

Numbers on PE-related mortality were only available in 3 studies.[11,13,17] Because follow-up time was dissimilar between these studies and not all mortality cases were adjudicated by an independent, blinded committee to determine the cause of death, we could not use PE-related mortality as an outcome of this analysis.

Ten studies provided data on adverse clinical outcome[6,8-13,15,16,18] of which 6 had NT-pro-BNP levels as outcome parameter.[6,8,12,13,15,18] Overall, criteria for adverse clinical outcome were comparable throughout all studies. In the BNP study group, 47 of 128 (37%, 95%CI 28-46) patients with elevated BNP levels had adverse advents during follow-up in comparison with 28 of 208 (13%, 95%CI 9.1-19) patients with normal plasma concentrations. High BNP levels were associated with a higher risk of occurrence of adverse clinical events (OR 6.3, 95%CI 3.6-11, Figure 3). This OR was even higher (9.5, 95%CI 3.5-25) after exclusion of 1 study with 6 months of follow-up,[9] thereby limiting the outcome to in hospital clinical course. Of the 318 patients with elevated NT-pro-BNP levels, 102 experienced short term adverse events (32%, 95%CI 27-38) as compared to 12 of 225 (5.3%, 95%CI 2.8-9.1) patients with normal NT-pro-BNP levels. Patients with high NT-pro-BNP serum concentration were at higher risk of complicated in-hospital course compared to patients with normal levels (OR 7.5, 95%CI 3.8-15, Figure 3). Pooled data of all assays showed elevated (NT-pro-) BNP levels in 52% of the patients with a risk of 23% (209/909, 95%CI 20-26) and an OR of 6.8 (95%CI 4.4-10) towards complicated clinical course.

Right ventricular dysfunction

Data on right ventricular dysfunction were reported in 6 studies (Figure 4). Four studies were evaluating BNP (243 patients)[7,11,14,16] and 2 studies evaluated NT-pro-BNP levels

Overall mortality

Study	High BNP n/N	Normal BNP n/N	OR (random) 95% CI	Weight %	OR (random) 95% CI
Ray et al[10]	3/29	0/19		14.82	5.15 (0.25-106)
Pieralli et al[11]	4/41	0/20		15.31	4.92 (0.25-96.0)
Krüger et al[14]	1/17	1/25		16.72	1.50 (0.09-25.8)
Ten Wolde et al[17]	9/36	2/74		53.15	12.0 (2.44-59.1)
Total	**17/123**	**3/138**		**100**	**6.52 (2.04-20.9)**

Test for heterogeneity: Chi²=1.65, df=3 (p=0.65), I²=0%
Test for overall effect: Z=3.16 (p=0.002)

	High NT-pro-BNP n/N	Normal NT-pro-BNP n/N			
Puls et al[8]	9/53	2/54		52.20	5.32 (1.09-25.9)
Binder et al[12]	7/67	0/57		15.73	14.3 (0.80-255)
Kostrubiec et al[13]	15/72	0/28		16.10	15.4 (0.89-266)
Pruszczyk et al[15]	15/58	0/21		15.97	15.3 (0.87-268)
Total	**46/250**	**2/60**		**100**	**8.72 (2.78-27.4)**

Test for heterogeneity: Chi²=0.85, df=3 (p=0.84), I²=0%
Test for overall effect: Z=3.71 (p=0.0002)

Combined total	**63/373**	**5/298**		**100**	**7.56 (3.35-17.1)**

Test for heterogeneity: Chi²=2.62, df=7 (p=0.92), I²=0%
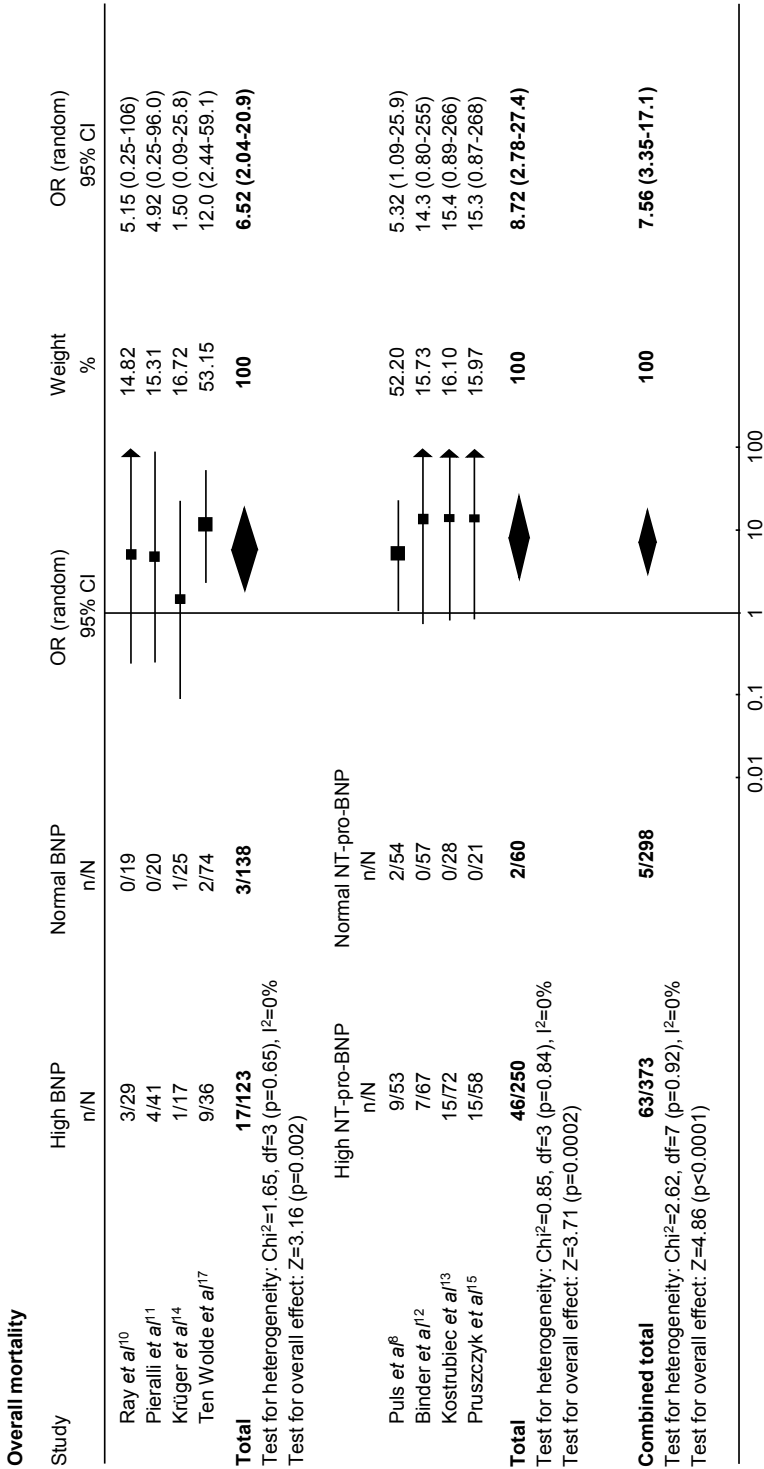Test for overall effect: Z=4.86 (p<0.0001)

Figure 2. Odds ratio (OR) for overall mortality based on elevated (NT-pro-) BNP levels. Different cutoffs were used for different studies; Mantel-Haenszel methods for combining trials were used for weighting the studies. CI: confidence interval.

Figure 3. Odds ratio (OR) for adverse clinical outcome based on elevated (NT-pro-) BNP levels. Different cutoffs were used for different studies; Mantel-Haenszel methods for combining trials were used for weighting the studies. CI: confidence interval.
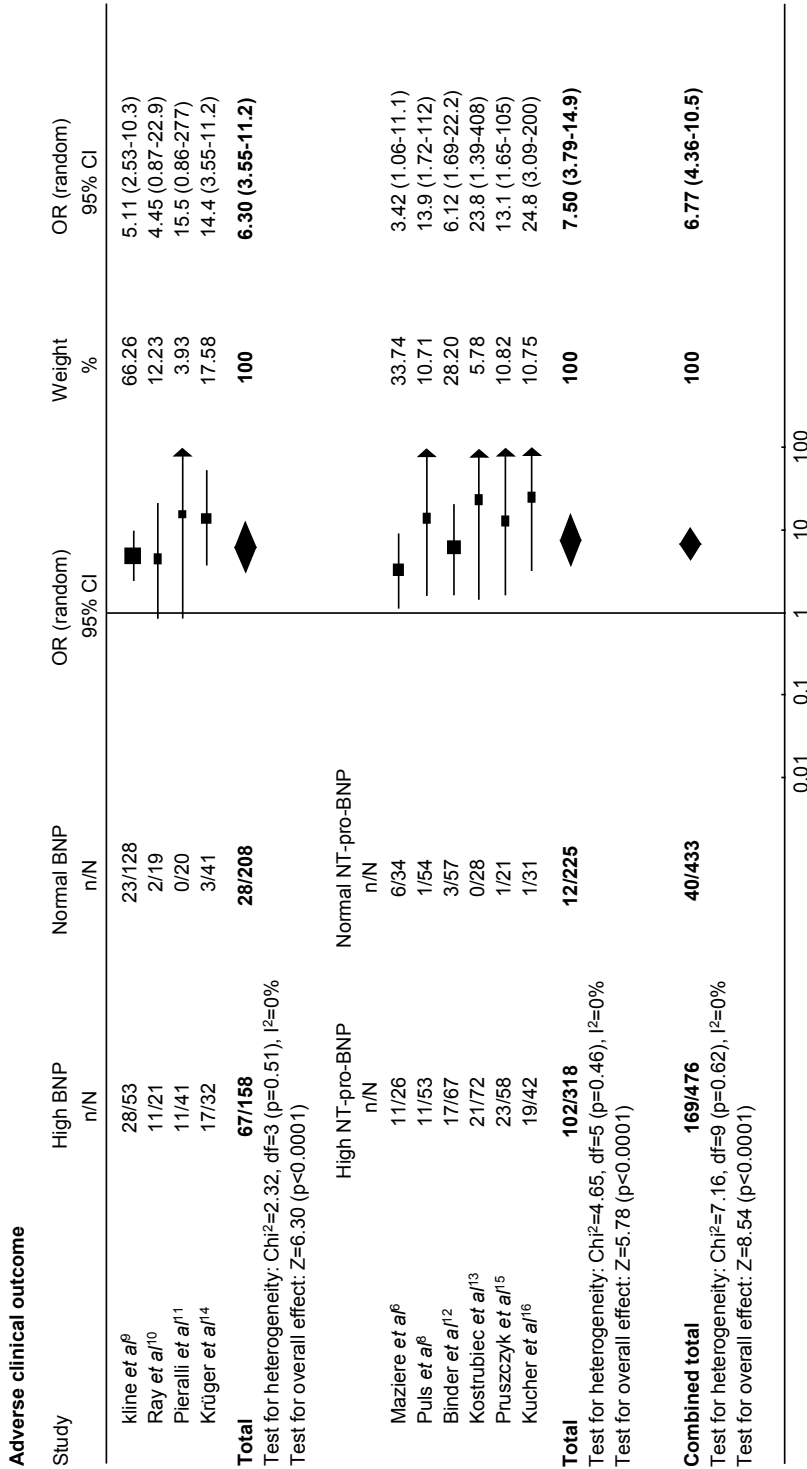
(197 patients).[12,18] The incidence of right ventricular dysfunction was 85% (116 of 137 patients; 95%CI 78-90) and 12% (13 of 106 patients; 95%CI 6.7-20) in patients with and without elevated BNP levels respectively (p<0.0001). A positive association was found between increased concentration of BNP and the presence of right ventricular dysfunction (OR 81; 95%CI 27-238). In NT-pro-BNP studies, the incidence of right ventricular dysfunction was 45% (49 of 109 patients; 95%CI 35-55) in patients with elevated NT-pro-BNP levels compared with 4.5% (4 of 88 patients; 95%CI 1.3-11) in patients with normal NT-pro-BNP levels. Elevated NT-pro-BNP levels were associated with the presence of right ventricular dysfunction (OR, 16.81; 95% CI 5.73 to 49.37). Pooled data of all assays revealed a combined OR of 39 (95%CI 17-89).

DISCUSSION

This meta-analysis demonstrates a significant relation between high levels of (NT-pro-) BNP and deterioration of clinical condition in patients with acute PE. This is physiologically plausible since BNP is released as a reaction to right ventricular stress, which has been shown to predict a non-benign course in patients with PE.[1-3] This relation is also demonstrated in this analysis: we found a very strong correlation between increased levels of (NT-pro-) BNP and right ventricular dysfunction on echocardiography (Figure 4).

There are some points for discussion if (NT-pro-) BNP levels would be incorporated in clinical treatment strategies for patients with acute PE. First, timing of blood sampling has consequences for the established BNP-concentration. The BNP prohormone (pro-BNP) in normal ventricular myocytes is not stored to a significant amount. As a consequence, it takes several hours for the plasma natriuretic peptide levels to increase significantly after the onset of acute myocardial stretch.[20] A very recent onset of complaints could therefore result in false-negative (NT-pro-) BNP test results. Second, many different cutoff levels for (NT-pro-) BNP are proposed in the literature.[21,22] The variation may be related to patient selection, different gender and age.[22] Despite the different cutoff levels and different assays, the prognostic value of both NT-pro- BNP and BNP was consistent in all included studies.

What are the potential implications of our findings? First, normal levels of BNP have a high negative predictive value for unfavorable outcome. Patients with normal levels of (NT-pro-) BNP have low risks for death as well as hemodynamic deterioration resulting in any adverse events. Conversely, elevated concentrations of B-type natriuretic peptides are a nonspecific finding. An explanation for this phenomenon is the elevation of natriuretic peptides in a multitude of other conditions, including preexisting left ventricular dysfunction, higher age, renal impairment and chronic lung disease.[23] The combination of BNP with other clinical risk factors for adverse outcome may improve

Right ventricular dysfunction

Study	High BNP n/N	Normal BNP n/N	OR (random) 95% CI	Weigh %	OR (random) 95% CI
Logeart et al[7]	36/47	0/20		14.14	130 (7.23-2324)
Pieralli et al[11]	35/41	0/20		13.71	224 (12.0-4183)
Krüger et al[14]	16/17	9/25		24.74	28.4 (3.22-254)
Kuger et al[18]	29/32	4/41		47.42	59.4 (18.5-431)
Total	**116/137**	**13/106**		**100**	**80.6 (27.3-238)**

Test for heterogeneity: Chi²=1.47, df=3 (p=0.69), I²=0%
Test for overall effect: Z=7.94 (p<0.0001)

Study	High NT-pro-BNP n/N	Normal NT-pro-BNP n/N		Weigh %	OR (random) 95% CI
Binder et al[12]	30/67	3/57		73.26	14.6 (4.15-51.4)
Kucher et al[16]	19/42	1/31		26.74	24.8 (3.09-199)
Total	**49/109**	**4/88**		**100**	**16.8 (5.73-49.4)**

Test for heterogeneity: Chi²=0.18, df=1 (p=0.67), I²=0%
Test for overall effect: Z=5.14 (p<0.0001)

| **Combined total** | **165/246** | **17/194** | | **100** | **38.6 (16.7-89.2)** |

Test for heterogeneity: Chi²=5.69, df=5 (p=0.34), I²=12.1%
Test for overall effect: Z=8.55 (p<0.0001)
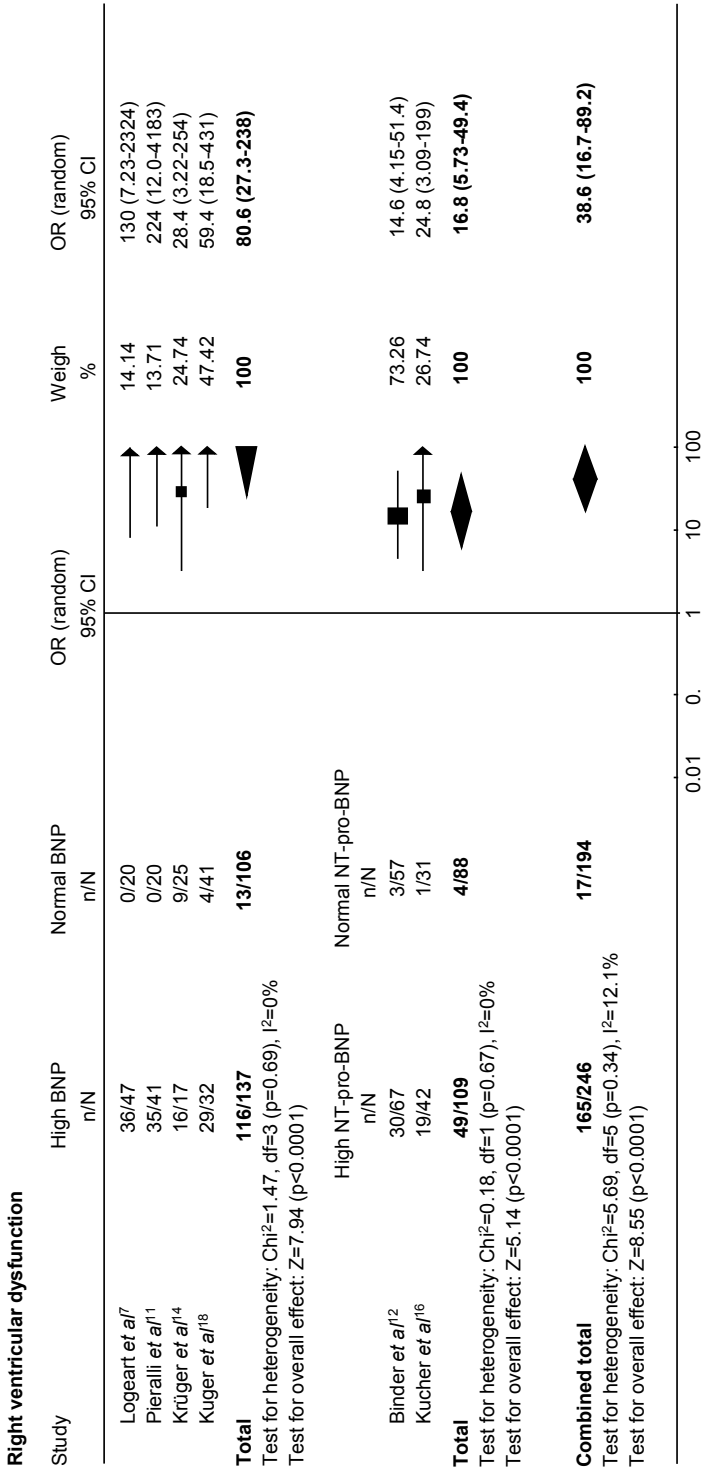
Axis: 0.01 0.1 1 10 100

Figure 4. Odds ratio (OR) for right ventricular dysfunction on echocardiography based on elevated (NT-pro-) BNP levels. Different cutoffs were used for different studies; Mantel-Haenszel methods for combining trials were used for weighting the studies. CI: confidence interval.

sensitivity and positive predictive value for clinical deterioration. Such algorithms for risk stratification would be clinically useful if they were able to identify patients eligible for outpatient management, for standard or intensive in-hospital treatment. Proposals for such algorithms including (bio)markers of right ventricular function, e.g. (NT-pro-)BNP, troponin[4] or heart-type fatty acid-binding protein,[24,25] have been made but not validated prospectively in clinical outcome studies yet.[12,13,26] Future studies are required to determine the clinical benefits of more aggressive treatments in patients with adverse prognosis as identified by these risk stratifications and less intensive treatment including out of hospital treatment in patients with normal values of BNP.

This meta-analysis has limitations. First, included studies used different assays with different retrospectively calculated cutoff points. Second, duration of follow-up and definitions of endpoints varied among the studies. In addition, most studies did not mention completeness of follow-up. Nonetheless, we have included a large cohort of prospectively followed patients (n=1128) and our analysis showed no evidence of heterogeneity between the outcomes of the incorporated studies. Third, the relative risk for mortality is not adjusted for confounding factors, thus part of the effect ascribed to high BNP values may be related to clinical conditions associated with PE. Fourth, we could not determine the ideal cutoff for the two BNP tests because we did not have the raw data to do ROC curves and other analyses. Finally, in the included studies it is not stated whether thrombolytic therapy or ICU admission were the result of the clinical condition or a high (NT-pro-)BNP value.

In summary, an elevated level of (NT-pro-)BNP is a risk factor for short-term mortality, overall short term complicated clinical outcome and an indicator of right ventricular dysfunction in patients with acute PE. It remains to be demonstrated whether it could play a role in risk stratification algorithms identifying patients that could benefit from differentiated forms of therapy, of which thrombolytic therapy and home treatment are two poles of the spectrum.

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