

Acute HEArt Failure Database

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for AHEAD investigators

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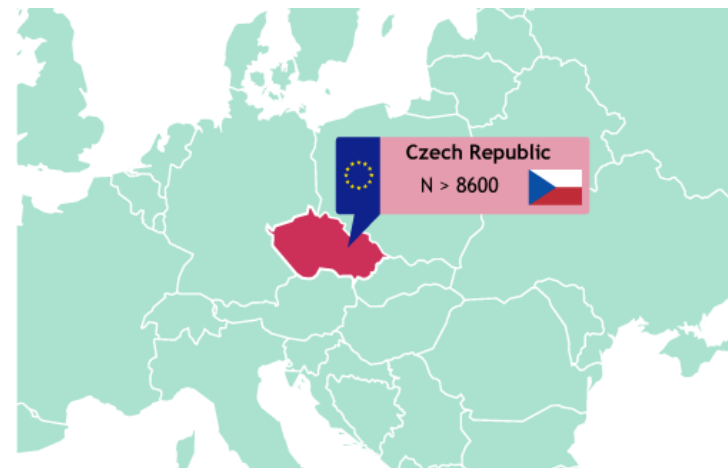
AHEAD database



- AHEAD (Acute HEArt Failure Database)* is database aimed on the monitoring of the ways of diagnosis and treatment of acute heart failure in the Czech Republic
- The data were collected between 2006-2012 and the database consist of two parts with more than 8600 records of Czech patients, 5 859 were first hospitalisation
 - The AHEAD main registry includes consecutive patients in seven centres with a 24-h Cath Lab service and centralized care for patients with acute coronary syndromes (ACS) from a region of about 3 million inhabitants.
 - The AHEAD network also includes five regional hospitals without a Cath Lab service.

* Spinar,; Parenica, J; Vitovec, J; Widimsky, P; Linhart, A; Fedorco, M; Malek, F; Cihalik, C; Spinarova, L; Miklik, R; Felsoci, M; Bambuch, M; Dusek, L; Jarkovsky, J. Baseline characteristics and hospital mortality in the Acute Heart Failure Database (AHEAD) Main registry. CRITICAL CARE Volume: 15 Issue: 6 Article Number: R291 DOI: 10.1186/cc10584 Published: 2011

<http://ahead.registry.cz/>





Acute Heart Failure Database

OK[Project AHEAD](#)[Involved centres](#)[Technology](#)[Results](#)[Subprojects](#)[Contacts](#)[Sitemap](#)[Registry entry](#)

About the project

The [acute heart failure](#) is a clinical syndrome which can be a result of many diseases. The symptoms can occur in patients with or without a previous heart disease. The heart failure can be systolic, diastolic or mixed; the impaired heart function can be manifested even by heart rhythm disorders. This condition is often life-threatening and requires an immediate medical intervention.

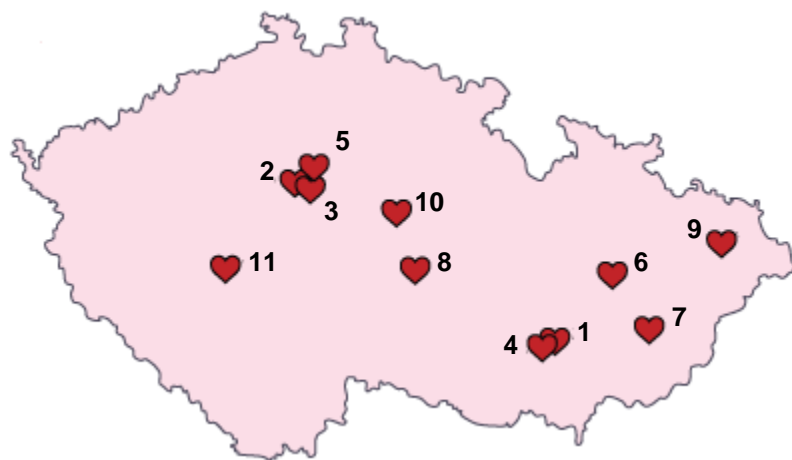
The treatment methods are mostly based on empirical knowledge and are not supported by extensive mortality studies. Registries and databases which were developed in the United States (e.g. ADHERE^[1]) and in Europe (e.g. EHFS^[2]) demonstrate a big diversity of this syndrome and differ significantly in the most important parameter - the mortality.

We have decided to monitor the ways of diagnosis and treatment of acute heart failures in the Czech Republic. We have established the database AHEAD (Acute HEArt Failure Database) which is progressively filled with the patients' records from five big clinical centres: [General University Hospital in Prague](#), [IKEM Prague](#), [St Anne's University Hospital Brno](#), [University Hospital Brno](#) and [University Hospital Olomouc](#). This produces a pilot registry of health care facilities having an angiography unit and concentrating the acute coronary syndrome. The registry includes both heart transplant centres as well.



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1.2 Included centers



2 900/milion AHF hospiti.
2 800/milion ACS hospiti.



Site

1. Faculty Hospital Brno
 2. IKEM Prague
 3. Faculty Hospital Kralovske Vinohrady, Prague
 4. St Anne Faculty Hospital, Brno
 5. General Faculty Hospital, Prague
 6. Faculty Hospital, Olomouc
 7. Zlín
 8. Havlíčkův Brod
 9. Frýdek Místek
 10. Čáslav
 11. Příbram
-

RESEARCH

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Baseline characteristics and hospital mortality in the Acute Heart Failure Database (AHEAD) Main registry

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Abstract

Introduction: The prognosis of patients hospitalized with acute heart failure (AHF) is poor and risk stratification may help clinicians guide care. The objectives of the Acute Heart Failure Database (AHEAD) registry are to assess patient characteristics, etiology, treatment and outcome of AHF.

Český registr AHEAD bodoval v Mnichově hned čtyřikrát

Na evropském kardiologickém kongresu v Mnichově byly hned čtyři české práce založeny na datech z registru AHEAD (Accute HEArt failure Database), který popisuje reálnou praxi péče o pacienty s akutním srdečním selháním. Spravuje jej Institut biostatistiky a analýz Masarykovy univerzity.

U zrodu registru AHEAD stála především Fakultní nemocnice Brno, v současnosti na něm participuje jedenáct nemocnic z celé České republiky. Tato databáze sleduje v každém záznamu asi sto parametrů důležitých pro krátkodobou a dlouhodobou prognózu nemocných.

Mezi ústními sděleními zazněly v Mnichově dvě takové práce, první přednesla RNDr. Simona Littnerová z Institutu biostatistiky a analýz. Týkala se především metodologie, konkrétně pak propensity

score u tzv. obezitologického paradoxu. Ačkoli je obezita potvrzena jako nezávislý prognostický faktor progresu srdečního selhání, podle řady recentních studií je u obézních pacientů riziko úmrtí nižší. Propensity score představuje statistickou metodu, která dovede odlišit, do jaké míry je jeden parametr (tělesná hmotnost) ovlivnitelný parametrem jiným (např. pohlaví).

Prof. MUDr. Lenka Špinarová, Ph.D., z I. interní kardiologické kliniky FN u sv. Anny se zaměřila na jiný tako-

vý paradox – cholesterolový. Ten ukazuje, že na rozdíl od zdravé populace nebo populace hypertoniků pacienti se srdečním selháním a nízkým cholesterolem mají mnohem horší prognózu než pacienti s cholesterolem vysokým, a to nejen pokud jde o krátkodobou hospitalizační mortalitu, ale i o mortalitu dlouhodobou. Unikátní na sdělení prof. Špinarové byla jednak velikost souboru, jednak délka sledování. Ta u některých pacientů dosahuje až sedm let, prezentace na ESC pracovala s daty nemocných z let 2006 až 2009.

Cholesterolový paradox je popsán poměrně dobře, stále však nejsou jasné jeho klinické implikace. „V žádném případě by to nemělo vést k tomu, abychom nemocným se srdečním selháním vysazovali statiny. Nezdá se také, že by to byla pouze otáz-

ka kachexie. Nepodařilo se prokázat, že by pacienti s nízkým cholesterolem měli body mass index například pod dvacet. Takže zatím jde opravdu o paradox,“ komentuje to prof. MUDr. Jindřich Špinar, CSc., přednosta Interní kardiologické kliniky FN Brno.

On sám se svým týmem v posterové sekci prezentoval hlavní závěry plynoucí z registru AHEAD: „Nejsou překvapující, jde spíše o rozvíjení známých faktů, že hodnoty natriuretických peptidů, ejekční frakce, renální funkce a věk jsou základní rizikové faktory.“

V posterové sekci MUDr. Jan Václavík, Ph.D., z I. interní kliniky FN Olomouc využil data z registru AHEAD k analýze, jaké nálezy na EKG predikují špatnou prognózu u srdečního selhání. Nejsilnějšími negativními prognostickými faktory

byly prodloužení QRS komplexu a junkční rytmus, které oba nezávisle predikovaly jak hospitalizační, tak dlouhodobou mortalitu.

Délka sledování je obrovskou devizou registru i do budoucna. „Víceméně se potvrzuje, že prognóza pacientů se srdečním selháním je mnohdy horší než u nemocných s malignitou. Registr je nástrojem, jenž nám může pomoci s tímto stavem něco dělat. Pořád se zdůrazňuje, že kardiologie je vítězný, úspěšný obor. Tito nemocní dostávají pět šest léků, a přesto jich do roka 30 % zemře. Když člověk sleduje vlastní pacienty, má pocit, že situace není tak zlá. Když se však vezme neselektovaná populace úplně všech, kteří jsou pak předáváni mimo kliniku (většinou starší a polymorbidní), zjistí se, že třetina do roka zemřela,“ uvedl prof. Špinar. lon

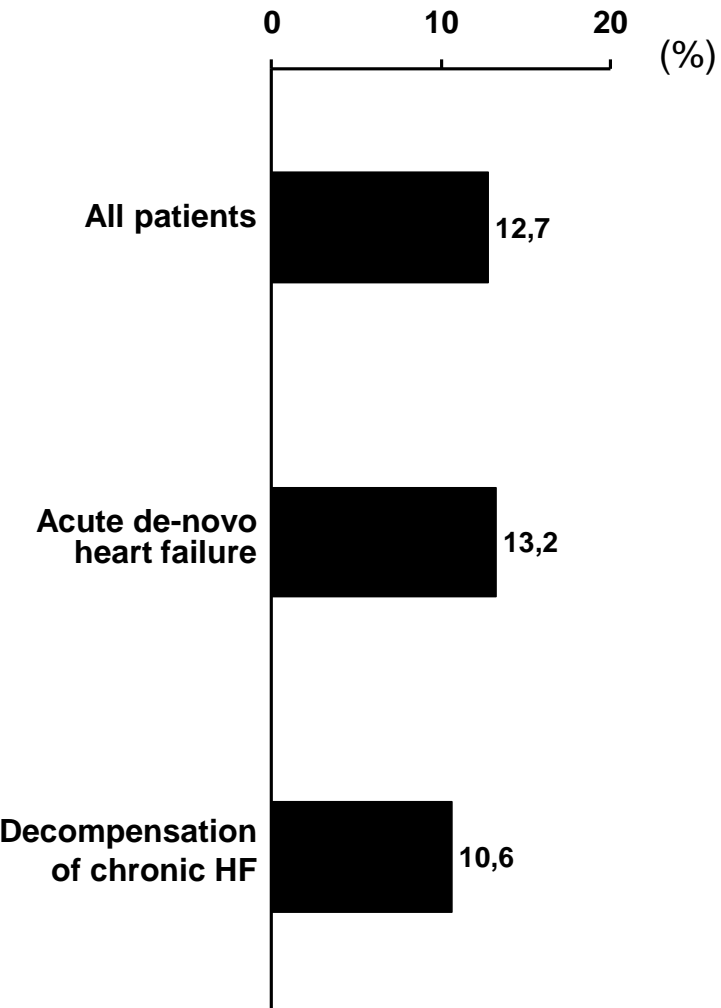
Results and publications

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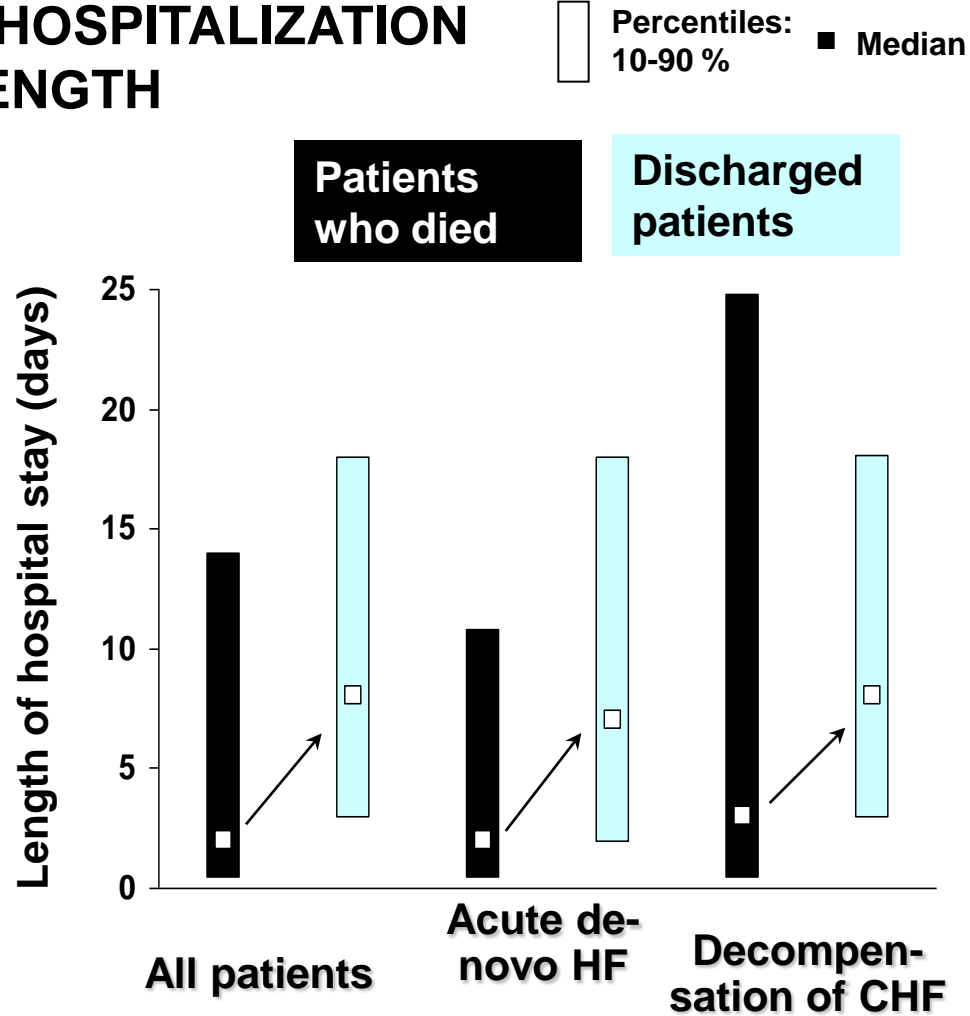
3.1 Key parameters: in-hospital mortality and length of hospital stay



❖ TOTAL MORTALITY



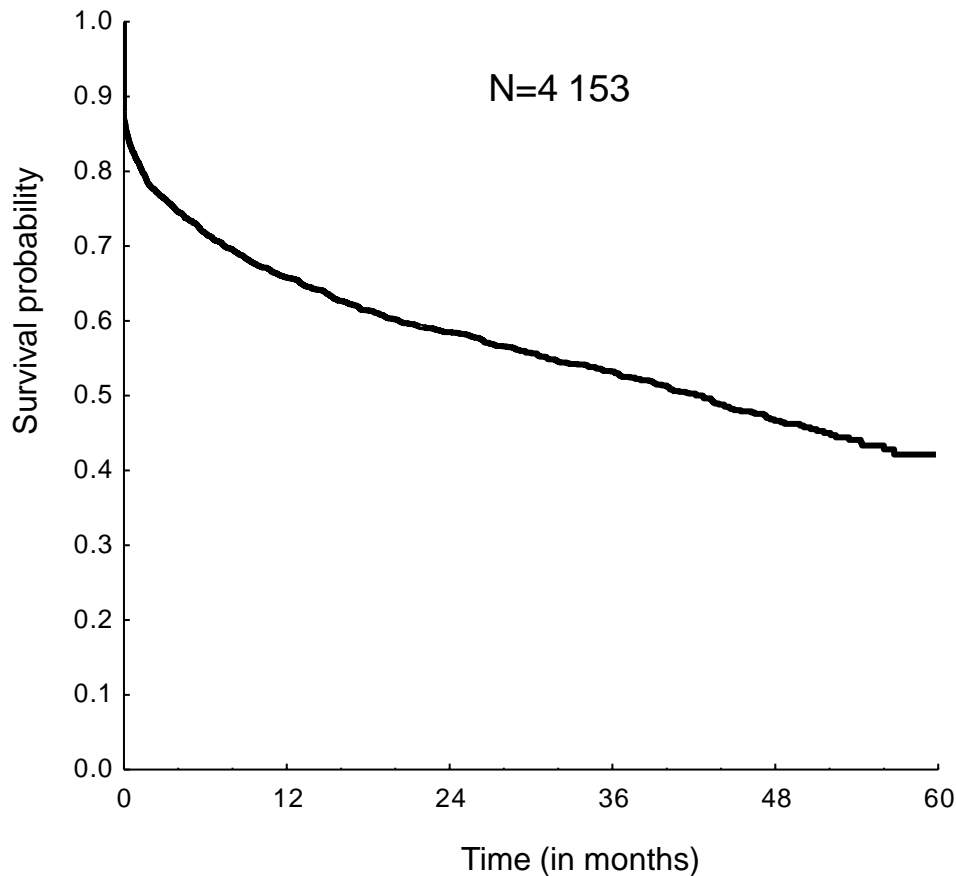
❖ HOSPITALIZATION LENGTH



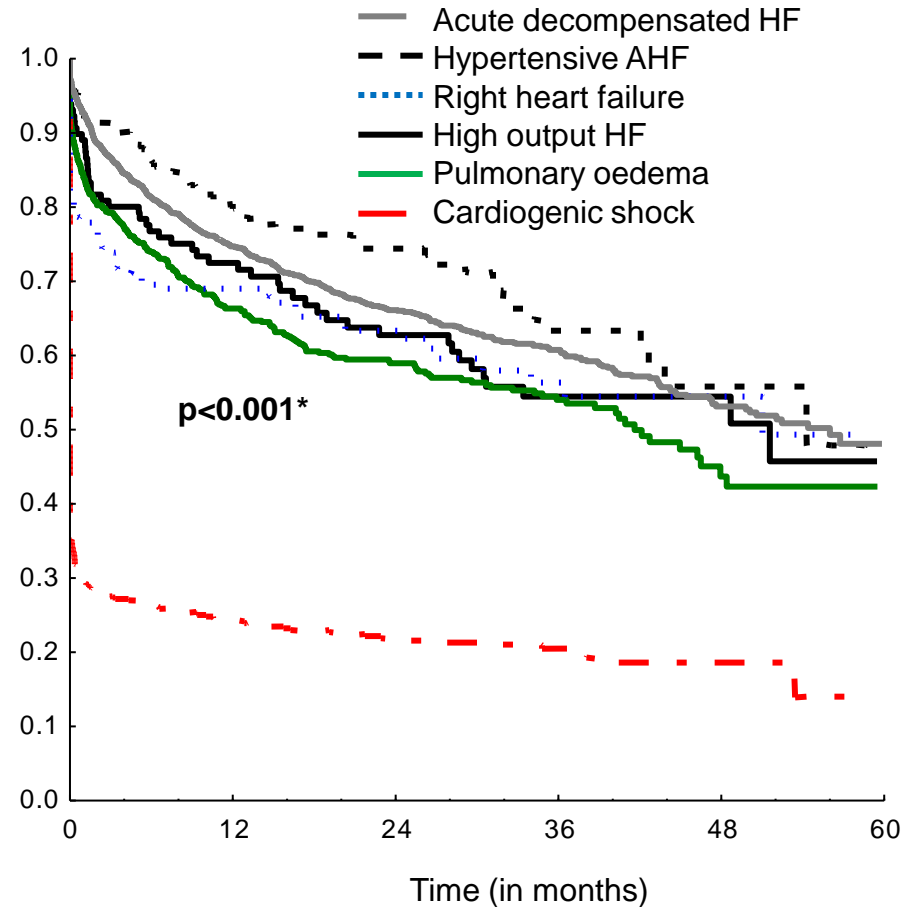
Median of hospital stay was 2 days by patients who died during hospitalization and 8 days by discharged patients (p<0.001).

6.1 Overall long-term survival stratified by syndromes in the whole dataset.

A) Overall survival

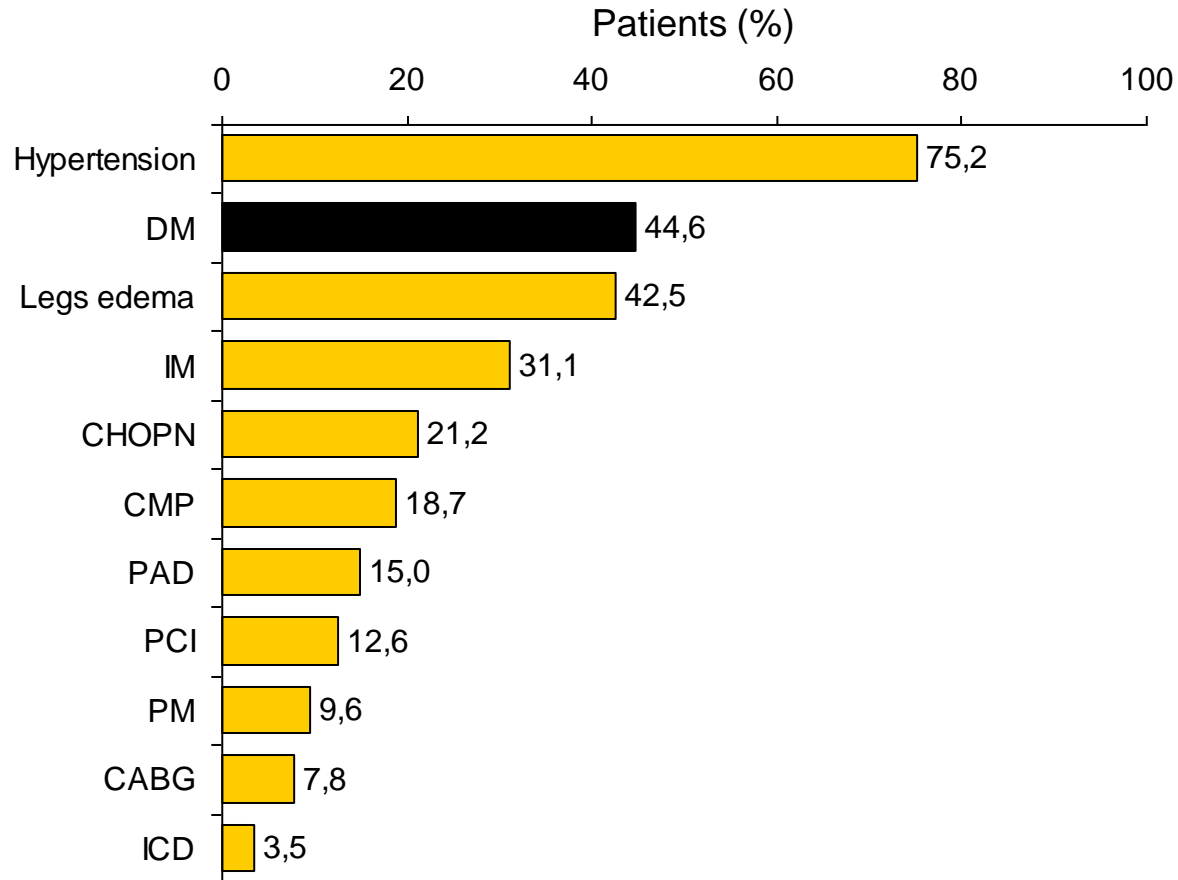


B) Survival by syndromes



Medical history

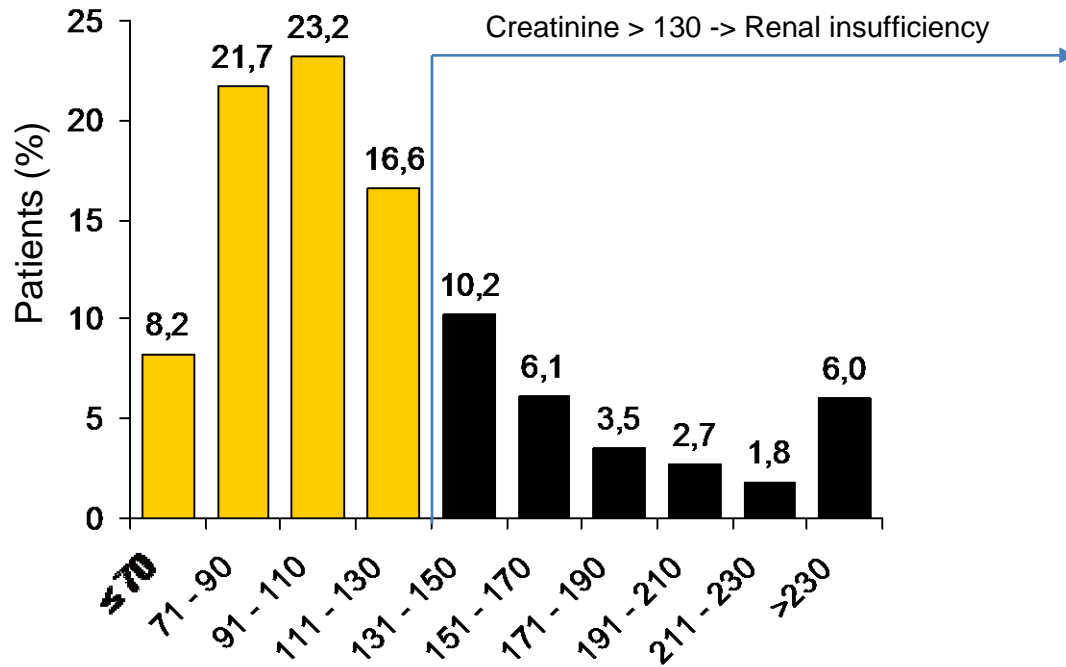
N=5859



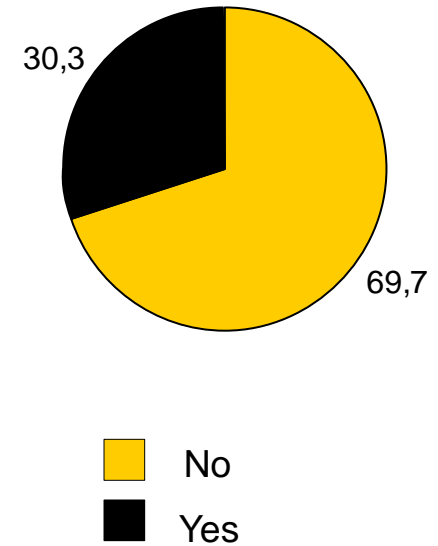
Creatinine and kidney function

N=5859

Creatinine at admission

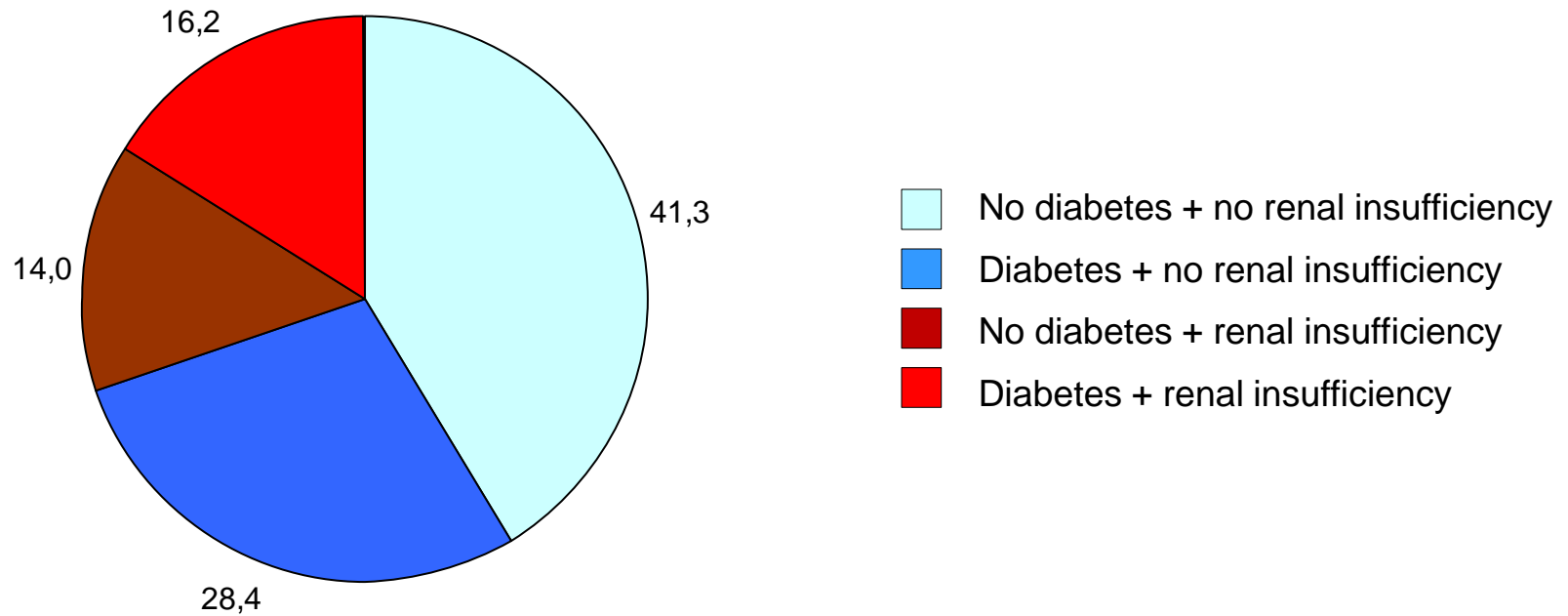


Renal insufficiency



Diabetes and renal insufficiency

N=5859

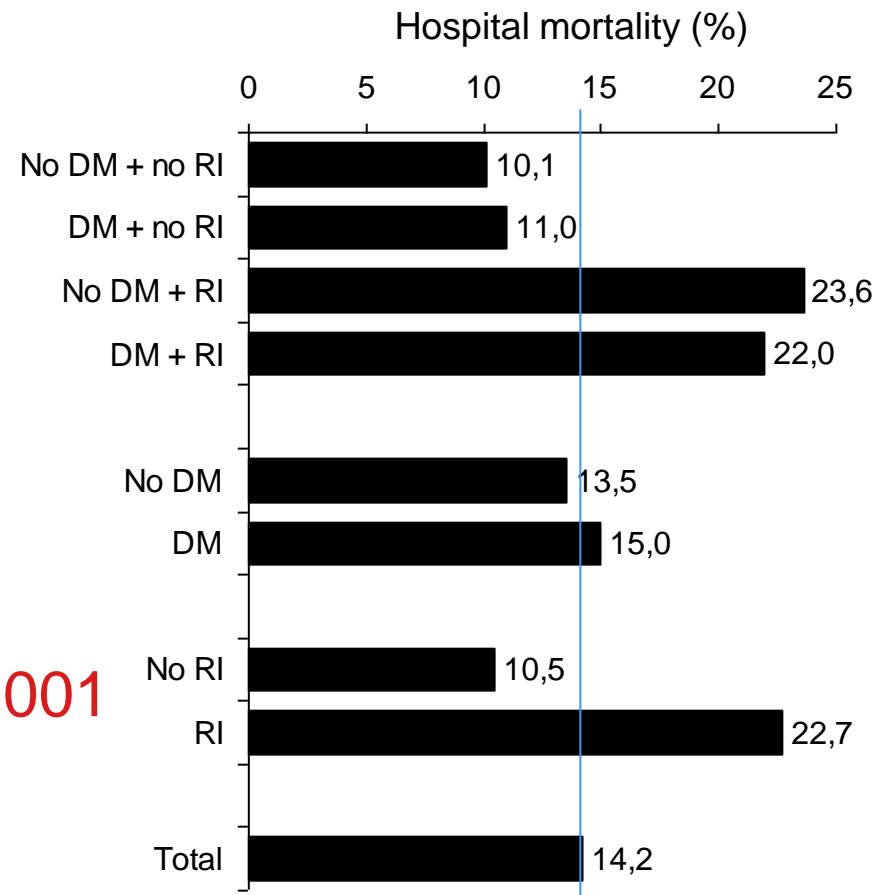
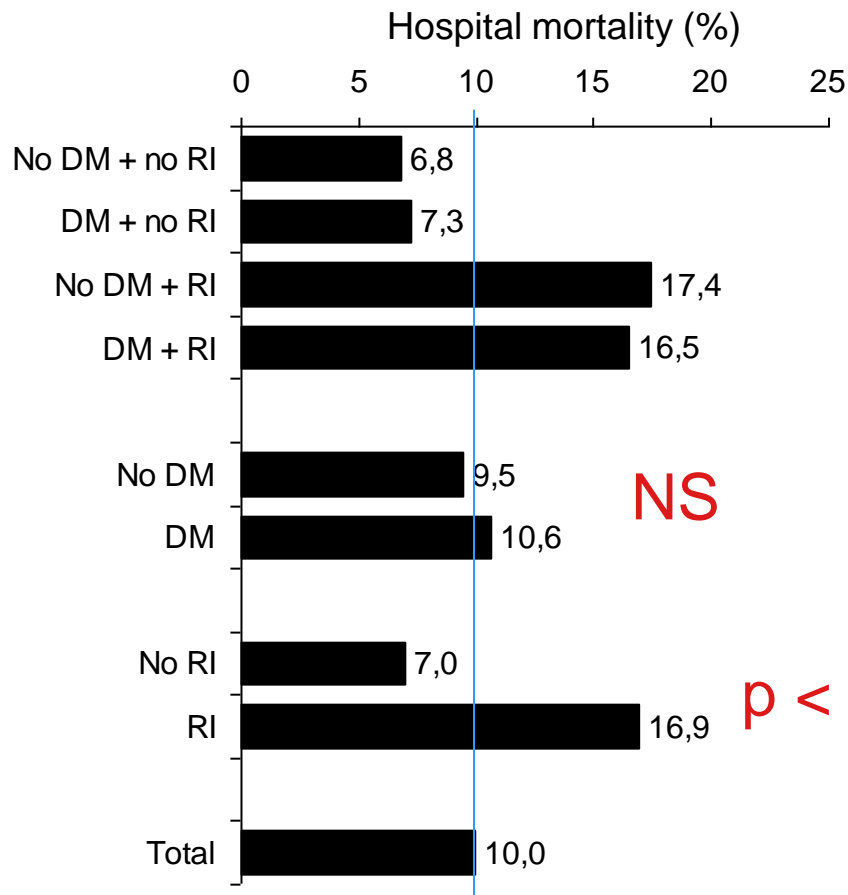


Hospitalisation and 30th day mortality according to diabetes and renal insufficiency

N=5859

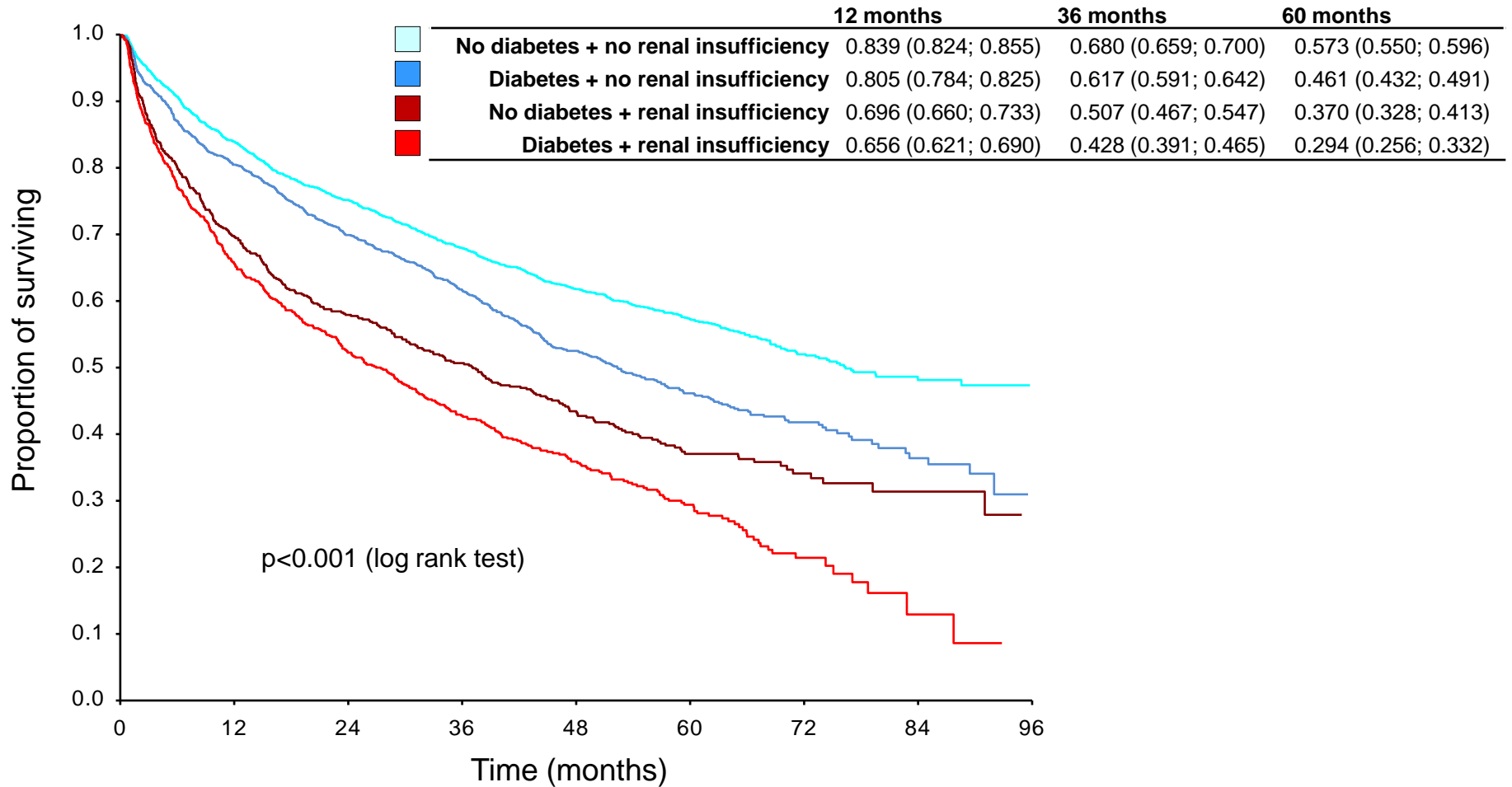
Hospital mortality

30th day mortality



Survival after 30th day according to diabetes and renal insufficiency

N=5028 (patients surviving after 30th day)



AKIN



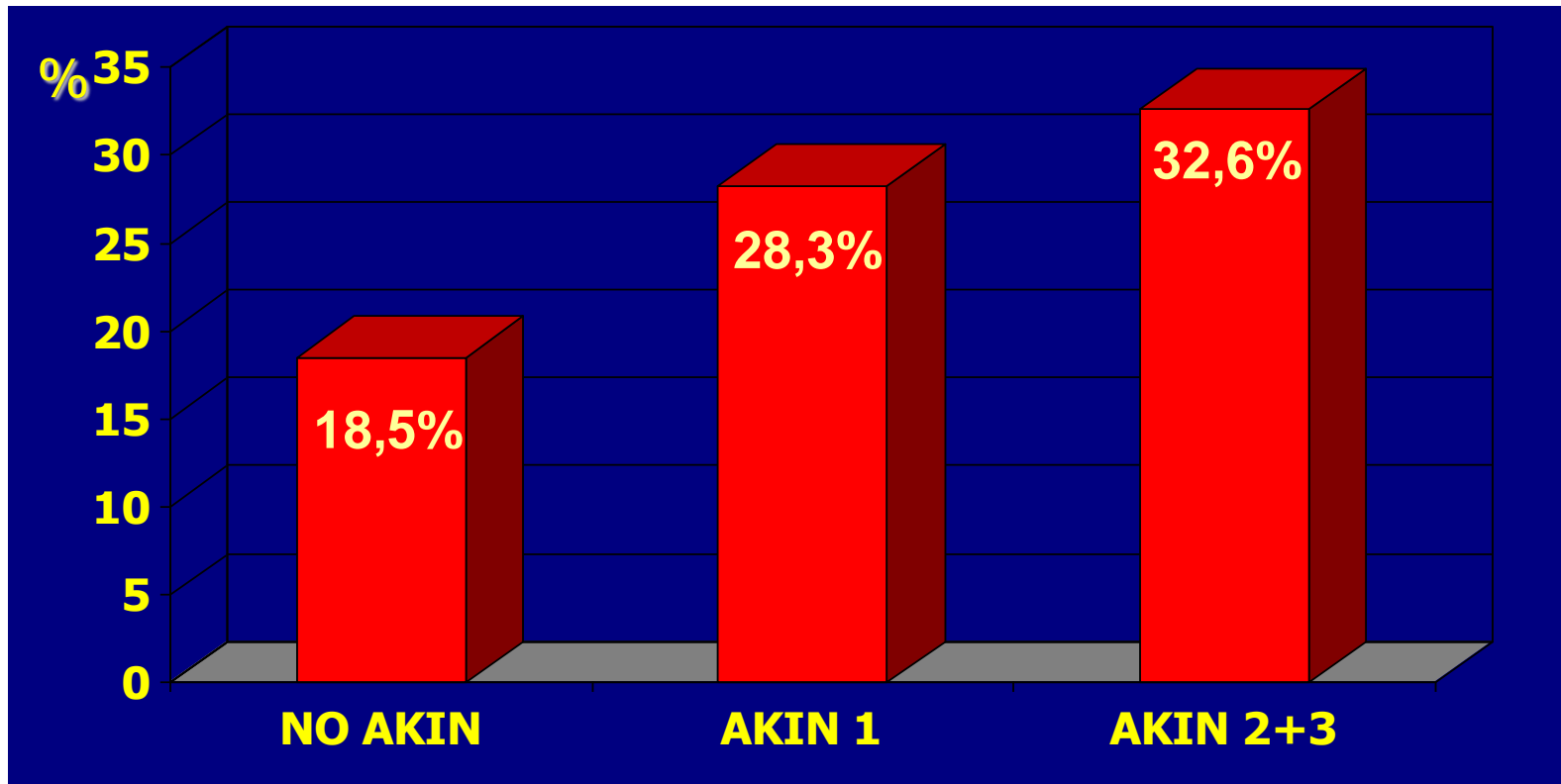
Definition and classification/staging system for acute kidney injury (AKI)*.

AKI stage	Creatinine criteria	Urine output criteria
AKI stage I	Increase of serum creatinine by ≥ 0.3 mg/dl (≥ 26.4 $\mu\text{mol/L}$) or increase to $\geq 150\%$ – 200% from baseline	Urine output < 0.5 ml/kg/hour for > 6 hours
AKI stage II	Increase of serum creatinine to $> 200\%$ – 300% from baseline	Urine output < 0.5 ml/kg/hour for > 12 hours
AKI stage III	Increase of serum creatinine to $> 300\%$ from baseline or serum creatinine ≥ 4.0 mg/dl (≥ 354 $\mu\text{mol/L}$) after a rise of at least 44 $\mu\text{mol/L}$ or treatment with renal replacement therapy	Urine output < 0.3 ml/kg/hour for > 24 hours or anuria for 12 hours

Diagnostic criteria for AKI includes an abrupt (within 48 hours) reduction in kidney function defined as an absolute increase in serum creatinine of either 0.3 mg/dl or more (≥ 26.4 $\mu\text{mol/L}$) or a percentage increase of 50% or more (1.5 fold from baseline) or a reduction in urine output.

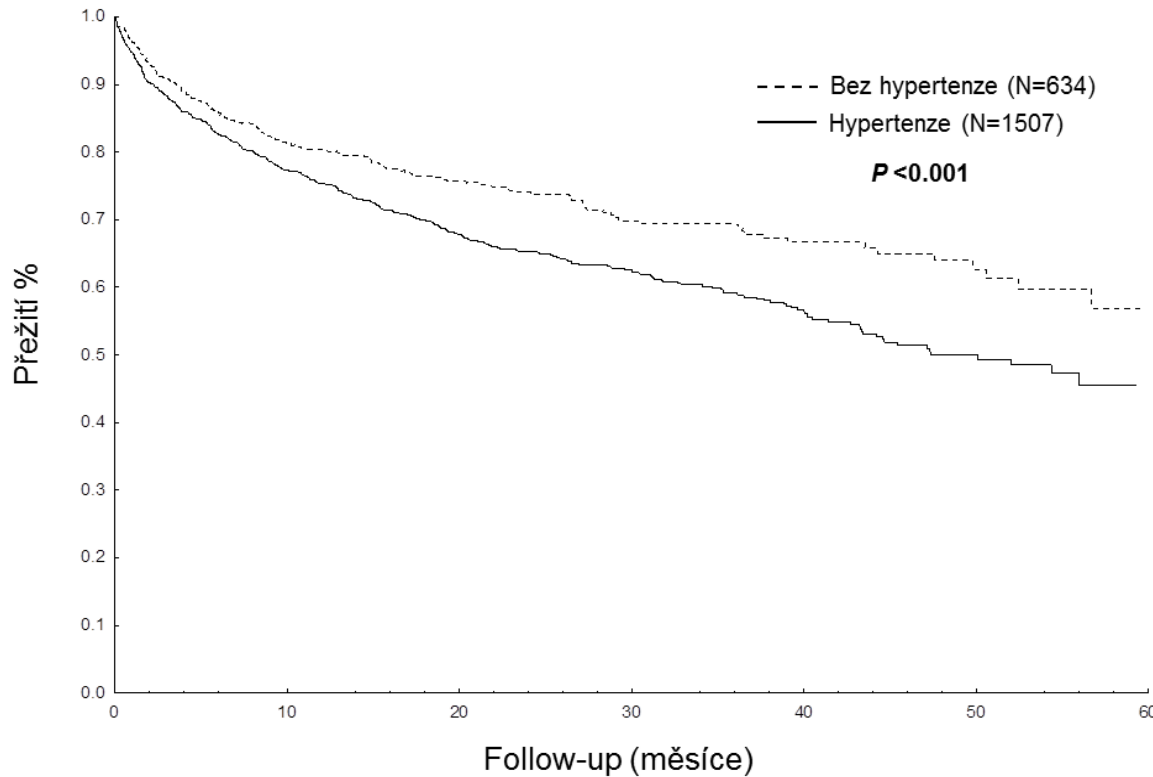
*according to Mehta and colleagues [3]

AHEAD (n = 3 941) (One year mortality)



HYPERTENZE

- Střednědobé (1, 2, 3-leté) přežívání – Killip II



non-HT

80.7% / 74.2% / 69.8%

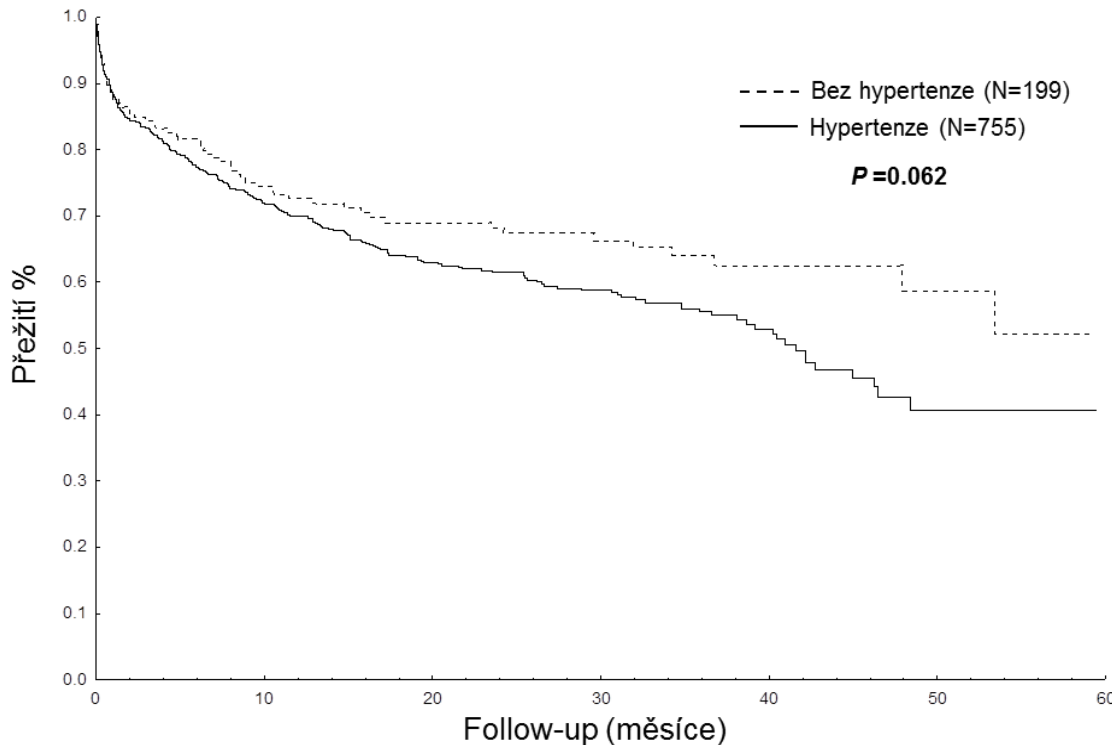
HT

75.6% / 65.9% / 58.7%

$P < 0.001$

HYPERTENZE

- Střednědobé (1, 2, 3-leté) přežívání – **Killip III+IV**



non-HT

72.6% / 68.2% / 64.0%

HT

70.0% / 61.5% / 55.5%

P=0.062

Prediktory mortality Killip II

	30 denní mortalita		Středně-dobá mortalita	
	OR (95% CI)	P	HR (95% CI)	P
Věk > 70 let	1.64 (1.12 - 2.40)	0.012	1.86 (1.51 - 2.28)	<0.001
BMI ≥ 25 (kg/m ²)	-	-	0.68 (0.56 - 0.83)	<0.001
De novo akutní SS	-	-	0.62 (0.51 - 0.75)	<0.001
Diabetes mellitus	-	-	1.45 (1.20 - 1.74)	<0.001
Systolický TK při přijetí ≤ 100 (mmHg)	1.70 (0.97 - 2.95)	0.062	1.54 (1.14 - 2.07)	0.005
Diastolický TK při přijetí ≤ 60 (mmHg)	1.90 (1.15 - 3.14)	0.012	-	-
Ejekční frakce levé kormory ≤ 30%	-	-	1.28 (1.04 - 1.56)	0.017
Kreatinin při přijetí ≥ 120 (μmol/l)	1.44 (1.01 - 2.06)	0.044	1.634 (1.35 - 1.98)	<0.001
Vstupní Na ⁺ ≤ 130 (mmol/l)	4.52 (2.64 - 7.76)	<0.001	1.78 (1.20 - 2.61)	0.004
Vstupní hemoglobin < 120 (ženy) /130 (muži) (g/l)	-	-	1.40 (1.16 - 1.70)	<0.001
<i>Přítomnost dlouhodobé hypertenze v anamnéze</i>	1.32 (0.87 – 2.01)	0.197	1.20 (0.81 – 1.76)	0.366

Prediktory mortality Killip III+IV

	30 denní mortalita		Středně dobá mortalita	
	OR (95% CI)	P	HR (95% CI)	P
Věk > 70 let	2.27 (1.65 - 3.12)	<0.001	1.68 (1.28 - 2.21)	<0.001
De-novo akutní srdeční selhání	-	-	0.71 (0.55 - 0.92)	0.009
Tepová frekvence při příjmu > 100 (úderů/min)	0.65 (0.47 - 0.89)	0.007		
Fibrilace nebo flutter síní na vstupním EKG	-	-	1.36 (1.02 - 1.82)	0.036
Vstupní kreatinin ≥ 120 (μmol/l)	1.95 (1.44 - 2.64)	<0.001	1.79 (1.39 - 2.31)	<0.001
Vstupní hemoglobin < 120 (ženy) /130 (muži) (g/l)	-	-	1.59 (1.23 - 2.05)	<0.001
Ejekční frakce levé komory ≤ 30%	1.71 (1.26 - 2.33)	<0.001	-	-
Umělá plicní ventilace za hospitalizace	4.03 (2.92 - 5.57)	<0.001	-	-
Noradrenalin/dobutamin/dopamin/levosimendan za hospitalizace	4.80 (3.45 - 6.69)	<0.001	-	-
Přítomnost dlouhodobé hypertenze v anamnéze	0.88 (0.62 - 1.26)	0.496	1.08 (0.78 - 1.50)	0.642

HYPERTENZE

- Pacienti s dlouhodobou hypertenzí jsou **starší**, s vyšším zastoupením **žen**, i přes lepší systolickou funkci LK jsou v horší funkční třídě **NYHA**
- vyšší výskyt **přidružených onemocnění**, horší laboratorní profil (↓renální funkce)
- agresivnější farmakoterapie
- hypertenze není **nezávislý** prognostický faktor

5. Comorbidities: AHEAD score





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AHEAD score – Long-term risk clasification in acute heart failure☆

[Jindrich Spinar](#)¹, [Jiri Jarkovsky](#)¹, [Lenka Spinarova](#), [Alexandre Mebazaa](#), [Etienne Gayat](#), [Jiri Vitovec](#), [Ales Linhart](#), [Petr Widimsky](#), [Roman Miklik](#), [Kamil Zeman](#), [Jan Belohlavek](#), [Filip Malek](#), [Marian Felsoci](#), [Jiri Kettner](#), [Petr Ostadal](#), [Cestmir Cihalik](#), [Jan Vaclavik](#), [Miloš Táborský](#), [Ladislav Dusek](#), [Simona Littnerova](#), [Jiri Parenica](#)

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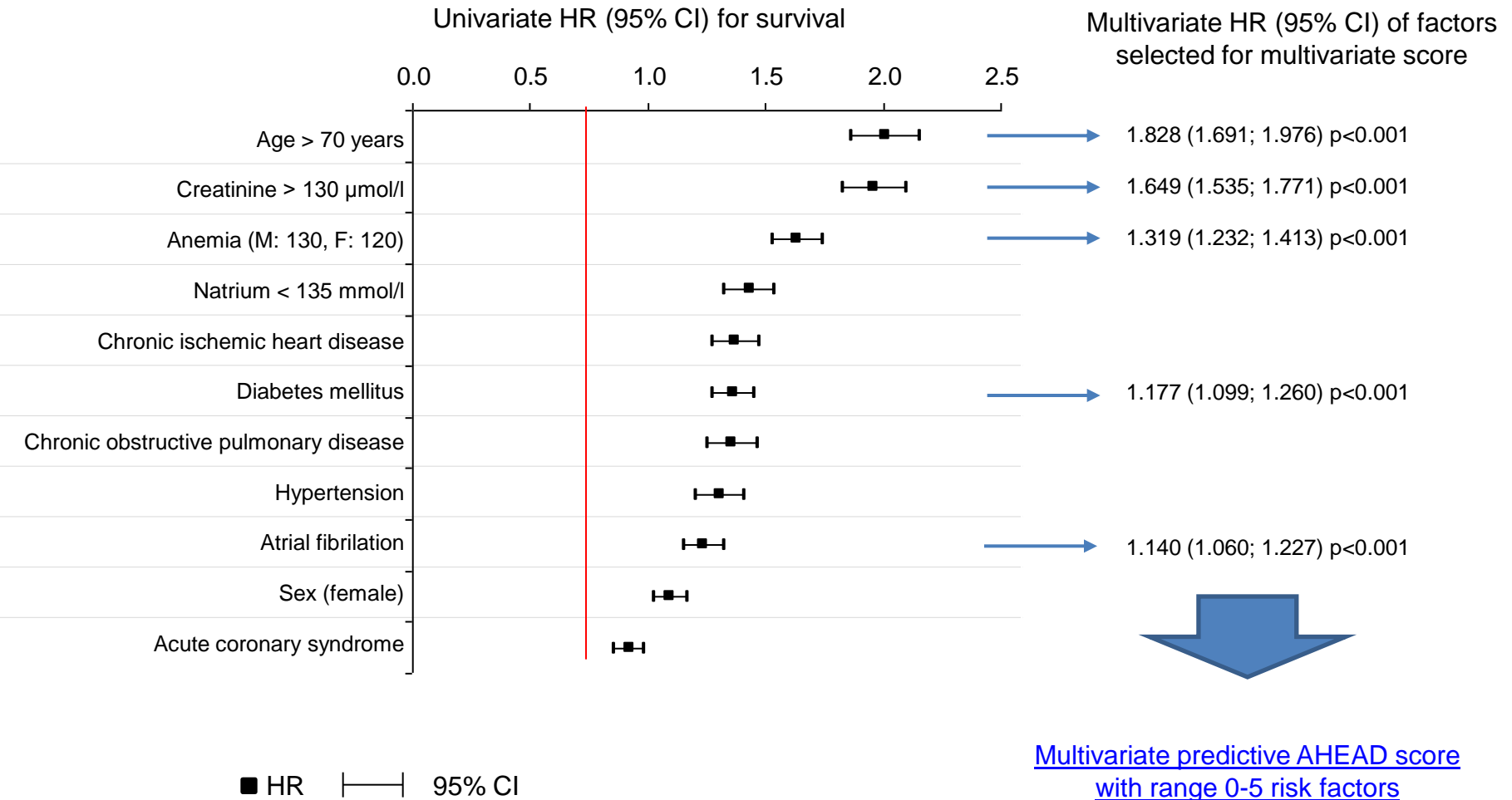
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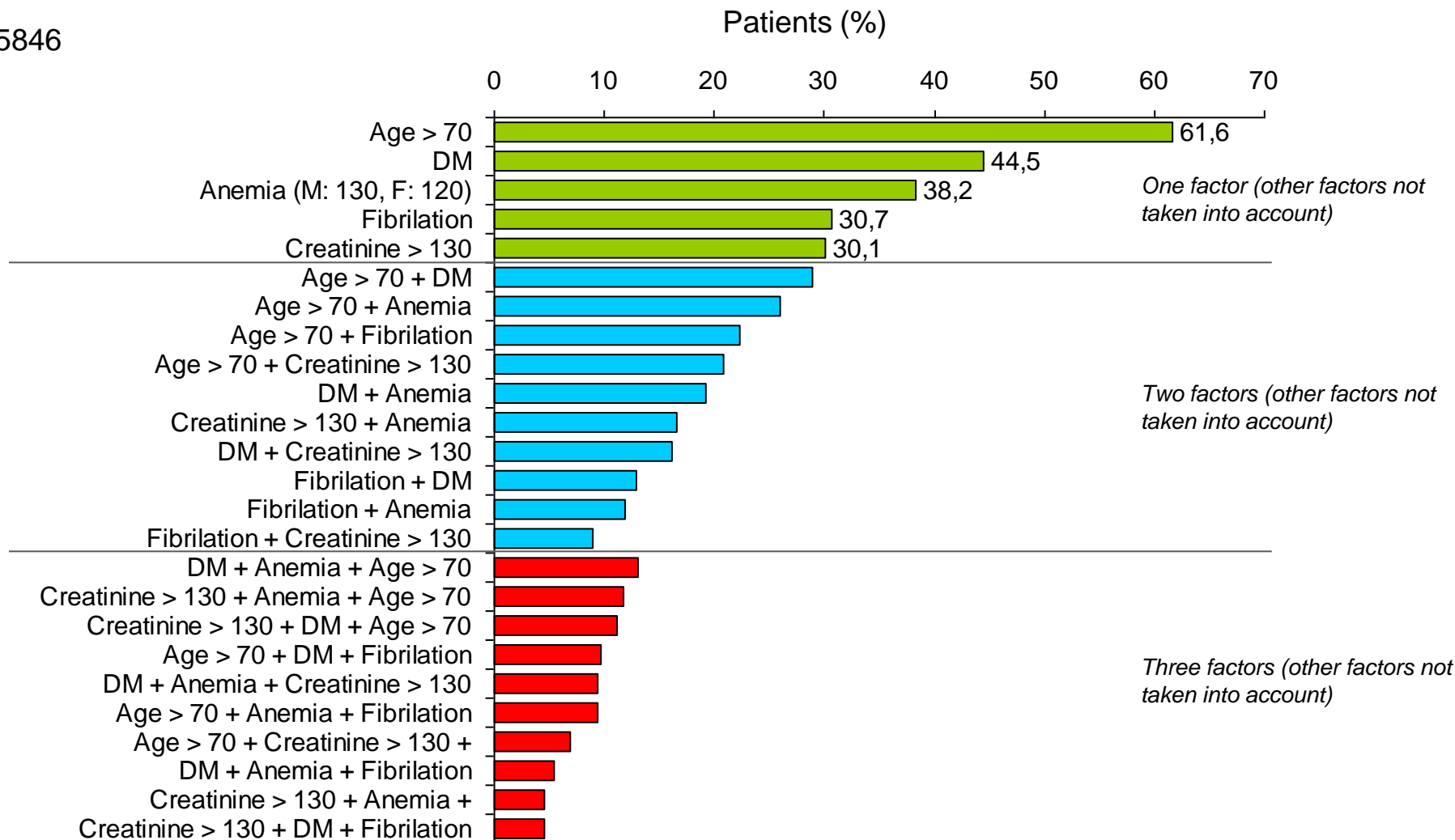
Identification of independent risk factors

N=5846



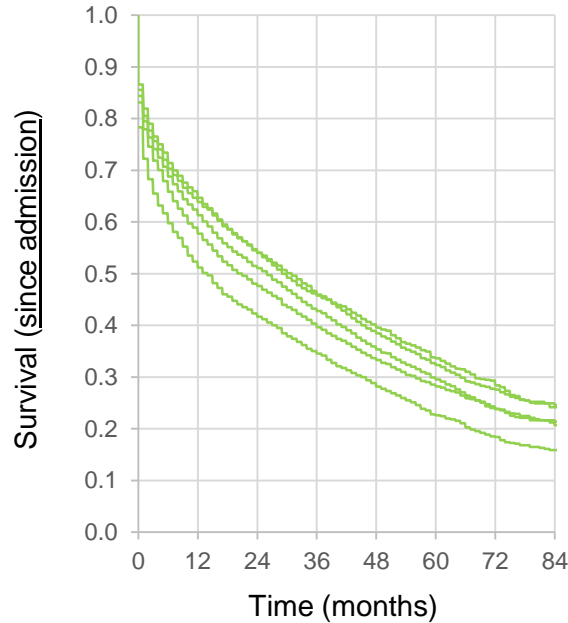
Frequency of selected risk factors combinations

N=5846

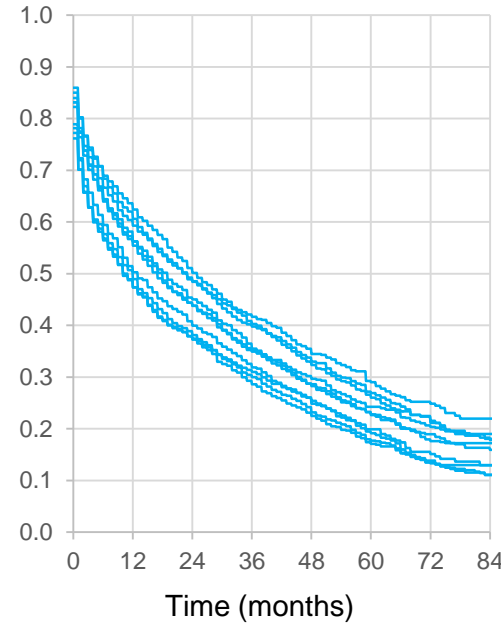


Survival of patients according to risk factors combinations I

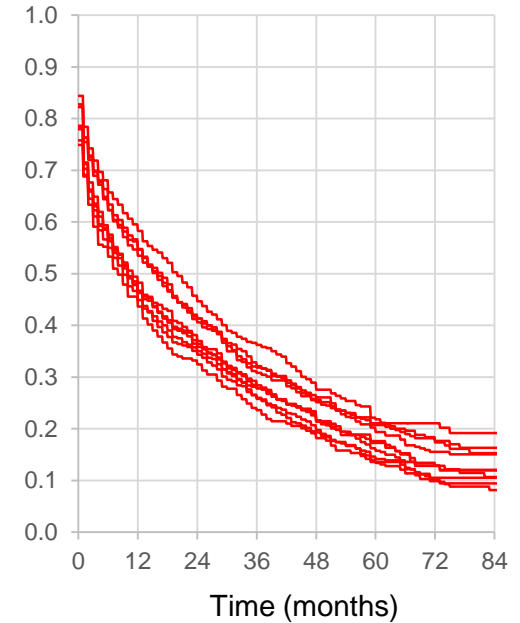
One factor (other factors not taken into account)



Two factors (other factors not taken into account)



Three factors (other factors not taken into account)



- Fibrillation
DM
Age > 70
Anemia
Creatinine > 130

- Fibrillation + DM
Age > 70 + Fibrillation
Age > 70 + DM
DM + Anemia
Fibrillation + Anemia
Age > 70 + Anemia
DM + Creatinine > 130
Fibrillation + Creatinine > 130
Creatinine > 130 + Anemia
Age > 70 + Creatinine > 130

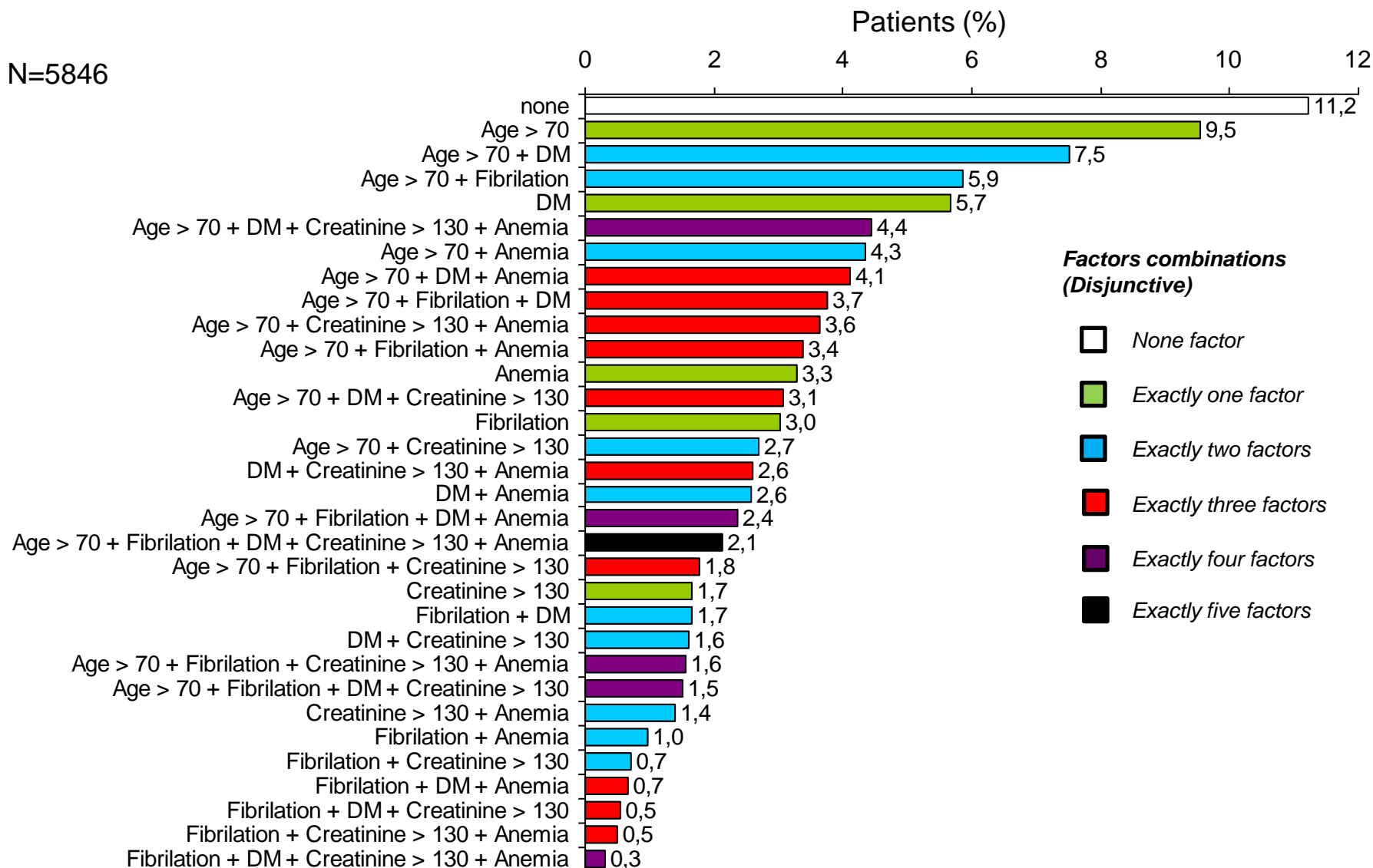
- Age > 70 + DM + Fibrillation
Age > 70 + Anemia + Fibrillation
DM + Anemia + Fibrillation
DM + Anemia + Age > 70
Creatinine > 130 + DM + Fibrillation
DM + Anemia + Creatinine > 130
Creatinine > 130 + DM + Age > 70
Age > 70 + Creatinine > 130 + Fibrillation
Creatinine > 130 + Anemia + Age > 70
Creatinine > 130 + Anemia + Fibrillation

Factors combinations ordered according to 1 year survival

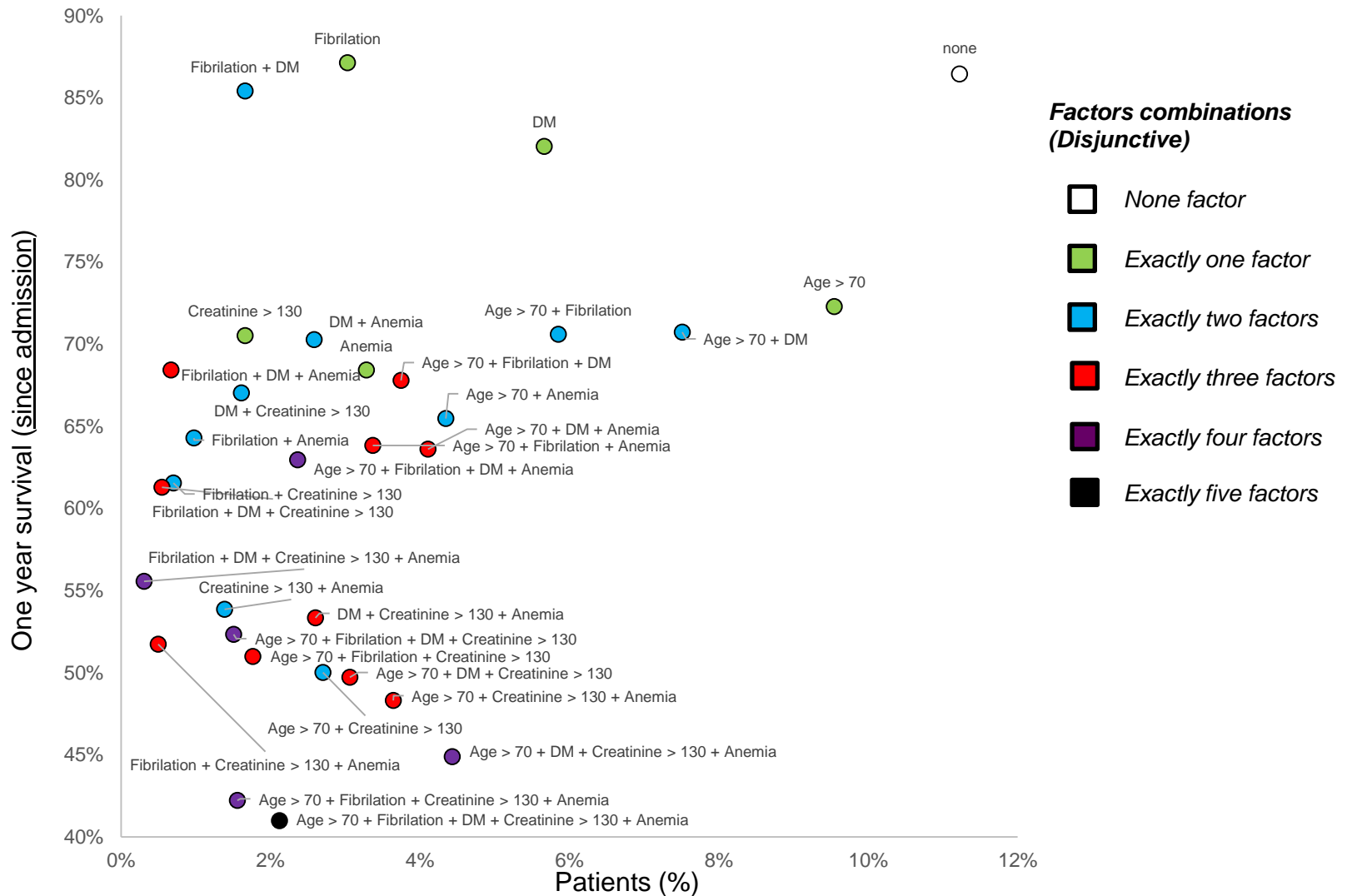


Frequency of selected risk factors combinations

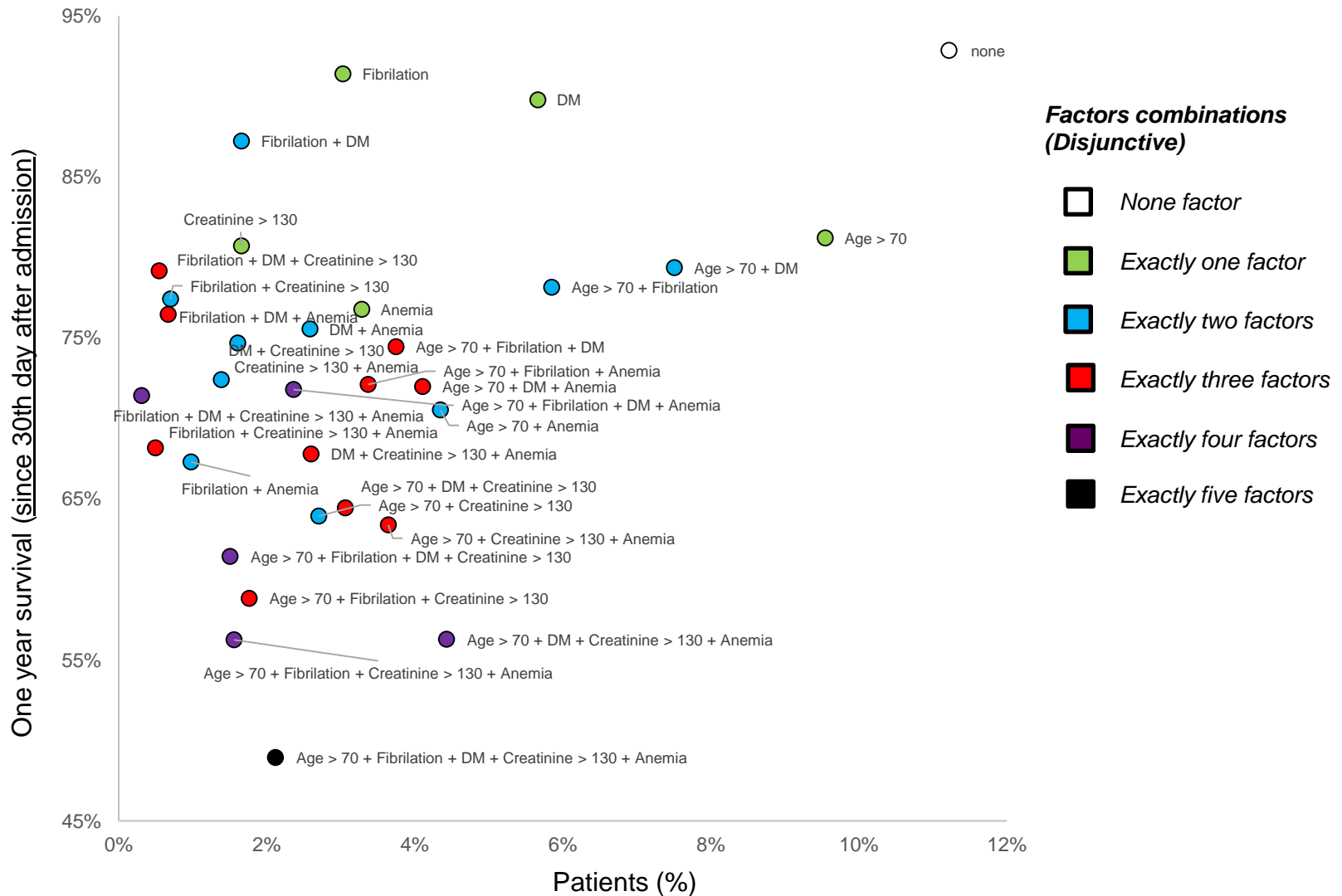
N=5846



Survival and frequency of patients according to risk factors combinations I



Survival and frequency of patients according to risk factors combinations II



A

A = atrial fibrilation

H

H = haemoglobin (< 120g/l F, < 130g/l M)

E

E = elderly > 70 years

A

A = abnorm. renal func. (creatinin > 130 umol/l)

D

D = diabetes mellitus



AHEAD score

The multivariate analysis identified 5 independent risk factors, for each risk factors the patient receive 1 point for the risk score computation:

A = Atrial fibrillation

H = Heamoglobin < 120 g/l (F)
130 g/l (M)

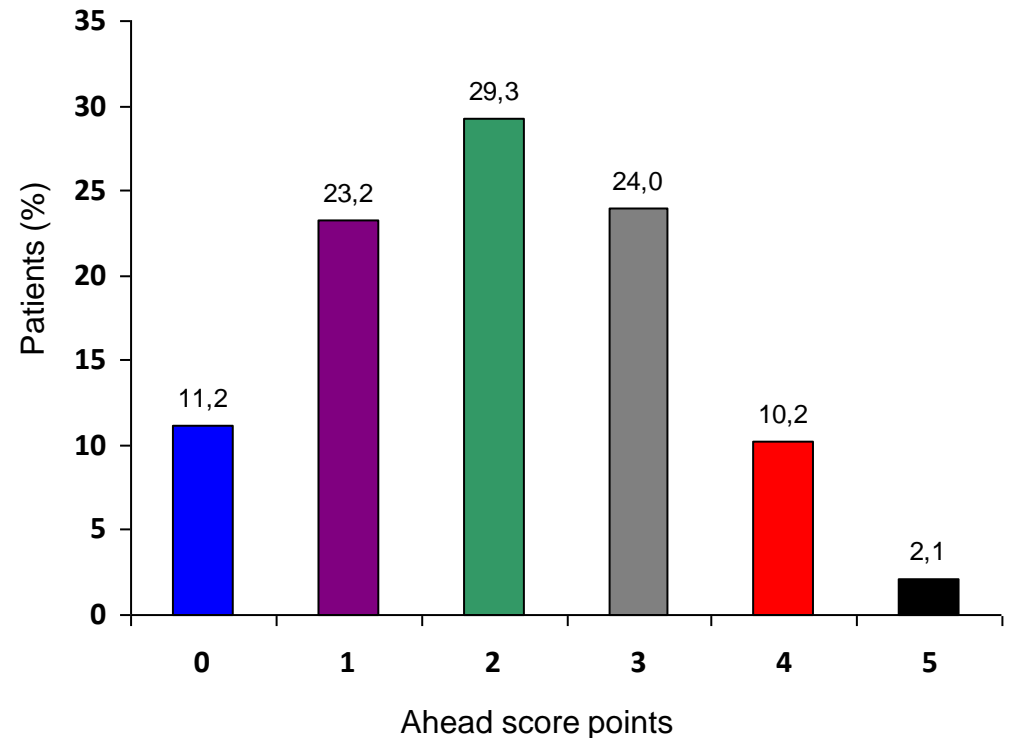
E = Elderly > 70 years

A = Abnormal renal functions
(creat > 130 umol/l)

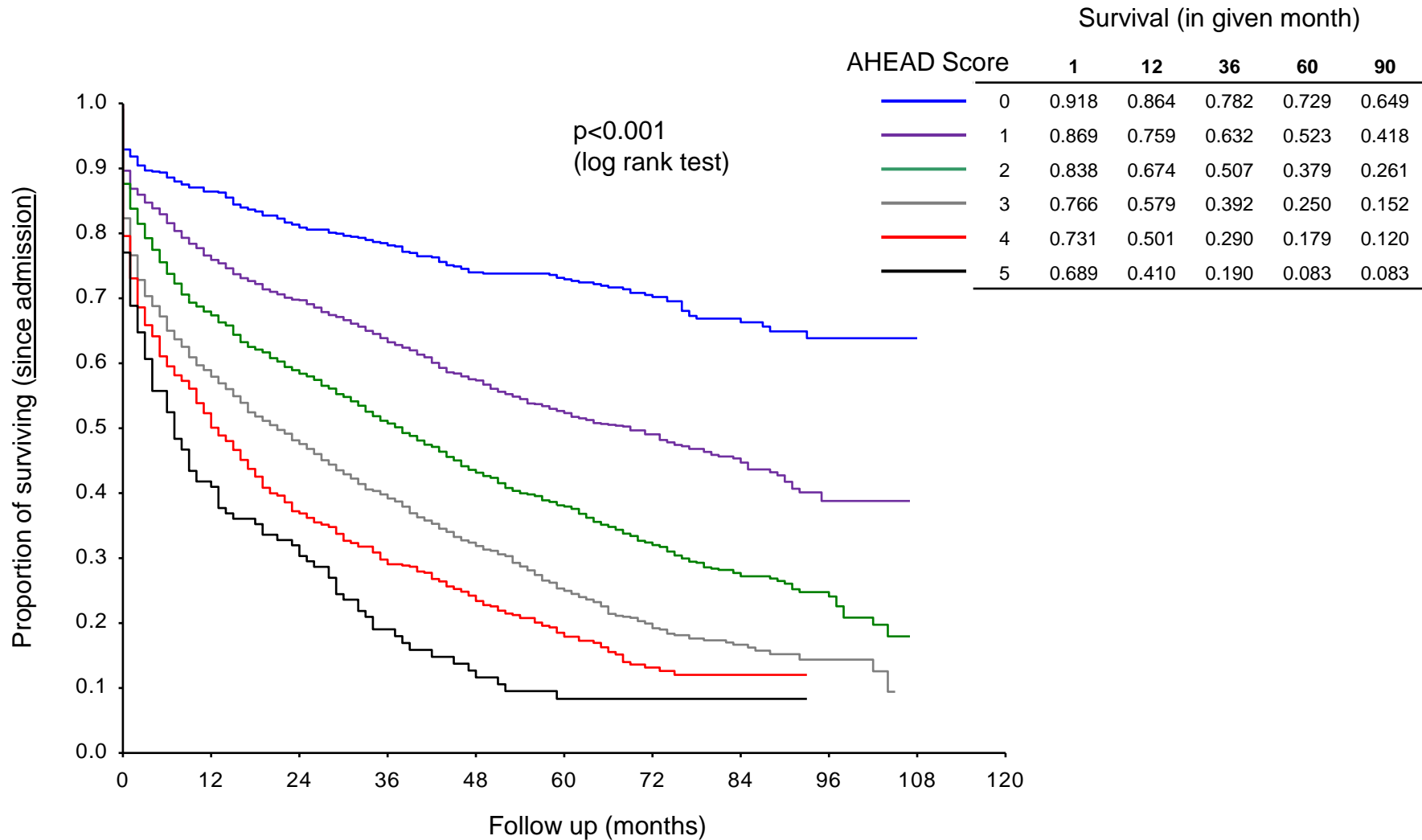
D = Diabetes mellitus

Ahead score is computed as a sum of points of risk factors in patients; its range is 0 (patients with no risk factors) to 5 points (all risk factors present).

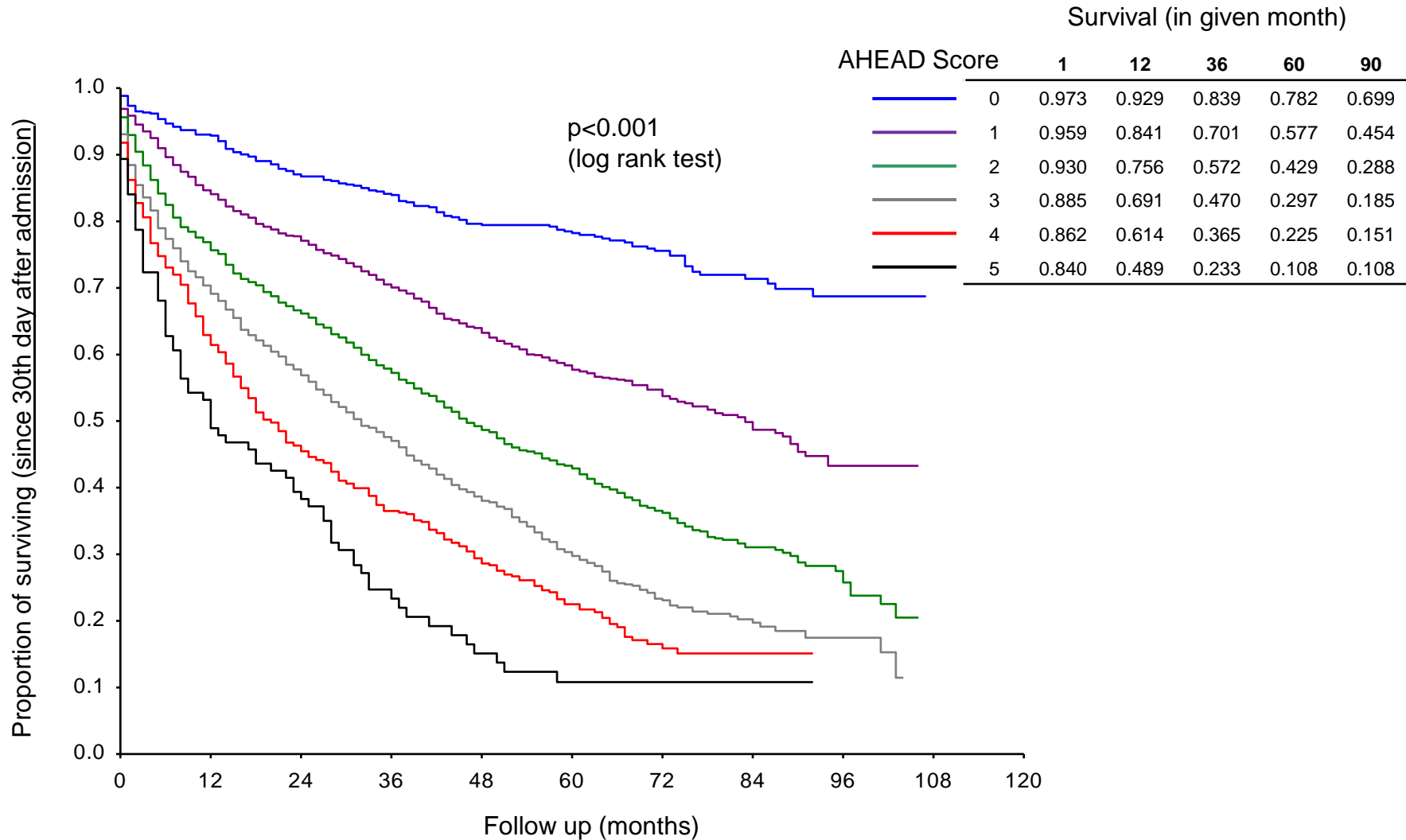
AHEAD score



Survival of patients according to AHEAD score I



Survival of patients according to AHEAD score II



AHEAD score VD studies
TRUE AHF - ularitide
RELAX - serelaxin

NO ACS

BPs > 110mmHg

N = 1648



AHEAD score AHEAD VD

The multivariate analysis identified 5 independent risk factors, for each risk factors the patient receive 1 point for the risk score computation:

A = Atrial fibrillation

H = Heamoglobin < 120 g/l (F)
130 g/l (M)

E = Elderly > 70 years

A = Abnormal renal functions
(creat > 130 umol/l)

D = Diabetes mellitus

Ahead score is computed as a sum of points of risk factors in patients; its range is 0 (patients with no risk factors) to 5 points (all risk factors present).

AHEAD score

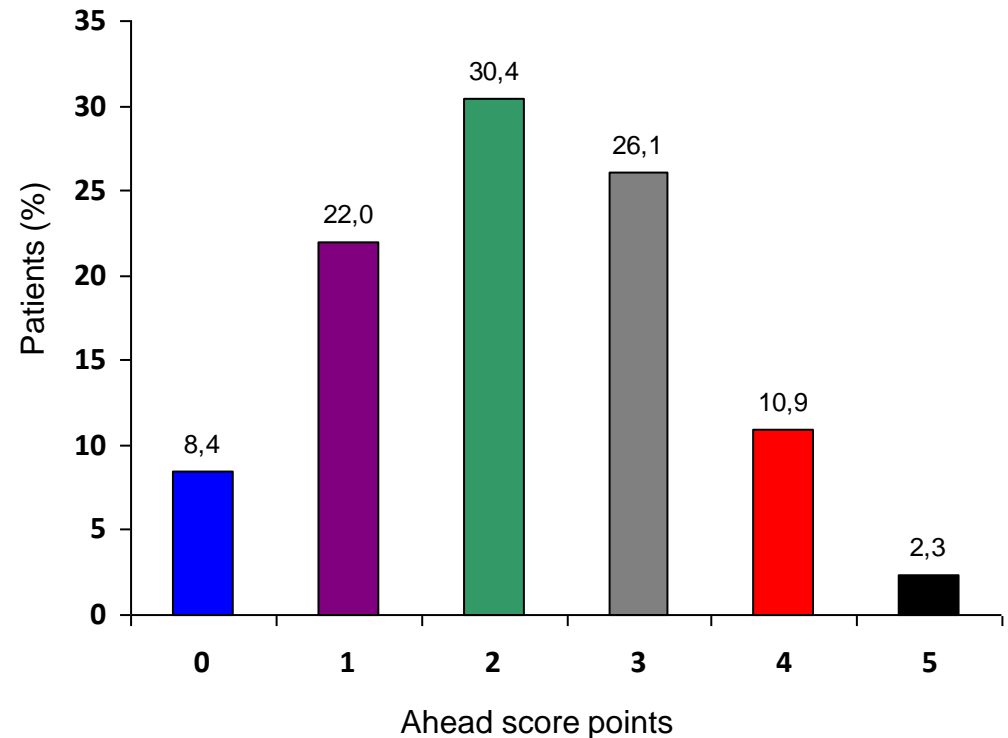


Figure 1 Analysis flowchart

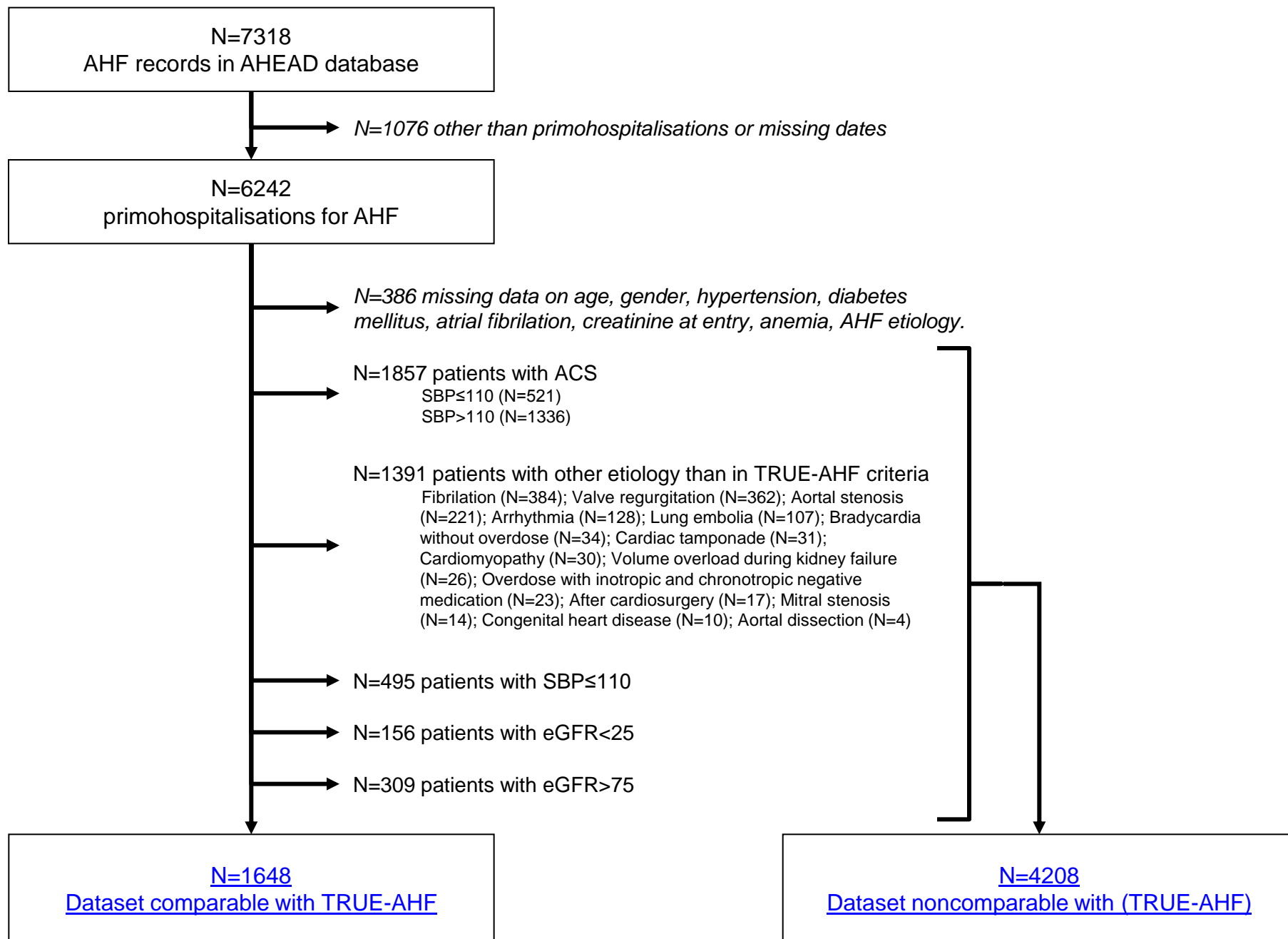


Figure 2 Survival since admission according to comparability with TRUE AHF study

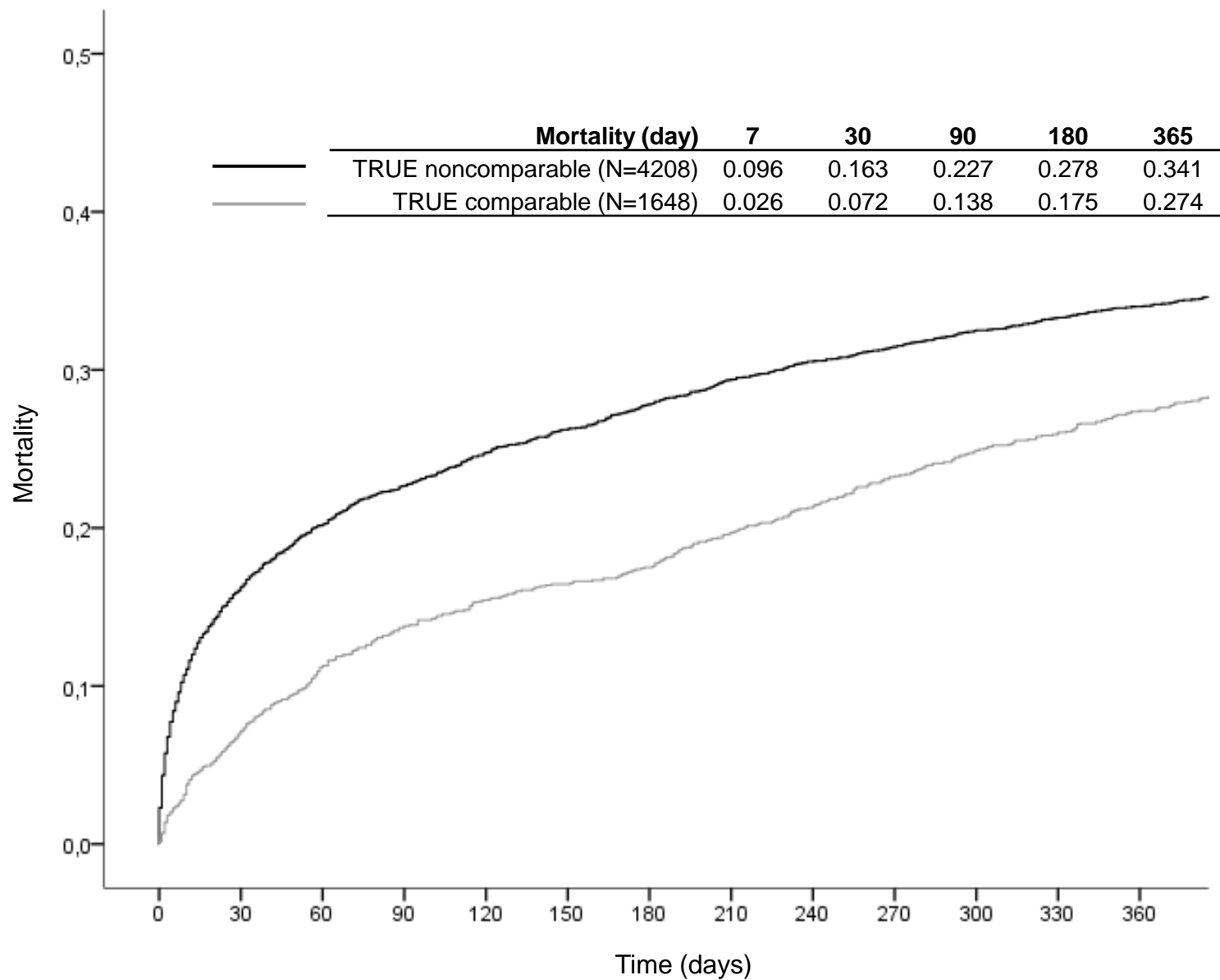


Figure 4 Systolic blood pressure as risk factor for in-hospital mortality

Dataset noncomparable with TRUE-AHF (N=4208)

Dataset comparable with TRUE-AHF (N=1648)

Systolic blood pressure

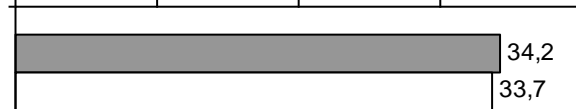
Inhospital mortality (%)

Inhospital mortality (%)

0 10 20 30 40

0 10 20 30 40

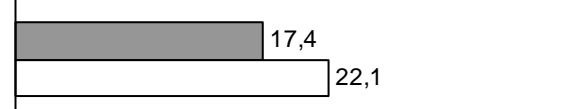
< 90 mm Hg



34,2

33,7

91-110 mm Hg



17,4

22,1

111-125 mm Hg



11,0

9,2

126-140 mm Hg



8,9

10,5

141-160 mm Hg



5,3

8,9

> 160 mm Hg



4,9

5,0

Men (N=893)

Women (N=755)

8,9

7,3

4,3

1,5

3,2

3,3

2,2

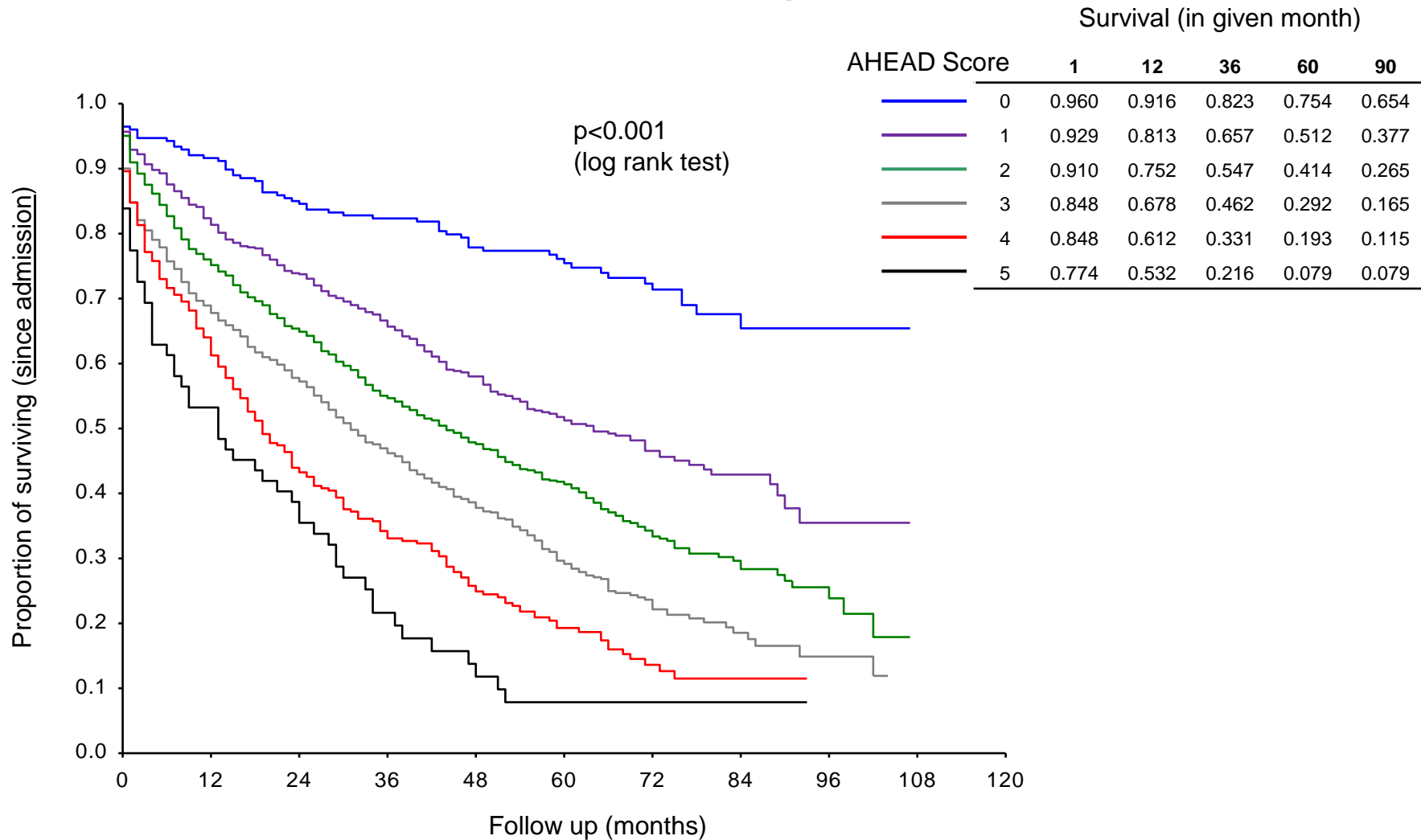
3,2

Men (N=2425)

Women (N=1783)

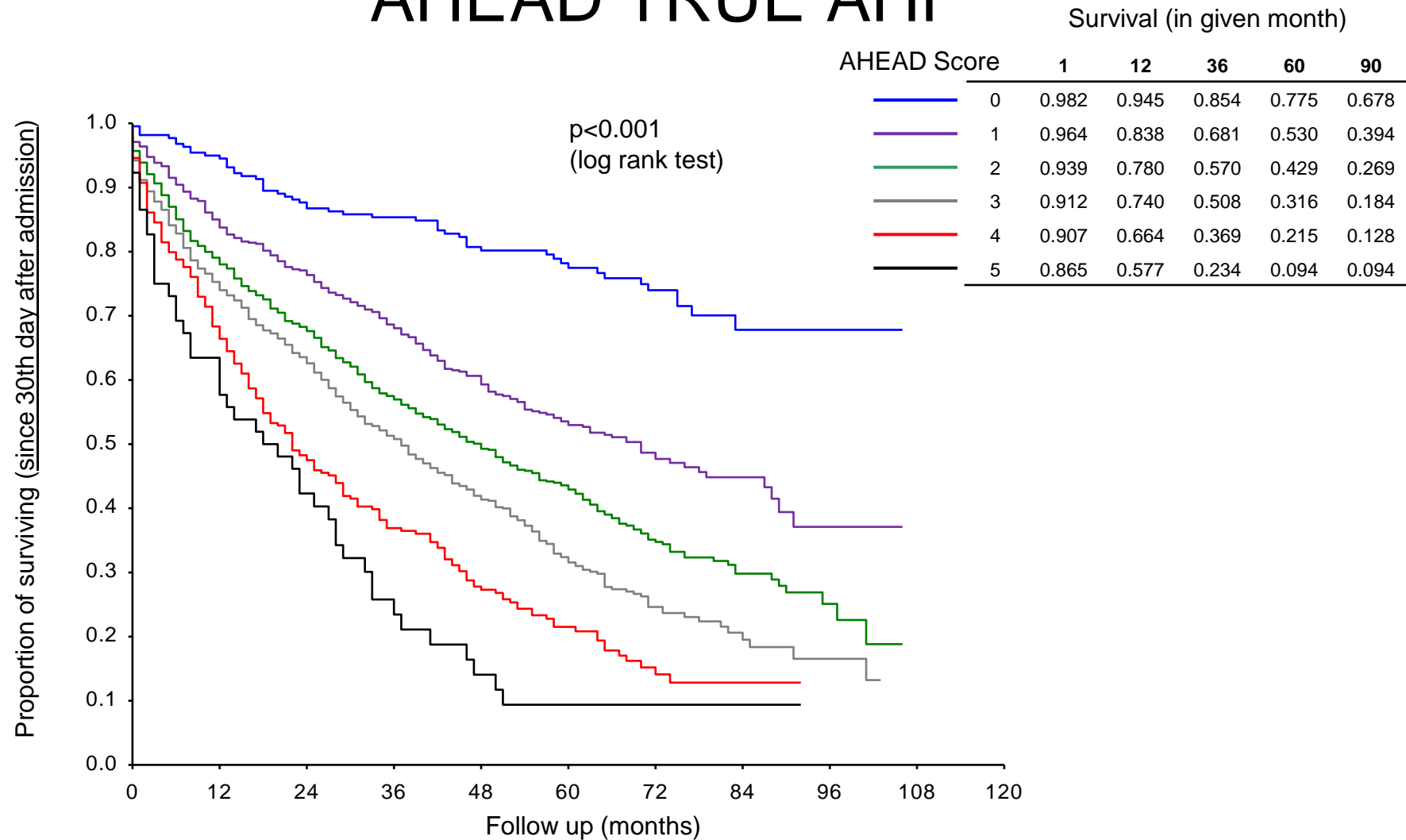
Survival of patients according to AHEAD score I

AHEAD TRUE AHF



Survival of patients according to AHEAD score II

AHEAD TRUE AHF



Děkuji za pozornost

