

Renal insufficiency increases mortality in acute coronary syndromes regardless of TIMI risk score

Dariusz Dudek, Bernadeta Chyrchel, Zbigniew Siudak, Rafał Depukat, Michał Chyrchel, Artur Dziewierz, Waldemar Mielecki, Tomasz Rakowski, Łukasz Rzeszutko, Jacek Dubiel

2nd Department of Cardiology, Institute of Cardiology, *Collegium Medicum*, Jagiellonian University, Krakow, Poland

Abstract

Background: Non ST-segment elevation acute coronary syndromes (NSTEMI ACS) are the most frequent cause of admission to intensive care units. Early risk assessment and implementation of optimal treatment are of special importance in these patients. Previous studies have demonstrated that renal insufficiency is an independent risk factor in patients with cardiovascular disease.

Aim: To assess the effects of renal function on the course of treatment and prognosis in patients with NSTEMI ACS admitted to hospitals without on-site invasive facilities but with a possibility of immediate transfer to a reference centre with a catheterisation laboratory.

Methods: Twenty-nine community hospitals without on-site invasive facilities participated in the Krakow Registry of Acute Coronary Syndromes – a prospective, multicentre, web-based, observational registry. Renal insufficiency (RI) was defined as creatinine clearance (CrCl) <60 ml/min.

Results: NSTEMI ACS was diagnosed in 1396 patients. Renal insufficiency was diagnosed in 34% of all patients. Only 17% of them had been diagnosed with RI prior to admission. Transfer for invasive treatment was undertaken in 10% of RI patients as compared to 16% of patients with CrCl >60 ml/min (NS). In-hospital mortality among patients remaining on conservative treatment in community hospitals was significantly higher among RI patients (4.0 vs. 0.6%; $p < 0.001$). Thienopyridines were less frequently used in RI patients (46 vs. 54%; $p < 0.05$). In-hospital mortality among RI patients remaining in community hospitals and treated conservatively was higher than among non-RI patients in each TIMI risk score group: 7.3 vs. 2.4% ($p < 0.05$) in the high risk group, 4.1 vs. 1.4% (NS) in the moderate and 3.6 vs. 0% ($p < 0.001$) in the low risk group. Multivariate logistic regression analysis identified reduced creatinine clearance and a history of heart failure as independent factors influencing mortality.

Conclusions: Renal insufficiency was present in one-third of NSTEMI ACS patients. Patients with renal insufficiency had worse clinical risk profile and received less aggressive treatment. Patients with NSTEMI ACS and renal insufficiency treated conservatively had higher in-hospital mortality. Renal insufficiency modifies mortality irrespective of the TIMI risk score. Creatinine clearance should be considered in modification of the TIMI risk score scale.

Key words: renal insufficiency, acute coronary syndromes, creatinine clearance, TIMI risk score

Kardiologia Polska 2008; 66: 28–34

Introduction

Non ST-segment elevation acute coronary syndromes (NSTEMI ACS) are the most frequent cause of admission to intensive care units [1, 2]. Early risk assessment, appropriate planning and implementation of optimal treatment are of particular importance in these patients [3].

Epidemiological studies have demonstrated that renal insufficiency (RI) is an independent risk factor in patients with cardiovascular disease, including coronary artery disease [4, 5]. Cardiovascular disease is also a major cause of death in patients with chronic kidney disease (CKD), including end-stage renal failure [6].

The purpose of the present study was to assess the effects of renal function on the course of treatment and prognosis in patients with NSTEMI ACS admitted to hospitals without on-site invasive facilities but with a possibility of immediate transfer to a reference centre with a catheterisation laboratory.

Methods

Krakow Registry of Acute Coronary Syndromes

The Krakow Registry of ACS was a prospective, multicentre, observational registry designed to examine current epidemiology, in-hospital management and

Address for correspondence:

Dariusz Dudek MD, PhD, Instytut Kardiologii, CM UJ, ul. Kopernika 17, 31-501 Kraków, tel.: +48 12 424 71 81, e-mail: mcdudek@cyf-kr.edu.pl

Received: 18 July 2007. **Accepted:** 17 October 2007.

outcome in patients with ACS in the Krakow Region of Poland with a population of 3.2 million. A total of 29 community hospitals without on-site invasive facilities participated in consecutive enrolment periods of the registry from 2002 until 2006. To minimise selection bias, all consecutive patients with a suspected diagnosis of ACS were included. During the index hospitalisation in the community hospital, data concerning baseline demographics, clinical characteristics, relevant laboratory results, pharmacotherapy and adverse cardiovascular outcomes were recorded on a standardised, electronic, web-based case report form (<https://www.cardio.pl/acs/index.php>). Standardised definitions were used for adverse events and final diagnosis. Data were collected in a central electronic database. The database was reviewed for completeness by an independent physician and site queries were generated if needed.

Assessment of renal function

Baseline creatinine clearance (CrCl) was calculated using the Cockcroft-Gault formula from serum creatinine [7]. Renal insufficiency was defined as CrCl <60 ml/min (renal insufficiency stage \geq III according to the National Kidney Foundation guidelines) [8].

Study groups

Patients were divided into two groups according to the level of CrCl: those with CrCl <60 ml/min and those with CrCl \geq 60 ml/min (patients with normal renal function; non-RI). Both groups were then subdivided according to the NSTEMI ACS TIMI risk score into low (0-2 points), moderate (3-4 points) and high risk (5-7 points) of death [3]. We also analysed demographic and clinical data, treatment modality and in-hospital mortality in patients treated conservatively.

Statistical analysis

Data were analysed according to the established standards of descriptive statistics. Categorical variables are given as percentages and were compared by a maximum likelihood (ML) χ^2 test. Continuous variables were assessed for normality and are reported as mean \pm standard deviation (SD). Continuous variables were compared by t-test or the two-tailed Mann-Whitney U-test as appropriate. In a subsequent step of the analysis, a multivariate logistic regression model was applied to search for independent predictors of in-hospital mortality (including baseline, clinical characteristics and applied pharmacotherapy). All independent variables considered potentially significant were initially included in the model followed by a stepwise deletion of the least significant variable down to a significance level of 0.05 or less. Statistical significance was defined as a p value <0.05.

The study was conducted in accordance with the Declaration of Helsinki and its later revisions.

Results

Of 2382 patients hospitalised for ACS and recorded in the registry database, 1396 were diagnosed with NSTEMI ACS. Baseline creatinine concentration was measured in 870 of these patients (62%) and further analyses were performed on that patient subset.

Patients with RI defined as CrCl <60 ml/min comprised 34% (n=295) of the NSTEMI ACS population. Only 17% of them had prior diagnosis of RI. There were significantly more women with reduced CrCl as well as elderly with history of prior angina, myocardial infarction, heart failure and brain stroke. Both groups were similar with respect to the prevalence of arterial hypertension, lipid disorders, percutaneous coronary intervention (PCI) and coronary artery bypass grafting surgery (CABG) in the past (Table I).

The study groups did not differ in the medications used: aspirin, low molecular and unfractionated heparins, GP IIb/IIIa inhibitors, ACE inhibitors and nitrates. Patients with reduced CrCl significantly less frequently received beta-blockers, statins or thienopyridines, whereas diuretics were more often administered (Table II).

Only 10% of RI patients were selected for urgent invasive diagnostic procedures and transferred to a PCI centre in comparison to 16% of those with preserved renal function (p=NS). The overall in-hospital mortality among patients remaining in the community hospitals for conservative treatment was 2%, and was significantly

Table I. Baseline characteristics and past medical history in NSTEMI ACS patients by presence or absence of impaired renal function (N=870)

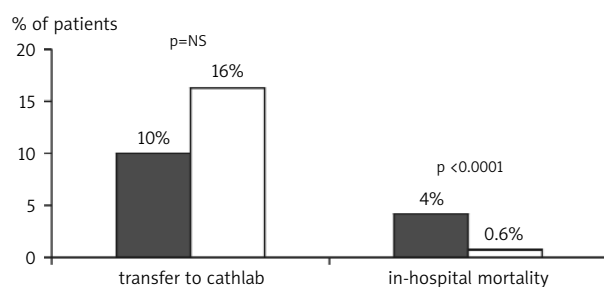
Parameter	Renal insufficiency (n=295)	Normal renal function (n=575)	p
Age [years]	69 \pm 12	63 \pm 10	<0.0001
Female gender [%]	55	35	<0.0001
Prior angina [%]	87	78	<0.0001
Previous myocardial infarction [%]	45	32	<0.0001
Heart failure [%]	43	15	<0.0001
Previous diagnosis of chronic kidney disease [%]	17	0.8	<0.0001
Arterial hypertension [%]	82	76	NS
Past stroke [%]	8	4	<0.05
Diabetes mellitus [%]	30	22	<0.05
Hyperlipidaemia [%]	49	49	NS
Current smokers [%]	18	31	<0.0001
Prior PCI [%]	6	7	NS
Prior CABG [%]	3.4	4	NS

Abbreviations: PCI – percutaneous coronary intervention, CABG – coronary artery bypass grafting surgery

Table II. In-hospital treatment in NSTEMI ACS patients stratified by renal function

Medication	Renal insufficiency (n=295)	Normal renal function (n=575)	p
Nitrates [%]	96.3	95	NS
Beta-blockers [%]	65	83	< 0.0001
Acetylsalicylic acid [%]	92.5	95	NS
Thienopyridines [%]	46	54	<0.05
Heparins (LMWH+UFH) [%]	85	85.6	NS
GP IIb/IIIa inhibitors [%]	0.3	0.5	NS
Diuretics [%]	54	31	< 0.0001
ACE inhibitors [%]	74	76	NS
Statins [%]	64	74	<0.05

Abbreviations: LMWH – low molecular weight heparin, UFH – unfractionated heparin

**Figure 1.** Transfer for invasive diagnosis and in-hospital mortality in entire registry population stratified by renal function; CrCl <60 ml/min – black bars, CrCl >60 ml/min – white bars**Table III.** Multivariate analysis for factors potentially influencing in-hospital mortality in NSTEMI ACS patients remaining for conservative treatment

Parameter	OR	95% CI	p
Creatinine clearance <60 ml/min	4.9949	1.1201-22.2739	0.035**
Male gender	0.6504	0.1931-2.1901	0.487
Age (per 1 additional year)	1.0060	0.9378-1.0781	0.861
Heart failure	3.0848	0.9039-10.5272	0.072*
Previous myocardial infarction	1.6063	0.4722-5.4645	0.448
Prior angina	0.9734	0.1038-9.1260	0.981
Current smoker	2.0627	0.6044-7.0393	0.248
Diabetes mellitus	1.4894	0.4555-4.8697	0.510
TIMI risk score (per additional point)	1.3540	0.8399-2.1829	0.213

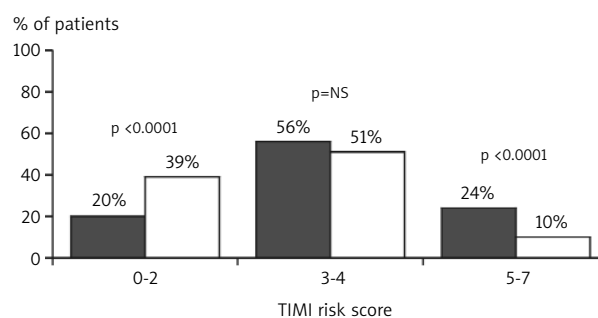
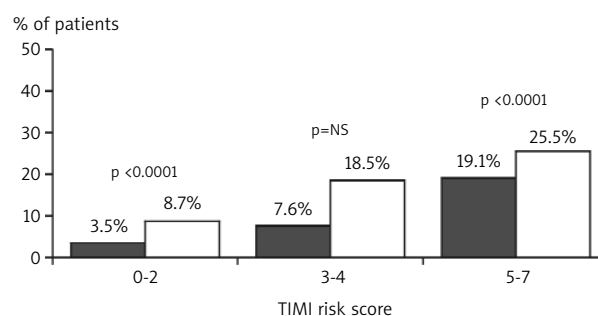
Abbreviations: OR – odds ratio, CI – confidence interval, * p < 0.1, ** p < 0.05

higher in RI patients than non-RI (4.0 vs. 0.6%, p < 0.0001) – Figure 1.

Multivariate logistic regression analysis identified reduced CrCl and a history of heart failure (borderline significance) as independent predictors influencing in-hospital mortality. Renal insufficiency was revealed to modify mortality irrespective of the TIMI risk score (Table III).

Patients with ACS were divided according to the TIMI risk score into low (TIMI risk score 0-2), moderate (3-4) and high risk (5-7) – Figure 2.

Patients with CrCl <60 ml/min in all TIMI risk score groups were less frequently transferred to the reference centre with cathlab facilities (Figure 3). In-hospital mortality among RI patients remaining in community hospitals and treated conservatively was higher in each TIMI risk score subgroup: 7.3 vs. 2.4% (p < 0.05) in the high risk group, 4.1 vs. 1.4% (p=NS) in the moderate and 3.6 vs. 0% (p < 0.001) in the low risk group (Figure 4).

**Figure 2.** TIMI risk score group assignment in renal insufficiency and normal renal function groups; CrCl <60 ml/min – black bars, CrCl >60 ml/min – white bars**Figure 3.** Transfer of NSTEMI ACS patients for invasive treatment to PCI centre in TIMI risk score subgroups; CrCl <60 ml/min – black bars, CrCl >60 ml/min – white bars

Discussion

The number of patients with RI is ever increasing [4]. In our Registry every third patient had renal function impairment. We also need to take into account that only one in seven of these patients had been diagnosed with RI prior to hospitalisation for NSTEMI ACS. Patients with RI were older, more often female and had more co-morbidities, thus confirming the findings of other investigators [9, 10]. This is probably associated with the fact that renal function and consequently CrCl declines with age [9]. The higher percentage of older women among those with reduced CrCl reflects the higher survival rate among women and consequently the higher proportion of women in the study group.

The optimal methods for the assessment of renal function in everyday clinical practice remain controversial. Glomerular filtration rate (GFR) is considered to be the best measure of kidney filtering capacity [10]. The level of GFR <60 ml/min is selected as the cut-off value for definition of RI because it represents a reduction by more than half of the normal value of ≈ 125 ml/min in young adults and this level is associated with the development of symptoms characteristic of RI [11, 12]. The National Kidney Foundation recommends estimating GFR from CrCl taking into account age, body weight and gender in order to avoid misclassification based on creatinine concentration alone [8].

Studies are continued to clarify the increased cardiovascular risk in patients with abnormal renal function. Decreased GFR may be associated with the prevalence of such traditional risk factors as arterial hypertension or dyslipidaemia, but also with other less understood factors such as hyperhomocysteinaemia. Decreased GFR is thought to be a marker of undiagnosed vascular disorders, especially in high-risk populations, and simultaneously an indicator of progressive remodelling and ventricular dysfunction [13]. Decreased GFR also strongly predisposes to the development of acute renal failure (ARF), which may per se increase mortality [12]. Diabetes mellitus (DM) is another important risk factor for the development of RI. In our study DM was significantly more frequent in patients with reduced CrCl.

What is even more pronounced is the fact that patients at higher risk are often treated less aggressively [14]. Although modern pharmacological and invasive treatment, so effective in the general population, may be safely and equally successfully used in patients with elevated risk, we may talk about therapeutic nihilism. The proportion of RI patients receiving appropriate pharmacological treatment that modifies the risk of death and referred for invasive therapy is lower than in the general population. Recent guidelines have mandated more aggressive lipid (statins) therapy, especially in patients with microalbuminuria, categorised as a high risk group.

The patients in the Krakow Registry of ACS with reduced CrCl were less often given beta blockers,

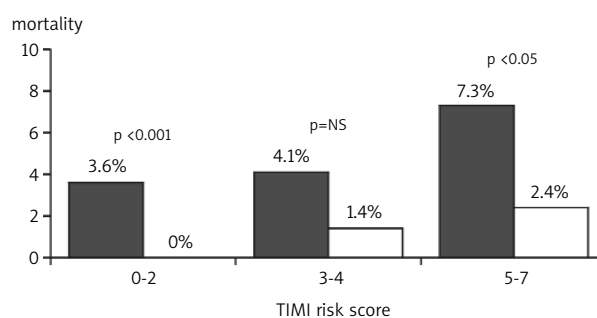


Figure 4. In-hospital mortality among NSTEMI ACS patients remaining for conservative treatment in TIMI risk score subgroups; CrCl <60 ml/min – black bars, CrCl >60 ml/min – white bars

thienopyridines and statins, but more frequently diuretics. This confirms the finding that patients with low level of GFR less often receive aggressive treatment with ACE inhibitors, beta-blockers, aspirin, platelet inhibitors or invasive procedures [13-16]. This trend is especially strong in the elderly over 65 years of age [17]. A similar tendency occurs in patients with end-stage renal disease, of whom less than half receive multi-drug treatment with ASA, beta-blockers, ACE inhibitors and statins [18]. It is important to modify this approach, especially for beta-blockers and statins, which have been found to have beneficial effects in patients with chronic renal disease [18, 19].

Patients with reduced CrCl are less often referred to invasive cardiology centres irrespective of TIMI risk score. The unwillingness to transfer patients for invasive treatment is probably a result of concern about increased risk of death and complications, including ARF associated both with invasive cardiology procedures (contrast medium usage) and CABG. However, it is high-risk patients that derive most benefit from aggressive invasive treatment [20-22].

Renal insufficiency in patients with NSTEMI ACS is an independent risk factor for mortality. In patients with reduced CrCl, mortality rate is significantly higher irrespective of stratification by TIMI risk score. It is of particular importance that in-hospital mortality in all TIMI risk score groups is significantly higher among RI patients, which further emphasises that renal function is an independent predictor in NSTEMI ACS. Therefore, CrCl should be considered in modification of the widely applied Antman's TIMI risk score scale [3]. The multivariate models confirm this association. In-hospital mortality in patients with ACS is higher if they have RI at presentation and depends on its severity; an increased mortality rate is also observed at long-term follow-up [11, 23].

The widely used TIMI risk score does not take into account reduced CrCl as a factor influencing final outcome in patients with NSTEMI ACS. There is probably an urgent need to establish a new scheme or modify the

existing one, paying special attention to RI in NSTEMI ACS patients when estimating the risk of death and other serious ischaemic complications, because current studies confirm the unfavourable effect of reduced CrCl on outcomes in these patients, leading to a paradoxical situation in which mortality for NSTEMI ACS through low, moderate and high risk TIMI score is similarly high.

Over recent years, equations estimating GFR on the basis of serum creatinine concentration and other obtainable patient data have been developed and validated. The widely used Cockcroft-Gault formula incorporates serum creatinine, gender, age and body weight [7]. However, weight measurements or estimations make calculation and reporting of Cockcroft-Gault results by laboratories problematic. In our analysis the body weight was precisely measured at admission to community hospitals, either by the nurse or by the doctor in the emergency room.

Study limitation

Use of the Cockcroft-Gault formula may be considered a limitation of the study due to the fact that new American Heart Association guidelines for detection of CKD in patients with or at increased risk of cardiovascular disease advise the use of a novel model – MDRD [24]. The Cockcroft-Gault formula was advised at the time of our registry and was the most popular method for the assessment of kidney function [8].

The major limitation in the interpretation of the results is the fact that it was a registry, with all the very well known disadvantages and drawbacks of registries. However, there seems to be positive acceptance of and a need for high quality registry results as they are certainly complimentary to the results of randomised clinical trials, and convey a message that is sometimes omitted in the latter. Nevertheless, we tried to avoid group selection potential bias by applying statistical tools like multivariate regression analysis.

Conclusions

Every third patient with NSTEMI ACS had RI. Only one in seven of these patients had prior diagnosis of CKD. Patients with NSTEMI ACS treated conservatively and RI had higher in-hospital mortality. In patients with NSTEMI ACS renal function provides valuable information about prognosis and may supplement the widely used TIMI risk score. The combination of the two may be a useful tool for genuine estimation of risk of death and complications in NSTEMI ACS.

References

1. Steg PG, Goldberg RJ, Gore JM, et al. GRACE Investigators. Baseline characteristics, management practices, and in-hospital outcomes of patients hospitalized with acute coronary syndromes in the Global Registry of Acute Coronary Events (GRACE). *Am J Cardiol* 2002; 90: 358-63.
2. Hasdai D, Behar S, Wallentin L, et al. A prospective survey of the characteristics, treatments and outcomes of patients with acute coronary syndromes in Europe and the Mediterranean basin; the Euro Heart Survey of Acute Coronary Syndromes (Euro Heart Survey ACS). *Eur Heart J* 2002; 23: 1190-201.
3. Antman EM, Cohen M, Bernink PJ, et al. The TIMI risk score for unstable angina/non-ST elevation MI: A method for prognostication and therapeutic decision making. *JAMA* 2000; 284: 835-42.
4. Foley RN, Parfrey PS, Sarnak MJ. Clinical epidemiology of cardiovascular disease in chronic renal disease. *Am J Kidney Dis* 1998; 32 (Suppl 3): S112-9.
5. Manjunath G, Tighiouart H, Ibrahim H, et al. Level of kidney function as a risk factor for atherosclerotic cardiovascular outcomes in the community. *J Am Coll Cardiol* 2003; 41: 47-55.
6. Herzog CA, Ma JZ, Collins AJ. Poor long-term survival after acute myocardial infarction among patients on long-term dialysis. *N Engl J Med* 1998; 339: 799-805.
7. Cockcroft DW, Gault MH. Prediction of creatinine clearance from serum creatinine. *Nephron* 1976; 16: 31-41.
8. National Kidney Foundation. K/DOQI clinical practice guidelines for chronic kidney disease: evaluation, classification and stratification. *Am J Kidney Dis* 2002; 39 (Suppl 1): S1-S266.
9. Al Suwaidi J, Reddan DN, Williams K, et al. for the GUSTO-IIb, GUSTO-III, PURSUIT, and PARAGON-A Investigators. Prognostic implications of abnormalities in renal function in patients with acute coronary syndromes. *Circulation* 2002; 106: 974-80.
10. Levey AS, Bosch JP, Lewis JB, et al. A more accurate method to estimate glomerular filtration rate from serum creatinine: a new prediction equation. Modification of Diet in Renal Disease Study Group. *Ann Intern Med* 1999; 130: 461-70.
11. Masoudi FA, Plomondon ME, Magid DJ, et al. Renal insufficiency and mortality from acute coronary syndromes. *Am Heart J* 2004; 147: 623-9.
12. McCullough PA, Nowak RM, Foreback C, et al. Emergency evaluation of chest pain in patients with advanced kidney disease. *Arch Intern Med* 2002; 162: 2464-8.
13. Hillege HL, van Gilst WH, van Veldhuisen DJ, et al. Accelerated decline and prognostic impact of renal function after myocardial infarction and the benefits of ACE inhibition: the CATS randomized trial. *Eur Heart J* 2003; 24: 412-20.
14. McCullough PA, Sandberg KR, Borzak S, et al. Benefits of aspirin and beta-blockade after myocardial infarction in patients with chronic kidney disease. *Am Heart J* 2002; 144: 226-32.
15. Freeman RV, Mehta RH, Al Badr W, et al. Influence of concurrent renal dysfunction on outcomes of patients with acute coronary syndromes and implications of the use of glycoprotein IIb/IIIa inhibitors. *J Am Coll Cardiol* 2003; 41: 718-24.
16. Reddan DN, O'Shea JC, Sarembock IJ, et al. Treatment effects of eptifibatid in planned coronary stent implantation in patients with chronic kidney disease (ESPRIT Trial). *Am J Cardiol* 2003; 91: 17-21.
17. Shlipak MG, Heidenreich PA, Noguchi H, et al. Association of renal insufficiency with treatment and outcomes after myocardial infarction in elderly patients. *Ann Intern Med* 2002; 137: 555-62.
18. Cice G, Ferrara L, D'Andrea A, et al. Carvedilol increases two-year survival in dialysis patients with dilated cardiomyopathy: a prospective, placebo-controlled trial. *J Am Coll Cardiol* 2003; 41: 1438-44.

19. Saltissi D, Morgan C, Rigby RJ, et al. Safety and efficacy of simvastatin in hypercholesterolemic patients undergoing chronic renal dialysis. *Am J Kidney Dis* 2002; 39: 283-90.
20. Szczech LA, Best PJ, Crowley E, et al. BARI Investigators. Outcomes of patients with chronic renal insufficiency in the bypass angioplasty revascularization investigation. *Circulation* 2002; 105: 2253-8.
21. Le Feuvre C, Borentain M, Beygui F, et al. Comparison of short- and long-term outcomes of coronary angioplasty in patients with and without diabetes mellitus and with and without hemodialysis. *Am J Cardiol* 2003; 92: 721-5.
22. Dacey LJ, Liu JY, Braxton JH, et al. Long-term survival of dialysis patients after coronary bypass grafting. *Ann Thorac Surg* 2002; 74: 458-62.
23. Beattie JN, Soman SS, Sandberg KR, et al. Determinants of mortality after myocardial infarction in patients with advanced renal dysfunction. *Am J Kidney Dis* 2001; 37: 1191-200. Erratum in: *Am J Kidney Dis* 2001; 38: 701.
24. Brosius FC 3rd, Hostetter TH, Kelepouris E, et al. Detection of chronic kidney disease in patients with or at increased risk of cardiovascular disease: a science advisory from the American Heart Association Kidney And Cardiovascular Disease Council; the Councils on High Blood Pressure Research, Cardiovascular Disease in the Young, and Epidemiology and Prevention; and the Quality of Care and Outcomes Research Interdisciplinary Working Group: developed in collaboration with the National Kidney Foundation. *Circulation* 2006; 114: 1083-7.

Niewydolność nerek zwiększa śmiertelność w ostrych zespołach wieńcowych niezależnie od skali ryzyka TIMI

Dariusz Dudek, Bernadeta Chyrchel, Zbigniew Siudak, Rafał Depukat, Michał Chyrchel, Artur Dziewierz, Waldemar Mielecki, Tomasz Rakowski, Łukasz Rzeszutko, Jacek Dubiel

II Klinika Kardiologii, Instytut Kardiologii, Collegium Medicum, Uniwersytet Jagielloński, Kraków

Streszczenie

Wstęp: Ostre zespoły wieńcowe bez uniesienia odcinka ST (NSTEMI ACS) są najczęstszą przyczyną hospitalizacji na oddziałach intensywnej opieki medycznej i kardiologicznej. Wczesna ocena ryzyka u chorych z NSTEMI ACS i włączenie właściwej, zgodnej z wytycznymi strategii leczniczej ma szczególne znaczenie dla rokowania tych chorych. Poprzednie badania wykazały, że niewydolność nerek jest niezależnym czynnikiem ryzyka u osób z chorobą wieńcową.

Cel: Oszacowanie wpływu funkcji nerek na przebieg leczenia i rokowanie chorych z NSTEMI ACS przyjętych do szpitala bez możliwości wykonania natychmiastowych zabiegów inwazyjnych, ale z możliwością szybkiego przetransportowania do szpitala z własną pracownią hemodynamiki.

Metodyka: W Małopolskim Rejestrze Ostrego Zespołu Wieńcowego – prospektywnym, wieloośrodkowym rejestrze obserwacyjnym, wzięło udział 29 szpitali powiatowych bez własnej pracowni hemodynamiki pracującej w trybie 24-godzinnym. Niewydolność nerek (RI) zdefiniowano jako klirens kreatyniny (CrCl) <60 ml/min.

Wyniki: U 1396 chorych potwierdzono NSTEMI ACS. Niewydolność nerek została potwierdzona u 34% chorych. Tylko 17% z nich miało zdiagnozowaną RI przed przyjęciem. Diagnostyce i leczeniu inwazyjnemu poddano 10% chorych z RI i 16% chorych z prawidłową funkcją nerek (p=NS). Śmiertelność wewnątrzszpitalna wśród chorych leczonych zachowawczo była istotnie wyższa w grupie chorych z RI w porównaniu z chorymi bez RI (4,0 vs 0,6%; p <0,001). Była ona wyższa w każdej z podgrup skali ryzyka TIMI – 7,3 vs 2,4% (p <0,05) w podgrupie wysokiego ryzyka, 4,1 vs 1,4% (p=NS) w podgrupie umiarkowanego ryzyka i 3,6 vs 0% (p <0,001) w podgrupie niskiego ryzyka. Tienopirydyny były rzadziej stosowane u chorych z RI (46 vs 54%; p <0,05). Analiza wieloczynnikowa pozwoliła wyodrębnić czynniki niezależnie wpływające na śmiertelność wewnątrzszpitalną, do których należą niewydolność krążenia w wywiadzie i obniżony klirens kreatyniny (<60 ml/min).

Wnioski: Niewydolność nerek zaobserwowano u co trzeciego chorego z NSTEMI ACS. Pacjenci z niewydolnością nerek mieli częściej choroby współistniejące oraz rzadziej otrzymywali zalecane leczenie farmakologiczne. U chorych z NSTEMI ACS i RI stwierdzono wyższą śmiertelność wewnątrzszpitalną. Niewydolność nerek wpływa na śmiertelność niezależnie od powszechnie stosowanej skali ryzyka TIMI. Być może należałoby uwzględnić klirens kreatyniny w modyfikacji skali ryzyka TIMI.

Słowa kluczowe: niewydolność nerek, ostre zespoły wieńcowe, klirens kreatyniny, skala ryzyka TIMI

Kardiologia Pol 2008; 66: 28–34

Adres do korespondencji:

dr hab. n. med. Dariusz Dudek, Instytut Kardiologii, CM UJ, ul. Kopernika 17, 31-501 Kraków, tel.: +48 12 424 71 81, e-mail: mcdudek@cyf-kr.edu.pl

Praca wpłynęła: 18.07.2007. Zaakceptowana do druku: 17.10.2007.