Mortality among older patients admitted to the medical wards of Groote Schuur Hospital, Cape Town, South Africa, 2010 - 2013

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Background. Geriatric medicine is an evolving specialty in Africa, and little is known about mortality among older patients admitted to medical wards.

Objectives. To determine mortality rates and associated factors among older medical inpatients.

Methods. Electronic data on patients aged ≥ 60 years admitted to the medical wards of Groote Schuur Hospital, Cape Town, South Africa, between January 2010 and December 2013 were analysed. Data extracted included sex, age, causes of death, and length of stay from date of admission to discharge or death. Results of laboratory tests carried out during the admission were also obtained.

Results. In all, 11 254 older patients were admitted (mean (standard deviation) age 70.7 (7.9) years). There were 1 701 deaths (15.1%). The unadjusted mortality rate was 29.6 deaths per 1 000 patient-days (PD). The majority (87.5%) were admitted as emergency cases. Mortality in the first 24 hours was 32.4 deaths per 1 000 PD. There was a significant increase in mortality with increasing age (p<0.001). Stroke was the commonest cause of mortality (14.5%). The predictors of mortality were short length of stay on admission (odds ratio (OR) 1.047, 95% confidence interval (CI) 1.033 - 1.061), high white blood cell count (OR 1.064, 95% CI 1.054 - 1.074), low platelet count (OR 0.999, 95% CI 0.996 - 1.000), low haemoglobin (OR 0.940, 95% CI 0.917 - 0.964) and high blood urea (OR 1.042, 95% CI 1.033 - 1.051).

Conclusions. Mortality among older medical inpatients was high. Modifiable predictors of mortality, especially related to laboratory derangements, should be identified and addressed promptly.

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Older persons constitute a unique subgroup of adults in any population, not only due to advanced age but also a variety of physiological, social and demographic changes.^[1] With increasing age, a number of physiological variables decline substantially, compromising the individual's ability to respond to pathological insults.^[2] The determinants of mortality in older persons are numerous and can be grouped as non-modifiable or modifiable factors.^[5]

Globally, hospital admission and mortality rates among older persons are continually on the increase.^[4] This is partly due to the increase in the number of older persons in the population. Hospitalisation for medical illnesses is considered to be a risk factor for death among older persons.^[5] Furthermore, such illness usually has adverse effects on health and may lead to negative outcomes such as disability, loss of independence and autonomy, and social isolation. In addition, older patients have increased susceptibility to iatrogenic conditions and nosocomial as well as other infections.^[3] Research has shown that older patients accounted for 11.0 - 56.6% of all adult hospital medical ward admissions,^[5] with a mortality rate of 26.6 - 57.6%.^[5] In Africa, however, the mortality rate is expected to be higher owing to a lack of standard medical facilities.

Globally, older persons are the fastest-expanding age group, with approximately one million people attaining the age of 60 years every month.^[3] It is estimated that by the year 2025, eight of the 11 countries with the biggest populations of older persons will be developing countries.^[3] The 2014 report on the profile of persons

aged ≥ 60 years in South Africa (SA) showed a population of 4.1 million, representing 8.0% of the total population.^[6,7] This makes SA the African country with the second-largest proportion of older persons (after Mauritius).^[6,7] Furthermore, the number of persons aged ≥ 60 years in SA is expected to increase over 100-fold from 2.47 million in 1985 to 5.23 million by the year 2025.^[6] The projected increase is huge compared with the estimated growth of 74 - 83% among young adults.^[6,7]

Objectives

Old age is associated with both health and functional decline and the associated increased prevalence of chronic diseases with increased morbidity and hospitalisation. There is, however, a paucity of information on the mortality rate among hospitalised older patients in medical wards in Africa. Most of the existing information on older patients has been extracted or extrapolated from data on general adult patients admitted to medical wards in Africa,^[5] which revealed higher levels of hospitalisation and mortality among older persons compared with the young population.^[5,6] This study therefore sought to determine mortality rates, temporal trends and associated factors among older patients admitted in a large tertiary hospital in SA.

Methods

This was a retrospective study of all medical admissions of patients aged ≥ 60 years between 1 January 2010 and 31 December 2013 at Groote Schuur Hospital (GSH), Cape Town, SA. GSH became

operational in 1938 and functions as a tertiary and quaternary care hospital. The division of geriatric medicine is one of 18 clinical divisions in the Department of Medicine. Although the hospital has a division of geriatric medicine, there are no beds dedicated to geriatric medicine for acute admissions. Older patients requiring acute medical management are admitted to general medical wards together with other adult medical patients.

Electronic records of all older patients were obtained via the Clinicom Patient Administration System with approval from the chief operations officer of GSH and the Human Research Ethics Committee of the Faculty of Health Sciences at University of Cape Town (ref. no. 012/2015). All older patients who were admitted to the medical wards during the study period were included. We obtained information on their sex, age, length of stay, causes of death (as reported on the basis of clinical impression), and dates of admission and discharge. Laboratory data, which included complete blood counts and urea, creatinine and glucose levels, were obtained from the National Health Laboratory Service (NHLS) data warehouse. These are the investigations most routinely done on acute admission to the hospital. Admission details and laboratory data were linked by means of the patient's unique hospital number. Confidentiality of data was maintained with the exclusion of names and hospital numbers from the analysis, leaving only the serial numbers allotted to the patients.

Data were analysed using the Statistical Package for the Social Sciences version 22 (IBM, USA). Descriptive and inferential statistics were used. Appropriate charts were used to illustrate categorical variables. Unadjusted mortality rates and rate ratios were calculated in patient-days (PD) with 95% confidence intervals (CIs). The Cochrane-Armitage test for temporal trend was used to test for linearity of the mortality trend over the study period. Logistic regression analysis was used to explore factors that were significantly associated with mortality, and the level of significance was set at p<0.05.

Results

A total of 11 254 older patients were admitted to the medical wards during the study period (males n=5 089, females n=6 165). The mean (standard deviation (SD)) age of the patients was 70.7 (8.0) years (females v. males 71.7 (8.3) years v. 69.5 (7.4) years) (t=14.580; *p*<0.001). The total number of deaths was 1 701 (15.1%), with 802 (15.8%) and 899 (14.6%) deaths occurring among males and females, respectively. The mean (SD) age of males who died was significantly higher than the mean age of males who were discharged, at 70.3 (7.6) years v. 69.4 (7.4) years (t=3.20; p=0.012). A similar trend was observed among females, with ages 72.5 (8.5) years v. 71.5 (8.2) years (t=4.35; p<0.0001). The majority (87.5%) of the patients were admitted as emergency cases. Mortality in the first 24 hours of admission was 32.4 deaths (95% CI 29.2 - 35.9) per 1 000 PD, which was not significantly higher among the males at 35.4 deaths (95% CI 30.5 - 40.8) per 1 000 PD compared with the females at 30.0 deaths (95% CI 25.9 - 34.6) per 1 000 PD (*p*=0.11).

Overall, the unadjusted mortality rate was 29.6 (95% CI 28.2 - 31.0) per 1 000 PD. The mortality rate was highest in 2010 (31.8 (95% CI 28.9 - 34.8) per 1 000 PD) and decreased steadily over the years to 27.0 (95% CI 24.4 - 29.7) per 1 000 PD in 2013. Table 1 shows the unadjusted mortality rates from January 2010 to December 2013 by gender and age groups. Overall, the mortality rate was higher among men compared with women, without a statistical difference (30.3 per 1 000 PD v. 29.0 per 1 000 PD; p=0.40). However, the only significant difference between mortality rate and gender was observed in the year 2010.

			M	Males				Fe	Females			
			Age at									
			death		Unadjusted			Age at death				
			(years),		mortality rate, per			(years),		Unadjusted mortality rate, Mortality rate ratio,	Mortality rate ratio,	
	Deaths, n	PD, n	mean (SD)	Patients, n	1 000 PD (95% CI)	Deaths, n PD, n	PD, n	mean (SD)	Patients, n	Patients, <i>n</i> per 1 000 PD (95% CI)	female/male (95% CI)	<i>p</i> -value
Year of admission												
2010	231	6515	69.8 (7.4)	1 336	35.5 (31.0 - 40.3)	235	8 161	71.5 (8.0)	1 687	28.8 (25.2 - 32.7)	1.23 (1.02 - 1.48)	0.03*
2011	183	6 191	69.2 (7.4)	1 230	29.6 (25.4 - 34.2)	203	7 216	71.8 (8.3)	1 460	28.1 (24.4 - 32.3)	1.05 (0.86 - 1.29)	0.66
2012	201	6 299	69.4 (7.3)	1 237	31.9 (27.7 - 36.6)	240	7 966	71.7 (8.4)	1 518	30.1 (26.4 - 34.2)	1.06 (0.87 - 1.28)	0.58
2013	187	7 490	69.6 (7.7)	1 286	25.0 (21.5 - 28.8)	221	7 645	71.7 (8.5)	1500	28.9 (25.2 - 33.0)	0.86 (0.71 - 1.05)	0.15
Age group (years)												
60 - 64	226	8 657	61.9 (1.5)	1 593	26.1 (22.8 - 29.7)	187	8 469	62.0(1.4)	1 483	22.1 (19.0 - 25.5)	1.18 (0.97 - 1.44)	0.10
65 - 69	194	7 018	66.8(1.4)	1 292	27.6 (23.9 - 31.8)	178	7 213	67.0(1.4)	1 304	24.7 (21.2 - 28.6)	1.12 (0.91 - 1.38)	0.30
70 - 74	152	4 495	71.9 (1.4)	934	33.8 (28.7 - 39.6)	171	5 981	72.0 (1.4)	1 198	28.6 (24.5 - 33.2)	1.18 (0.94 - 1.48)	0.15
75 - 79	119	$3\ 410$	76.8 (1.6)	682	34.9 (28.9 - 41.8)	165	4 653	76.9 (1.4)	1 051	35.5 (30.3 - 41.3)	0.98 (0.77 - 1.25)	0.94
80 - 84	77	1 902	81.7 (1.3)	385	40.5 (31.9 - 50.6)	100	2 669	81.6 (1.4)	640	37.5 (30.5 - 45.6)	1.08 (0.79 - 1.47)	0.66
≥85	34	$1\ 013$	88.1 (2.1)	203	33.6 (23.2 - 46.9)	98	2 003	88.7 (3.3)	489	48.9 (39.7 - 59.6)	0.69 (0.45 - 1.02)	0.07
Total	802	26 495	69.5 (7.4)	5 089	30.3 (28.2 - 32.4)	899	30 988	71.7 (8.3)	6 165	29.0 (27.2 - 31.0)	1.04 (0.95 - 1.149)	0.40
PD = patient-days, SD = standard deviation; CI = confidence interval *Significant at 5% level.	standard deviati	on; CI = confic	dence interval.									

Table 1. Unadjusted mortality rates from January 2010 to December 2013 by gender and age groups

Fig. 1 describes the overall unadjusted mortality rates by age group. There was a significant increase in the mortality rate with age from 24.1 per 1 000 PD in the 60 - 64-year age group to 43.8 per 1 000 PD in patients aged \geq 85 years (Cochrane-Armitage test for linear trend: χ^2 =64.57; *p*<0.0001).

(14.5%), heart failure (7.9%) and myocardial infarction (7.2%). Fig. 2 describes the classification of the diagnoses and proportion of deaths according to the *International Classification of Primary Care*, 2nd edition (ICPC-2).^[8] The highest proportion of deaths were related to the general body system (22.2%), while no death was related to the psychological and eye systems.

Diagnoses were obtained in 6 143 (54.6%) of the patients. As shown in Table 2, the commonest reasons for admission were myocardial infarction (19.6%), heart failure (7.4%) and stroke (7.3%). However, the highest proportions of deaths were found in patients with stroke

Table 3 shows the laboratory values of the patients by mortality. Mean white blood cell counts and serum creatinine, blood urea and blood glucose levels were significantly higher among patients who

Table 2. Morbidity pattern by proportion of	deaths		
Morbidities	Died (<i>N</i> =455), <i>n</i> (%)	Discharged (N=5 688), n (%)	Total (N=6 143), n (%)
Myocardial infarction	33 (7.2)	1 168 (20.5)	1 201 (19.6)
Heart failure	36 (7.9)	416 (7.3)	452 (7.4)
Stroke	66 (14.5)	385 (6.8)	451 (7.3)
Chronic obstructive pulmonary disease	22 (4.8)	335 (5.9)	357 (5.8)
Hypertension	15 (3.3)	243 (4.3)	258 (4.2)
Pneumonia	31 (6.8)	222 (3.9)	253 (4.1)
Ischaemic heart disease	13 (2.9)	172 (3.0)	185 (3.0)
Atrioventricular block	7 (1.5)	175 (3.1)	182 (3.0)
Paroxysmal tachycardia	10 (2.2)	169 (3.0)	179 (2.9)
Delirium	7 (1.5)	162 (2.8)	169 (2.8)
Disorders of digestive system (unspecified)	15 (3.3)	145 (2.5)	160 (2.6)
Mental disorder (unspecified)	0	153 (2.7)	153 (2.5)
Cardiac disease (unspecified)	6 (1.3)	142 (2.5)	148 (2.4)
Dementia	14 (3.1)	127 (2.2)	141 (2.3)
Malignancy with metastasis	31 (6.8)	97 (1.7)	128 (2.1)
Diabetes mellitus	4 (0.9)	99 (1.7)	103 (1.7)
Chronic kidney disease	13 (2.9)	74 (1.3)	87 (1.4)
Dermatitis	4 (0.9)	75 (1.3)	79 (1.3)
Bronchial asthma	10 (2.2)	62 (1.1)	72 (1.2)
Pulmonary tuberculosis	10 (2.2)	52 (0.9)	62 (1.0)
Acute renal failure	16 (3.5)	45 (0.8)	61 (1.0)
Acute upper respiratory tract infection	2 (0.4)	55 (1.0)	57 (0.9)
Epilepsy	1 (0.2)	52 (0.9)	53 (0.9)
Peptic ulcer disease	2 (0.4)	50 (0.9)	52 (0.8)
Gastroenteritis	3 (0.6)	46 (0.8)	49 (0.8)
Disorders of urinary system (unspecified)	4 (0.9)	45 (0.8)	49 (0.8)
Septicaemia	14 (3.1)	34 (0.6)	48 (0.8)
Rheumatic heart disease	0	45 (0.8)	45 (0.7)
Anaemia	1 (0.2)	42 (0.7)	43 (0.7)
Non-rheumatic heart disease	1 (0.2)	39 (0.7)	40 (0.7)
Pulmonary heart disease	0	38 (0.7)	38 (0.6)
Atrial fibrillation	1 (0.2)	36 (0.6)	37 (0.6)
Cellulitis	2 (0.4)	32 (0.6)	34 (0.6)
Chronic liver disease	4 (0.9)	30 (0.5)	34 (0.6)
Psoriasis	0	28 (0.5)	28 (0.5)
Cardiomyopathy	1 (0.2)	25 (0.4)	26 (0.4)
Phlebitis	0	20 (0.4)	20 (0.3)
Systemic poisoning	2 (0.4)	16 (0.3)	18 (0.3)
Psychosis	0	18 (0.3)	18 (0.3)
Transient ischaemic attack	1 (0.2)	17 (0.3)	18 (0.3)
Neuropathies	0	16 (0.3)	16 (0.3)
Multiple myeloma	2 (0.4)	13 (0.2)	15 (0.2)
Arterial diseases	0	13 (0.2)	13 (0.2)
Metabolic imbalance	0	13 (0.2)	13 (0.2)
Parkinson's disease	0	13 (0.2)	13 (0.2)
Ulcerative colitis	1 (0.2)	12 (0.2)	13 (0.2)
Others	50 (11.0)	422 (7.4)	472 (7.9)

died during the hospital admission compared with patients who were discharged. The mean haemoglobin concentration was significantly lower in patients who died compared with patients who were discharged. Although notably lower platelet counts were observed among patients who died than in those who survived, this finding was not statistically significant.

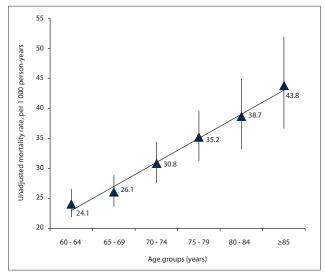


Fig. 1. Unadjusted mortality rate by age groups (Cochrane-Armitage test for linear trend: $\chi^2 = 64.57$; p<0.0001).

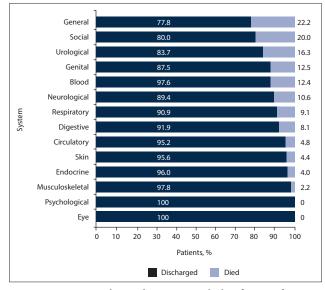


Fig. 2. Diagnoses according to the International Classification of Primary Care, *2nd edition*,^[8] *by proportion of deaths.*

Table 4 presents a logistic regression analysis of the factors associated with mortality. The logistic model was found to be statistically significant ($\chi^2(8)=642.994$; p<0.0001). The model explained 13.5% (Nagelkerke R^2) of the variance in mortality. It also correctly classified 84.2% of cases. The most significant factors associated with mortality were short length of stay on admission (odds ratio (OR) 1.047, 95% CI 1.033 - 1.061; p<0.0001), high white blood cell count (OR 1.064, 95% CI 1.054 - 1.074; p<0.0001), low platelet count (OR 0.999, 95% CI 0.996 - 1.000; p=0.006), low haemoglobin concentration (OR 0.940, 95% CI 0.917 - 0.964; p<0.0001) and high blood urea level (OR 1.042, 95% CI 1.033 - 1.051; p<0.0001).

Discussion

Factors associated with increased mortality in older patients admitted to medical wards were male sex (on bivariate analysis), high white blood cell count, low haemoglobin concentration, high urea level and a short hospital stay. This study was conducted in a large tertiary hospital in SA and deaths were reported in almost one-sixth (15.1%) of older patients. This proportion of deaths was similar to that reported in a systematic review of in-hospital mortality among elderly patients in medical wards in Africa^[5] and South America.^[9] However, mortality was higher than that reported from Europe (8.4%).^[10] Available mortality data on hospitalised older patients from Nigeria (26.7 - 31.7%)^[11,12] showed a higher proportion of deaths than the findings of the present study.^[5] A previous retrospective study (2002 - 2009) carried out in the same setting as our study found a proportion of deaths of 12 - 17%, although this population included young adults.^[13]

The proportion of deaths in the present study was higher among males than among females, though without statistical significance. This finding was not unexpected, as male sex has been reported to be an independent risk factor associated with inpatient mortality in medical wards in other studies.^[5,13] Other factors that work singly or jointly have been adduced for the mortality difference between male and female older patients.^[14] Older females have longer life expectancy, better health-seeking behaviour, better lifestyle habits and less exposure to stress. In addition, cardiocirculatory protection of females due to the effect of hormones during their fertile years is a major contribution.^[14]

Advancing age has been reported as an immutable risk factor for mortality in hospitalised older patients, and our observation was no different.^[5,13,15] A significant linear increase in mortality rate with increasing age was observed in both sexes. Also, there were some differences across the years of study. A possible explanation for the reduction in mortality between 2010 and 2013 may be the increase in the number of medical beds, which provided space for improved patient assessment. Medical beds were increased from 142 in 2010 to 153 in 2013. Similarly, the medical teams (called firms) were reorganised, resulting in improved efficiency. The medical firms were each provided with a dedicated ward in 2013, whereas in

	n	Dead, mean (SD)	Discharged, mean (SD)	t	p
White blood cell count (× $10^{9}/L$)	9 522	15.0 (0.1)	11.2 (0.1)	12.713	< 0.0001*
Platelet count (× 10 ⁹ /L)	8 758	311.2 (0.2)	312.0 (0.1)	-0.019	0.985
Haemoglobin (g/dL)	9 559	11.5 (2.9)	12.3 (2.6)	-9.194	< 0.0001*
Serum creatinine (mmol/L)	10 519	219.8 (26.4)	134.1 (15.3)	12.569	< 0.0001*
Urea (mmol/L)	9 382	17.6 (1.6)	10.7 (0.9)	15.804	< 0.0001*
Blood glucose (mmol/L)	365	11.2 (0.9)	8.2 (0.7)	2.077	0.042*
SD = standard deviation. *Significant at 5% level.					

				95% CI for OR	
riables	Beta coefficient	<i>p</i> -value	OR	Lower	Upper
e	0.004	0.318	1.004	0.996	1.012
ngth of stay	0.046	< 0.0001*	1.047	1.033	1.061
hite cell count	0.001	< 0.0001*	1.064	1.054	1.074
itelet count	-0.001	0.006*	0.999	0.996	1.000
emoglobin	-0.062	< 0.0001*	0.940	0.917	0.964
eatinine	0.002	0.366	1.000	0.999	1.000
ea	0.041	< 0.0001*	1.042	1.033	1.051

Table 4. Logistic regression analysis of the factors associated with mortality

previous years, medical wards were shared as an open space by the different medical firms. Having space dedicated to each team may have impacted positively on care of older patients in the medical wards by providing focused patient care and improved handovers. The improvement in mortality occurred despite improved access to tertiary care through the expanded primary healthcare platform. This platform started in 1994 and has improved access to care for older patients with advanced morbidities.^[13]

The commonest reasons for hospitalisation in this study were myocardial infarction, heart failure and stroke. Acute illnesses such as stroke and acute exacerbation of respiratory disease were common causes of mortality, as has been found in other studies in Africa.^[5,16,17] Older people are facing the double burden of communicable and non-communicable diseases as part of the health transition in Africa.^[18] Acute exacerbation of chronic diseases, infections and deranged metabolic factors are therefore common reasons for hospitalisation and mortality.^[5,16] Not surprisingly, the leading causes of death in this study were septicaemia, acute renal failure and malignancies with metastasis.^[5,16,17] These were also reflected in the predictors of mortality found, as the risk of mortality increased by 6.4% for every 1×10^{9} /L increase in the white blood cell count. Anaemia from causes including malignancies with metastasis was a predictor, as there was a 6.0% increase in mortality with every 1 g/dL decrease in the haemoglobin concentration. Serum creatinine was not found to be a predictor of mortality, although there was a significant association between mortality and serum creatinine level on bivariate analysis. However, blood urea was a predictor, with a 4.2% increase in mortality with every 1 mmol/L increase in the urea level. Serum creatinine may not have been an independent predictor of mortality, as creatinine is a poor screening test for renal failure in older patients, who commonly have reduced muscle mass.^[19] Increased length of stay has been associated with increased mortality in a number of studies, particularly in patients with delirium.^[9] In our study, reduced length of stay was associated with increased mortality. This is an indication of the high mortality rate (32.4 deaths per 1 000 PD) in the first 24 hours of admission. The high rate is probably due to patients presenting at an advanced stage of the illness, and an inadequate service provided at the primary level of care.[5,9]

Study strengths and limitations

The present study is of significance in that it is among the few studies in Africa to evaluate mortality and associated factors in the older population in Africa. It has limitations in that it was retrospective and a diagnosis was available in the dataset for just over half of the patients. Prehospital functional assessment and nutritional status, which have an impact on mortality in older people, could not be obtained. The study was limited to one centre and is therefore not generalisable to other settings in SA.

Conclusions

There was a high mortality rate among older patients admitted to the medical wards of a large tertiary hospital in SA, with a male preponderance. The high proportion of patients presenting on an emergency basis calls for effective and more frequent outpatient appointments. This extra effort will facilitate early diagnosis and management of acute exacerbations and complications of morbidities. Lastly, the association of mortality with laboratory biomarkers shows the need for adequate investigation and management of morbidities with which these biomarkers are associated.

Declaration. None.

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Conflicts of interest. None.

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