

Risk Factors Related to Hospital Mortality in Kenyan Patients with Traumatic Intracranial Haematomas

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Background: The aim of this study was to investigate the factors which influence mortality of patients with traumatic intracranial hematomas (ICH).

Methods: Following ethical approval, the records of patients admitted to the Kenyatta National Hospital neurosurgical unit between January 2000 and December 2009 and who were diagnosed with traumatic ICH were retrieved and reviewed. The outcome measured was mortality during admission. Analysis was done using Statistical Package for Social Sciences (SPSS) version 12.

Results: During the study period, 608 patients were admitted with a diagnosis of traumatic intracranial haematoma. 89.3% were male and 10.7% female with a mean age was 35.33 years (+/- 17.420). 59.3% of patients with preoperative GCS scores of <8 died, while only 11% and 3% deaths occurred in the patients with moderate and mild head injury respectively. 70.4% of patients with unreactive pupils died as compared to 7.5% of patients with bilaterally symmetrical reactive pupils ($p=0.002$). There was a statistically significant increase in mortality in patients who did not undergo surgical intervention (26.1%; $p=0.000$) as compared to those who were operated (15.7%; $p=0.000$). The mean time from accident to surgery was 3 days. Patients who were operated on more than 4 days after the initial trauma had a mortality of 42.1% as compared to 9.3% for patients operated on within 24 hours.

Conclusion: An increased risk of death was observed in patients who are over 61 years of age, have lower preoperative GCS, the presence of pupillary abnormalities and a long interval between trauma and decompression.

Introduction

Traumatic intracranial haematomas (ICH) are some of the most common traumatic neurosurgical emergencies which often require surgical intervention. It is estimated that intracranial haematomas occur in 25–45% of severe traumatic brain injuries, 3–12% of moderate cases, and approximately 1 in 500 patients with mild head injury^{1, 2, 3}. Although there have been drastic developments and improvements in neurotraumatology, emergency medical service systems, neuro-intensive monitoring and treatment, ICH is still a disorder with a very high mortality rate and extremely poor prognosis among traumatic brain injuries^{4, 5, 6}.

Therefore, identifying reliable prognostic factors for traumatic ICH to improve the surgical results in these patients is important. However relatively few studies have focused on the factors that affect the outcome of patients with surgically treated traumatic ICH. In addition, to the authors' knowledge there has been no other study that addressed Intracranial Haematomas among Kenyan patients. Kenyatta National Hospital is a national teaching and referral hospital with a bed capacity of 2000 and the largest neurosurgical unit in the Great Lakes region. Through this center, the great majority of neurosurgical referrals are managed. We reviewed records of patients who were diagnosed with traumatic ICH to elucidate which factors are related to mortality of this lethal disorder with a view to improve functional outcome.

Patients and Methods

Following ethical approval, the records of 608 patients who were admitted to the Kenyatta National Hospital and were diagnosed with traumatic ICH between January 2000 and December 2009 were reviewed. We categorized all variables which might have been related to the functional recovery and mortality into three groups : 1) clinical variables; gender, age, mechanism of injury, preoperative GCS scores, preoperative pupillary abnormalities 2) computerized tomography (CT) variables; location of hematoma, midline shift 3) surgical variables; type of surgery, time elapsed from accident to surgery. The GCS score is usually determined on admission and all patients were divided patients into three

groups; those with GCS scores of 3 to 8, 9 to 13 and 13 to 15 for statistical analysis. For logistic regression analysis, patients were classified as having 0, 1, or 2 reactive pupils.

Data was collected in pre-formed questionnaires, coded and analysis carried out using Statistical Package for Social Sciences (SPSS) version 11.5. Frequencies and means were computed for description of the various variables, discrete variables compared using the Chi-square test and continuous variables compared using the Students' t-test. Logistic and univariate linear regression model were run to determine which variables are independently associated with functional recovery and mortality. A p value <0.05 was considered statistically significant.

Results

During the study period, 608 patients were admitted at Kenyatta National Hospital with a diagnosis of traumatic intracranial haematoma. Five hundred and forty three patients (89.3%) were male and 65 (10.7%) female. The mean age was 35.33 years (± 17.420) with a range from 6 months to 96 years. Majority of the patients (49%) were aged between 26 and 45 years while 5.6% and 9.4% were aged below 13 years and older than 61 years respectively. The most common cause of injury was assault (48%) with road traffic and falls accounting for 28% and 24% respectively.

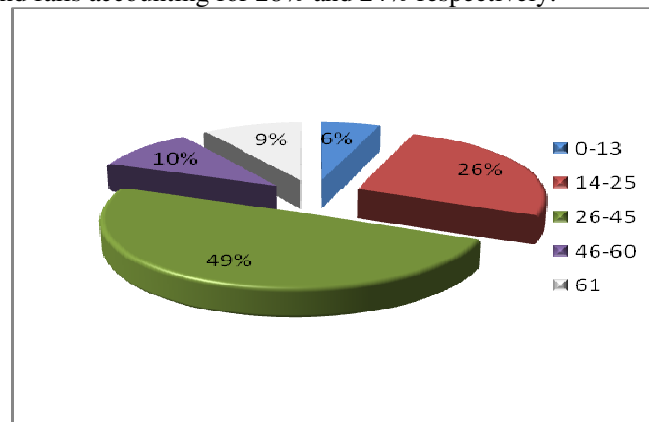


Figure 1. The Age Distribution of Patients with Intracranial Haematomas

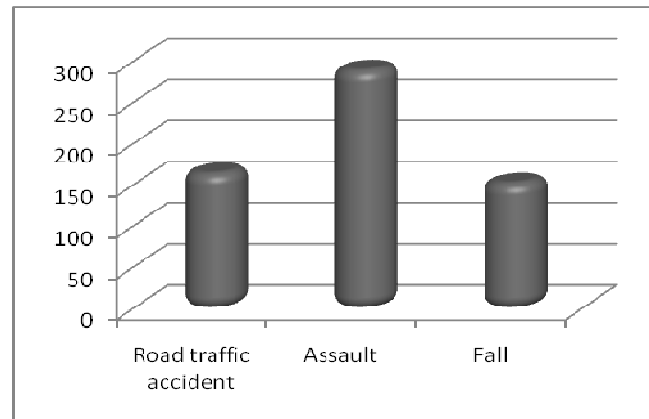


Figure 2. Aetiology of intracranial hematoma

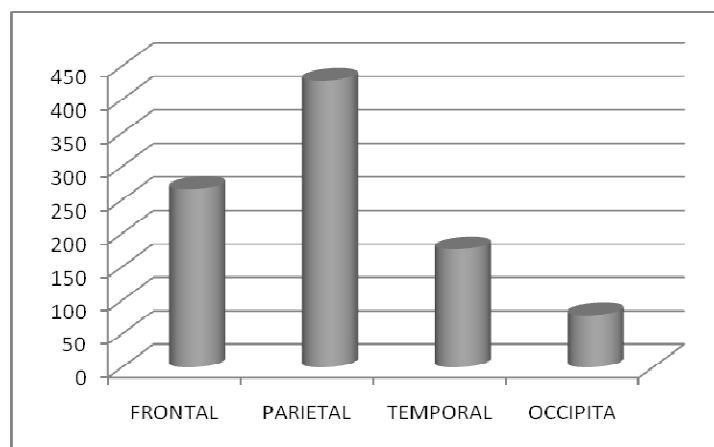


Figure 3. A representation of the anatomical locations of post-traumatic hematomas

A total of 290 patients (47.7%) presented with an extradural haematoma and 259 (42.6%) with a subdural haematoma while 24 (3.9%) patients presented with both haematomas. Other post-traumatic lesions included intracerebral hemorrhage (126; 20.7%), subarachnoid haematomas (21; 3.5%), contusions (49; 8.1) and skull fractures (169; 27.8%). The parietal lobe was the most common location of intracranial hematomas accounting for 68% of the cases while the occipital lobe was the least common with 12.8%.

Of the 608 patients in our series, 112 patients died while hospitalized and the overall mortality rate was 18.4%. The mean age of these patients was 38.37 years $\pm$ 16.518 with a range from 2 to 96 years. Eleven contributory variables to death were analyzed separately in univariate analyses. Consequently, eight factors were found to be significantly related to death (Table 1). A significantly higher mortality rate of 23.3% and 26.3 % was observed amongst patients aged between 46 to 60 years and older than 61 years respectively ($p=0.025$). It was also observed that the mortality rate increased in direct proportion to an increase in age.

With regards to sex, one hundred and six (94.6%) of the patients who passed away during admission were men while 6 (5.4%) were women ($p=0.026$). In addition, there was a higher mortality for patients with subarachnoid hemorrhage (43%) with the lowest mortality being observed among patients with extradural hematoma (15%). The admission GCS score was highly correlated with hospital mortality ($p=0.000$). Of the 148 patients with preoperative GCS scores of 8 or less, 83 (59.3%) died. By contrast, only 22 (11%) and 7 (3%) deaths occurred in the patients with scores ranging from 9 to 12 and greater than 13 respectively. Details of the various components of the Glasgow Coma score were available for 492 patients. The mortality rate for patients with a motor score of 6 was 4.8 % as compared to 72.7% amongst patients with a motor score of 1 ($p=0.000$). In addition, higher vocal and eye responses were associated with statistically significant lower mortality.

Details of pupillary reaction to light were documented for 589 patients as follows: 466 patients (79.1%) had symmetrical reactive pupils, 69 (11.7%) had anisocoria but reactive pupils and 54 (9.2%) had homolateral unreactive mydriasis. When pupillary characteristics were cross tabulated with outcome, 70.4% of patients with unreactive pupils died as compared to 7.5% of patients with bilaterally symmetrical reactive pupils ($p=0.002$). Further, of the 406 patients with a history of loss of consciousness, 24.9% died during admission as compared to 5.4% mortality in patients with no history of loss of consciousness ($p=0.000$).

A history of convulsions was associated with a higher mortality 26.6% as compared to 17.3% of patients who did not have such a history ($p=0.157$).

Table 1. Factors Significantly Related to Death

Clinical variables	No. of patients	Mortality (%)	<i>p</i> value
Total	608	112 (18.4%)	
Sex Distribution			0.026
Male	543	106 (19.5%)	
Female	65	6 (9.2%)	
Age Distribution (Years)			0.025
<13	34	4 (5.9%)	
14-25	157	17 (10.2%)	
26-45	298	63 (21.1%)	
46-60	60	14 (23.3%)	
>61	57	15 (26.3%)	
Mechanism of Injury			0.018
Assault	173	15 (8.7%)	
Motor vehicle collision	94	19 (20.2%)	
Fall	87	3 (3.4%)	
GCS			0.000
>13	235	7 (3%)	
9-12	200	22 (11%)	
<8	148	83 (59.3%)	
Pupillary Abnormalities			0.002
Reactive to light	466	35 (7.5%)	
Anisocore but reactive	69	38 (55.1%)	
Areactive to light	54	38 (70.4%)	
History of loss of Consciousness			0.000
Yes	406	101 (24.9%)	
No	202	11 (5.4%)	
History of Convulsions			0.157
Yes	64	17 (26.6%)	
No	538	93 (17.3%)	
Mode of Referral (N=566)			0.612
Directly from accident scene	272	47 (17.3%)	
From another health facility	294	53 (18.0%)	
Surgery Done			0.000
Yes	447	70 (15.7%)	
No	161	42 (26.1%)	
Type of Surgery			0.095
Craniotomy and evacuation	337	62 (18.4%)	
Burr hole evacuation	83	5 (6%)	
Craniotomy and elevation of depressed skull fracture	23	0	
Elevation of depressed skull fracture only	4	1	
Time Interval from Trauma to Surgery			0.087
<24 hours	227	21 (9.3%)	
1 to 2 days	101	23 (22.8%)	
3 to 4 days	223	44 (19.7%)	
>4 days	57	24 (42.1%)	
ICU Stay			0.000
Yes	116	56 (48.3%)	
No	492	16 (3.3%)	

Regarding the relationship between the haematoma location and mortality, our data showed a higher mortality rate for patients with posterior fossa haematomas (34.04%), although the statistical analysis indicated that the difference was not significant ($p=0.072$). In addition, there were 447 patients who underwent surgical evacuation of hematomas. There was a statistically significant increase in mortality in patients who did not undergo surgical intervention (26.1%; $p=0.000$) as compared to those who were operated (15.7%; $p=0.00$). The mean time elapsing from accident to surgery was 3 days. Patients who were operated on more than 4 days after the initial trauma had a mortality of 42.1% as compared to 9.3% for patients operated on within 24 hours. When the entire study population was subjected to logistic regression analysis, sex, age, pupillary reactivity, admission GCS scores and mechanism of injury were found to be significantly independently predictive of functional recovery. Among these patients, type and timing of operative intervention did not significantly affect outcome.

Table 2. Results of logistic regression on factors affecting mortality among patients with intracranial haematomas

Explanatory variables*	Exp (B)† (95% CI)	P value
Age	-0.12 (-0.18 to -0.06)	0.035
Sex	-0.149 (-4.88 to 0.190)	0.854
Mechanism of injury	0.313 (0.169 to 0.457)	0.086
Glasgow coma score	0.879 (0.741 to 1.017)	0.000
Surgery done	-0.570 (-0.812 to -0.327)	0.000
Loss of consciousness	0.221 (-0.114 to 0.557)	0.009
Pupillary reactivity	-1.179 (-1352 to -1.006)	0.000

*Table 2 Variables are listed in order they entered forward stepwise selection process. Age was entered as continuous variable; Glasgow coma scale group as 1 (Glasgow coma score 3-8), 2 (Glasgow coma score 9-12) and 3 (Glasgow coma score 13-15); systolic blood pressure as 1 (<90 mm Hg) and 2 (>90 mm Hg);

†Factor by which odds change when independent variable increases by one unit; values <0 indicate that odds increase as explanatory variable increases and values >0 indicate that odds decrease as explanatory variable increases.

Discussion

In spite of more rapid diagnosis and aggressive neurosurgical intervention, the mortality rate of traumatic intracranial hematomas is still high in majority of series ranging between 39%^{7,8} and 75%^{6,9}. In our series, the observed overall mortality was 18.4%. Although this figure may appear low, it must be interpreted in consideration of the admission GCS. Majority of the patients in this series 435 (74.6%) had an admission GCS above 9 and only 148 (25.4%) had severe head injury. Mwang'ombe and Kiboi¹⁰ reported an overall mortality of 56.2% amongst patients with severe head injury in a Kenyan population which is in accord with our observed 59.3% mortality of those patients in this series with severe head injury.

It has been established in literature that increasing age is associated with a higher mortality from traumatic brain injury. Howard *et al*¹¹ compared 33 young patients (aged 18-40 years) with old patients (aged over 65) and they reported significantly higher mortality rate in the older group (74% versus 18%). Mosenthal *et al*¹² observed that the mortality from isolated traumatic brain injury for the geriatric population was twice that of younger patients. Munro *et al*¹³ also found that patients aged 65 years and older had lower survival rates than patients less than 65 years old. Similar findings have been reported by other authors^{5,14,15}. In the study by Wilberger *et al*^{16,17}, the mean age of survivors was 41 years and of non survivors was 59 years. We observed a similar trend and found that age was an independent predictor of outcome in traumatic ICH. In our study, those patients aged younger than

13 years had a mortality rate of 5.9%, whereas patients above 61 years had a mortality rate of 26.3%. In addition, those patients older than 45 years showed significantly higher rate (OR=0.12, $p=0.035$) of mortality by multivariate logistic regression analysis. The mechanism by which age has such an effect on outcome is unknown, but suggestions include a poor regenerative capacity of the older brain and predisposition to develop a more lethal injury¹¹. Some of this increased mortality in the elderly may be explained by the intrinsic properties of the ageing brain, pre-existing co-morbidities and complications. Furthermore, the adverse effects of general anaesthesia and surgery may affect the respiratory and circulatory function of the elderly, increasing the severity of brain injury. Therefore, in addition to treating pre-existing diseases to decrease the risk of complications, improved long-term care should be emphasized for elderly surgical patients.

The preoperative GCS score was another important predictor of outcome. Many authors reported that there is highly significant correlation between outcome and GCS score at admission^{14, 18, 19, 20}. Of the 148 patients with preoperative GCS scores of 8 or less, 83 (59.3%) died. By contrast, only 22 (11%) and 7 (3%) deaths occurred in the patients with scores ranging from 9 to 12 and greater than 13 respectively. These findings confirm recent studies indicating that the severity of injury determines the outcome²¹. Further, Mwang'ombe and Kiboi¹⁰ also found that outcome in head injury patients was directly related to admission GCS.

Pupillary abnormalities are associated with a significantly worse outcome. In our series, patients who had bilateral areactive pupils had a mortality of 70.4% as compared to 55.1% and 7.5% for one reactive pupil and bilaterally symmetrical pupils respectively. Many authors reported that patients with bilateral fixed pupils at surgery had a mortality rate from 64 to 93%^{4, 5, 6, 16, 22}. Kyu-Hong *et al*²¹ reported that patients with one non-reacting pupil, had a mortality from 48 to 68%. This is in accordance with the findings of our series and is confirmed by other reports^{18, 19, 23}. In addition on logistic regression, pupillary abnormalities were strong predictors for mortality of patients with traumatic intracranial hematomas (OR= -1.179, $p= 0.000$). It has been postulated that pupillary dilatation is associated with decreased brainstem blood flow and that ischaemia rather than mechanical compression of the third cranial nerve is an important causal factor²⁴.

The time from the trauma until surgical decompression also affects the mortality. Some researchers have observed that the sooner surgery is performed in cases of acute head trauma, the better the final results are^{25, 26}. Seelig *et al*²⁷ in their study concluded that a delay from injury to operation was the factor of greatest therapeutic importance in traumatic ASDH. But the relationship between time to surgery and outcome is still controversial. Haselsberger *et al*¹⁹ reported that 47% died and 32% had a favorable outcome among the patients operated within two hours after the onset of coma. On the other hand, Stone *et al*²³ reported no difference in patients operated within 4 hours of injury compared with those operated later. In our series, the mean duration before surgical evaluation was more than most reports but with a low mortality rate.

In our study, only 9.3% of the patients undergoing surgery within 24 hours of the injury died, while those operated on over 4 days after injury had a 42.1% mortality rate. However in our series, due to the retrospective nature of the study, we were unable to analyse the length of the period of herniation or duration of operation as independent predictors.

Conclusion

We suggest that the patients in whom surgery is indicated, especially those with herniation, undergo surgery as soon as possible after trauma. Moreover, a rational surgical approach, effective removal of the haematoma and the avoidance of complications are key factors to reduce the postoperative mortality.

References

1. Servadei, F. Prognostic factors in severely head injured adult patients with acute subdural hematomas. *Acta Neurochir (Wien)*. 1997;139:279-285.
2. McKhann, GM, Copass, MK, Winn, RH. Prehospital care of the head-injured patient. In: *Neurotrauma*. New York: McGraw-Hill; 1996:103-118.
3. Meyer, SH, Tandall, MD, Chesnut, MD. Posttraumatic extra-axial mass lesions: subdural and extradural hematoma. In: *The practice of neurosurgery*. Vol. 2. Baltimore (MD): Williams and Wilkins; 1996:1443-1460.
4. Hatashita, S, Koga, N, Hosaka, Y, Takagi, S. Acute subdural hematoma: severity of injury, surgical intervention, and mortality. *Neurol Med Chir (Tokyo)*. 1993;33:13-18.
5. Phuenpathom, N, Choomuang, M, Ratanalert, S. Outcome and outcome prediction in acute subdural hematoma. *Surg Neurol*. 1993; 40:22- 25.
6. Sakas, DE, Bullock, MR, Teasdale, GM. One-year outcome following craniotomy for traumatic hematoma in patients with fixed dilated pupils. *J Neurosurg*. 1995; 82 : 961-965
7. Fell, DA, Fitzgerald, S, Moiel, RH, Caram, P. Acute subdural hematomas. Review of 144 cases. *J Neurosurg*. 1975;42:37-42.
8. Jang, HS, Lee, YB, Chung, C, Lee, KC, Park, YS, Mok, JH. Acute subdural hematoma: an analysis of 244 operated cases. *J Korean Neurosurg Soc*. 1996; 25:111-118.
9. Richards, T, Hoff, J. Factors affecting survival from acute subdural hematoma. *Surgery*. 1974;75:253-258.
10. Mwang'ombe NJM, Kiboi J. Factors influencing the outcome of severe head injury at Kenyatta National Hospital. *EAMJ* 2001;78: 238-41
11. Howard, MA, Gross, AS, Dacey, RG, Winn, HR. Acute subdural hematomas: an age-dependant clinical entity. *J Neurosurg*. 1989;71:858-863.
12. Mosenthal, AC, Lavery, RF, Addis, M, Kaul, S, Ross, S, Marburger, R. Isolated traumatic brain injury: age is an independent predictor of mortality and early outcome. *J Trauma*. 2002;52:907-911.
13. Munro, PT, Smith, RD, Parke, TR. Effect of patients' age on management of acute intracranial haematoma: prospective national study. *BMJ*. 2002;325:1001-1006.
14. Kotwica, Z, Brzezinski, J. Acute subdural hematoma in adults: an analysis of outcome in comatose patients. *Acta Neurochir (Wien)*. 1993;121:95-99.
15. Marshall, LF, Gautille, T, Klauber, MR. The outcome of severe closed head injury. *J Neurosurg*. 1991;75:528-536.
16. Wilberger, JE Jr, Harris, M, Diamond, DL. Acute subdural hematoma: morbidity, mortality, and operative timing. *J Neurosurg*. 1991;74:212- 218.
17. Wilberger, JE Jr, Harris, M, Diamond, DL. Acute subdural hematoma: morbidity and mortality related to timing of operative intervention. *J Trauma*. 1990;30:733-736.
18. Klun, B, Fettich, M. Factors influencing the outcome in acute subdural hematoma. A review of 330 cases. *Acta Neurochir (Wien)*. 1984;71:171-178.
19. Haselsberger, K, Pucher, R, Auer, LM. Prognosis after acute subdural or epidural hemorrhage. *Acta Neurochir (Wien)*. 1988;90:111-116.
20. Koc, RK, Akdemir, H, Oktem, IS, Meral, M, Menkü, A. Acute subdural hematoma: outcome and outcome prediction. *Neurosurg Rev*. 1997;20:239-244.
21. Kyu-Hong, K. Predictors for Functional Recovery and Mortality of Surgically Treated Traumatic Acute Subdural Hematomas in 256 Patients. *J Korean Neurosurg Soc*. 2009;45:143-150.
22. Stening, WA, Berry, G, Dan, NG, Kwok, B, Mandryk, JA, Ring, I. Experience with acute subdural hematomas in New South Wales. *Aust N Z J Surg*. 1986;56:549-556.
23. Stone, JL, Rifai, MH, Sugar, O, Lang, RG, Oldershaw, JB, Moody, RA. Acute subdural hematoma: progress in definition, clinical pathology, and therapy. *Surg Neurol*. 1983;19:216-231.
24. Ritter, AM, Muizelaar, JP. Brainstem blood flow, pupillary response and outcome in patients with severe head injuries. *Neurosurg*. 1999;44:941-948.
25. Bullock, MR, Chesnut, R, Ghajar, J, Gordon, D, Hartl, R, Newell, DW. Surgical management of acute subdural hematomas. *Neurosurg*. 2006;58(3):16-24.

26. Leach, P, Childs, C, Evans, J, Johnston, N, Protheroe, R, King, A. Transfer times for patients with extradural and subdural haematomas to neurosurgery in Greater Manchester. *Br J Neurosurg*. 2007;21:11-15.
27. Seelig, JM, Becker, DP, Miller, JD, Greenberg, RP, Ward, JD, Choi, SC. Traumatic acute subdural hematoma: major mortality reduction in comatose patients treated within four hours. *New Engl J Med*. 1981;304:1511-1518.