

OTHER

#P17 - Management of patients admitted for Acute liver failure in a subsaharan country

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Background & Aims

Acute liver failure (ALF) remains a rare pathology, with management that remains codified and the prognosis sometimes at stake. Very few studies have been carried out in Africa, more particularly in sub-Saharan Africa. It seemed important to conduct a study to assess these aspects in our context. To evaluate the management of patients admitted for ALF in a resource limited sub saharan country?

Methods

This was a descriptive and cross-sectional study, carried out at the General Hospital of Douala (HGD) over a period of 10 years from January 1, 2013 to December 31, 2022. The collection was made using of a questionnaire. Included were all patients admitted to the Resuscitation department and the Hepato-Enterology department of the Hospital for acute liver failure (ALF) aged over 18 years, of both sexes, and whose files were usable.

Results

We collected 27 patients with ALF out of a total of 617 patients registered for hepatopathy, giving an incidence of 4.37%. The average age of our patients was 39.4 ± 14 years with extremes ranging from 17 to 69 years. The most represented age group was those of 17-39 with a frequency of 63%. The male sex was the most represented with a sex ratio of 1.45. The toxic cause was found in almost all our patients, ie 74.12%, dominated by traditional pharmacopoeia products at 59.3%. The most common reason for consultation was jaundice. hepatomegaly and hepatic encephalopathy were the most found physical signs respectively at 77.8% and 66%. Nearly 18.5% of patients were admitted to intensive care and 81.5% to the medical department. Treatment was based on Lactulose 48.1% of cases, Rifaximin 7.4% of cases, antibiotic therapy 74.5% of cases, corticosteroid therapy 22.2% of cases, N-Acetyl-cysteine 7.4% of cases, and vitamin K 40.7% of cases. There was no liver transplantation. The most common complications were coagulopathy in 84% of cases, metabolic and hydro-electrolytic disorders in 85% of cases and hepatic encephalopathy in 66.7% of cases. The mortality rate was 40.7% of cases.

Conclusions

ALF is a rare but severe pathology. The treatment is symptomatic without access to liver transplantation. The mortality rate remains high at 40.7%.