Epidemiology and Risk Factors for Invasive Fungal Infections among patients with Hematological Malignancies in Colombia

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Background

Invasive fungal infection (IFI) is a potentially complication in patients lethal with hematological malignancies (HM).

Purpose: The current study aimed to investigate the epidemiology of IFI in patients with HM hospitalized in non-HEPAfiltered rooms (resource-limited settings) in a reference center in Colombia.

Methods

A cross-sectional, retrospective study was patients involving HM conducted and pulmonary infection hospitalized in a tertiary hospital in Bucaramanga, Colombia, between 2015-2020.

The primary outcome was proven/probable IFI according to the EORTC/MSGERC criteria. A descriptive and group comparison analysis was performed between patients with IFI and those with non-fungal infections. Multivariate stepwise logistic regression analysis identified the main risk factors for the development of IFI.

Va	ariable
Pr	ior history of pulmonary
in	fection, n (%)
In	-hospital chemotherapy, n (%)
Fe	brile neutropenia, n (%)
Du m	uration of neutropenia (days), edian (IQR)
Pr	ofound neutropenia, n (%)
Se	ptic shock, n (%)
La m	ctate-dehydrogenase U/L, edian (IQR)
To SD	tal serum protein g/dL, mean :
Se SC	rum albumin level g/dl, mean:
Тс (Ю	tal bilirubin mg/dl, median QR)
PI	atelets x103/µL, median (IQR)
м	ould-active prophylaxis, n (%)
Ar n	ntibiotic use more than 3 week (%)
He m	ospital length of stay (days), edian (IQR)
Ba	cteremia, n (%)

Table 1. Univariate analysis of patient characteristics
 compared between patients with invasive fungal infection and patients with non-fungal infection.



Results

• In 201 patients, the prevalence of proven/probable IFI was 21.39% (43 cases). • The most common IFI was caused by *Aspergillus spp.* (41.8%), followed by *Candida spp.* (34.8%), *Mucor spp.* (6.9%), *Penicillium spp.* (4.6%) and *Cryptococcus neoformans* (4.6%).

• In hospital-mortality was 77.1% (155 cases).

	Total (n=201; 100%)	Invasive fungal infection (n=43;	Non-fungal infection	p value
		21.39%)	(n=158; 78.61%)	
	23 (11.44%)	11 (25.58%)	12 (7.59%)	0.001
	105 (52.5%)	29 (67.44%)	76 (48.41%)	0.027
	109 (54.23%)	29 (67.44%)	80 (50.63%)	0.050
	14 (5-25)	25 (15-39)	10 (4-19)	0.000
	70 (34.83%)	25 (58.14%)	45 (28.48%)	0.000
	109 (54.23%)	31 (72.09%)	78 (49.37%)	0.008
	353 (231-671)	490 (324-954)	313 (217-583)	0.000
:	5.97 (5.01-6.71)	5.4 (4.85-6.1)	6.03 (5.31-6.86)	0.027
:	2.98 (0.69)	2.79 (0.67)	3.04 (0.69)	0.04
	0.66 (0.43-1.37)	0.95 (0.6-1.55)	0.6 (0.41-1.33)	0.012
	55 (21-142)	33 (16-93)	68.5 (24-160)	0.016
	45 (22.38%)	3 (6.97%)	42 (26.58%)	0.000
•	48 (24%)	22 (51.16%)	26 (16.56%)	0.000
	35 (17-56)	43 (33-78)	30 (15-53.5)	0.001
	88 (43.78%)	25 (58.14%)	63 (39.87%)	0.032

Mould-active prophylaxis	ŀ	0.20
Antibiotic use more than 3 weeks		
History of pulmonary infection		
Neutropenia duration ≥ 21 days		
	0.05	0.25

Associated factors to develop invasive fungal infections (IFIs) in patients with HM in the multivariate analysis.

Conclusions

- Patients with HM in resource-limited settings have a high prevalence of IFI with elevated mortality.
- The use of mould-active prophylaxis is associated with a significantly lower occurrence of IFI.
- Cost-effective strategies for prevention and early diagnosis of IFI are required to improve survival in patients with HM.



Odds ratio

