Respiratory Viral Infections in Patients with Hematologic Malignancies

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Results

Background

→ Community acquired-respiratory virus infections (CA-RVI) are an important cause of lower respiratory tract infection (LRTI) with increased risk of morbidity and mortality in patient with hematologic malignancies (HM).

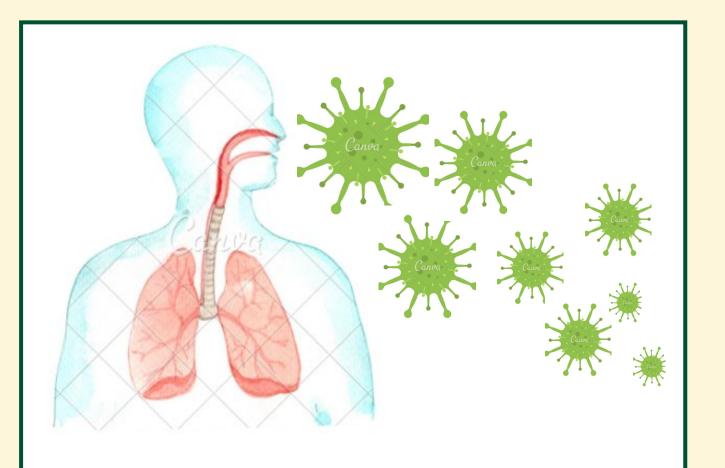
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HM	n (%)		
Age range (median)	23-92 (61) years		
n= 104			
Table 1. Patient characteristics			

Table 2. Patient outcomes / Table 3. LRTI risk factors

Outcomes	Total	Exposure to chemotherapy (n=80)	No exposure to chemotherapy (n=24)
L RTI (%)	31/84 (36.90%)	24 (30%)	7 (29.17%)
Requiring hospital			
admission within 7 days of RV diagnosis (%)	37/75 (49.33%)	28 (35%)	9 (37.5%)
Admission to ICU with respiratory failure (%)	7/37	5 (6.25%)	2 (28.57%)
Requiring mec. vent. (%)	(18.92%) 4/7 (57.14%)	3 (3.75%)	1 (14.28%)
All-cause mortality at 30 days (%)	5 (4.81%)	4 (5%)	1 (14.28%)
Risk Factors	LRTI (n=31/84)	No LRTI (n=53/84)	
Age median (range)	61 (23-92)	61 (24-82)	
Disease status RV diagn. Remission (c/p) Refractory/Relapsed	7 (22.58%) 17 (54.84%)	5 (9.43%) 3 (5.66%)	
ANC (<500 10 ³ cells/uL)	12 (38.71%)	2 (3.77%)	
ALC (<1000 10 ³ cells/uL)	20 (6.45%)	10 (18.87%)	

- Data on respiratory viral infections other than Influenza and RSV in this population is still lacking.
- → Goal: analyze trends of CA-RVI in our HM population



Methods

Study design:

- \rightarrow Retrospective cohort study
- → Analyzed data from multiplex qPCR-based respiratory viral panel (RVP) samples from January 2016-May 2022

Non Hodgkin's Lymphoma (%)	32 (30.77%)
Acute Myelogenous Leukemia (%)	19 (18.27%)
Hodgkin's Lymphoma (%)	4 (3.85%)
Chronic Lymphocytic Leukemia (%)	7 (6.73%)
Acute Lymphoblastic Leukemia (%)	11 (10.58%)
DLBCL + Hodgkin's Lymphoma (%)	3 (2.88%)
Myelofibrosis (%)	3 (2.88%)
Aplastic Anemia (%)	2 (1.92%)
Multiple Myeloma (%)	10 (9.62%)
T cell Lymphoma/Leukemia (%)	4 (3.85%)
Chronic Myelogenous Leukemia (%)	1 (0.96%)
Myelodysplastic Syndrome (%)	5 (4.81%)
Others (%)	3 (2.88%)

RV (Etiology)	n (%)
RHV/ENT (%)	53 (50.96%)
PIV 1-4 (%)	6 (5.77%)
Influenza A/B (%)	17 (16.35%)
RSV A-B (%)	15 (14.42%)
ADV (%)	2 (1.92%)
CoV (%) - <i>Non SARS-CoV 2</i>	7 (6.73%)
HMPV (%)	3 (2.88%)

 \rightarrow Data collected +/- 30 days from RV diag.

Study population:

- → Adult patients with HM with unique episode of CA-RVI
- → Diagnosed with CA-RVI at University of Miami/Sylvester Comprehensive **Cancer Center**

Statistical analysis (in progress)

→ Univariate/multivariate analysis to identify risk factors to develop LRTI and compared outcomes in patients with and without exposure to chemotherapy within 30 days prior to RVI

Other population characteristics	n (%)	
Disease status at time of RV diagnosis		
Remission (complete/partial) (%)	28 (26.92%)	
Refractory/Relapsed (%)	38 (36.54%)	
Other/Not reported (%)	38 (36.54%)	
		\rightarrow
Immunosuppressive agents		
Chemotherapy within 30 days (%)	80 (76.92%)	
Corticosteroids within 30 days (%)	36 (36.62%)	\rightarrow
Setting of diagnosis		
Community-acquired (%)	75 (72.12%)	
Hospital-acquired (%)	29 (27.88%)	\rightarrow
Source of PCR		
Nasal (%)	97 (93.27%)	
BAL (%)	7 (6.73%)	

Conclusions ymptomatic CA-RVI are a common cause of LRTI in patients vith HM (36.90%) emains unclear if exposure to chemotherapeutic agents vithin 30 days prior to RV diagnosis is associated with less avorable outcomes. A-RVI showed relatively low 30-day mortality even in this ulnerable population (4.80%)

arger studies are needed to further characterize outcomes in patients with CA-RVI and HM.

References

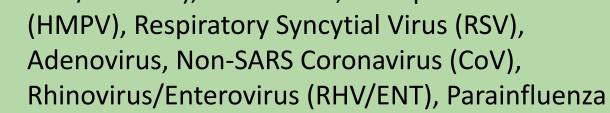
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Virologic testing

→ Nasopharyngeal and BAL specimens were tested using multiplex RT-PCR for the following respiratory viruses: Influenza A (including AH3/AH1200), Influenza B, Metapneumovirus

Exclusion criteria

- → Patients with concomitant stem cell transplantation
- → Patients with concomitant CAR-T cell therapy







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