

Réanimation des patients d'oncohématologie (POH)

Collège de Réanimation Médicale

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Dr Faten May

Sepsis chez les malades cancéreux en Réa: évolution clinique et facteurs pronostiques

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Critically ill patients with cancer and sepsis: Clinical course and prognostic factors ☆☆☆★

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Introduction

- POH: terrain d'immunosuppression
(chimiothérapie, irradiations, corticoïdes...)
- Sepsis: Risque important, motif
d'hospitalisation en Réa
- Morbidité, mortalité élevées

Objectif:

- Déterminer les caractéristiques et l'évolution clinique des malades ayant un cancer, hospitalisés en Réa pour sepsis
- Préciser les facteurs indépendamment associés à la mortalité chez ces malades

Patients et méthodes:

- Etude prospective: cohort study: Janvier 2003 et Juillet 2007

Critères d'inclusion:

- Adultes >18 ans
- Ayant un cancer admis en réa pour sepsis

Critères de non inclusion:

- Rémission complète >5 ans
- Séjour <24 h
- réadmissions

Analyse Statistique:

- Les variables continues sont exprimées en médiane ($\pm$ IQR).
- Une analyse univariée et multivariée (régression logistique) a été réalisée pour déterminer les variables associées à la mortalité chez ces malades.

Résultats:

Caractéristiques de la population étudiée et de l'infection en cause du sepsis

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Table 1 Main patients' characteristics and outcomes

Variables	All patients (n = 563)
Demographics	
Age (y)	59.2 ± 17.8
Male sex	301 (54%)
Comorbidity score (ACE-27)	
None-mild	424 (75%)
Moderate-severe	139 (25%)
Cancer-related characteristics	
Type of cancer	
Locoregional solid tumor	348 (62%)
Metastatic solid tumor	88 (16%)
Low-grade hematologic malignancy	39 (7%)
High-grade hematologic malignancy	88 (16%)
Cancer status	
Controlled/remission	260 (46%)
Active—newly diagnosed	200 (36%)
Active—recurrence/progression	103 (18%)
Performance status	
0-1	269 (48%)
2-4	294 (52%)
Neutropenia	71 (13%)
Severity of acute illness at ICU admission	
SAPS II score (points)	51.0 ± 16.1
SAPS II—predicted mortality (%)	48.4 ± 26.8
SOFA on the first day of ICU (points)	8 (5-11)
Acute organ failures (n)	2 (1-3)
Mechanical ventilation	489 (87%)
Dialysis	110 (20%)
Vasopressors	372 (64%)
Outcome data	
ICU LOS (d)	9 (4-18)
Hospital LOS (d)	23 (11-43)
End-of-life decisions at the ICU	165 (29%)
ICU mortality	289 (51%)
Hospital mortality	364 (65%)
6-mo mortality	407 (72%)

Results are expressed as mean ± SD, median (25%-75% interquartile range), and n (%). LOS indicates length of stay; ACE-27, Adult Comorbidity Evaluation—27.

Variables	All patients (n = 563)
Proof of infection	
Clinically suspected	180 (32%)
Microbiologically proven	383 (68%)
Severity	
Sepsis	48 (9%)
Severe sepsis	143 (25%)
Septic shock	372 (66%)
Acquisition	
Community acquired	227 (40%)
Nosocomial	336 (60%)
Positive blood cultures	114 (20%)
Pathogens^a	
Gram-positive bacteria	168 (30%)
Enterococci	92 (16%)
<i>Staphylococcus aureus</i>	61 (11%)
Streptococci group D	16 (3%)
Pneumococci	13 (2%)
Other	12 (3%)
Gram-negative bacteria	297 (53%)
<i>Escherichia coli</i>	91 (16%)
<i>Pseudomonas</i>	74 (13%)
<i>Klebsiella</i>	72 (13%)
<i>Enterobacter</i>	46 (8%)
<i>Proteus</i>	34 (6%)
<i>Morganella</i>	18 (4%)
<i>Stenotrophomonas maltophilia</i>	14 (3%)
Other	92 (16%)
Fungi	43 (8%)
<i>Candida</i> species	35 (6%)
Other fungi	8 (1%)
Other infectious agents	19 (3%)
Multiresistant bacteria	81 (14%)
Site of infection	
Lung	246 (44%)
Abdomen	172 (31%)
Urinary tract	42 (8%)
Skin/soft tissue	35 (6%)
Primary bloodstream infection	24 (4%)
Central nervous system	11 (2%)
Other/unknown	57 (10%)
More than 1 site of infection	24 (4%)

Results expressed as mean ± SD, median (25%-75% interquartile range), and n (%).

^a Three hundred eighty-three patients had microbiologically proven infections; patients could have more than 1 pathogen.

Résultats:

Analyse multivariée: facteurs associés à la mortalité

Table 3 Multivariate analysis of hospital mortality predictors

Variables	Coefficients	OR (95% CI)	P
Type of admission			
Emergency surgery		1.00	
Medical	0.784	2.19 (1.40-3.42)	.001
Cancer status			
Controlled/remission		1.00	
Active—newly diagnosed	0.565	1.76 (1.12-2.75)	.013
Active—recurrence/ progression	0.884	2.42 (1.35-4.35)	.003
Performance status			
0-1		1.00	
2-4	1.272	3.57 (2.36-5.97)	<.001
Site of infection			
Urinary tract infection		1.00	
Indeterminate or other site of infection	1.189	3.28 (1.57-6.86)	.002
SIRS criteria (n)			
At least 2		1	
3 or 4	0.589	1.80 (1.20-2.72)	.014
Organ dysfunctions			
Cardiovascular ^a	0.664	1.94 (1.27-2.94)	.008
Respiratory ^a	0.827	2.29 (1.24-4.23)	.002
Renal	0.750	2.12 (1.34-3.35)	.001
Constant	-3.568		

n = 563.

Area under receiver operating characteristic curve, 0.80 (95% CI, 0.76-0.84); Hosmer-Lemeshow goodness-of-fit ($\chi^2 = 6.106$, $P = .635$).

OR indicates odds ratio.

^a For these variables, the OR are also affected by the associated interaction term: cardiovascular dysfunction × respiratory dysfunction (coefficient, 1.839; odds ratio, 6.29 [1.19-33.32]; $P = .032$).

Résultats:

Courbes de survie de Kaplan-Meier

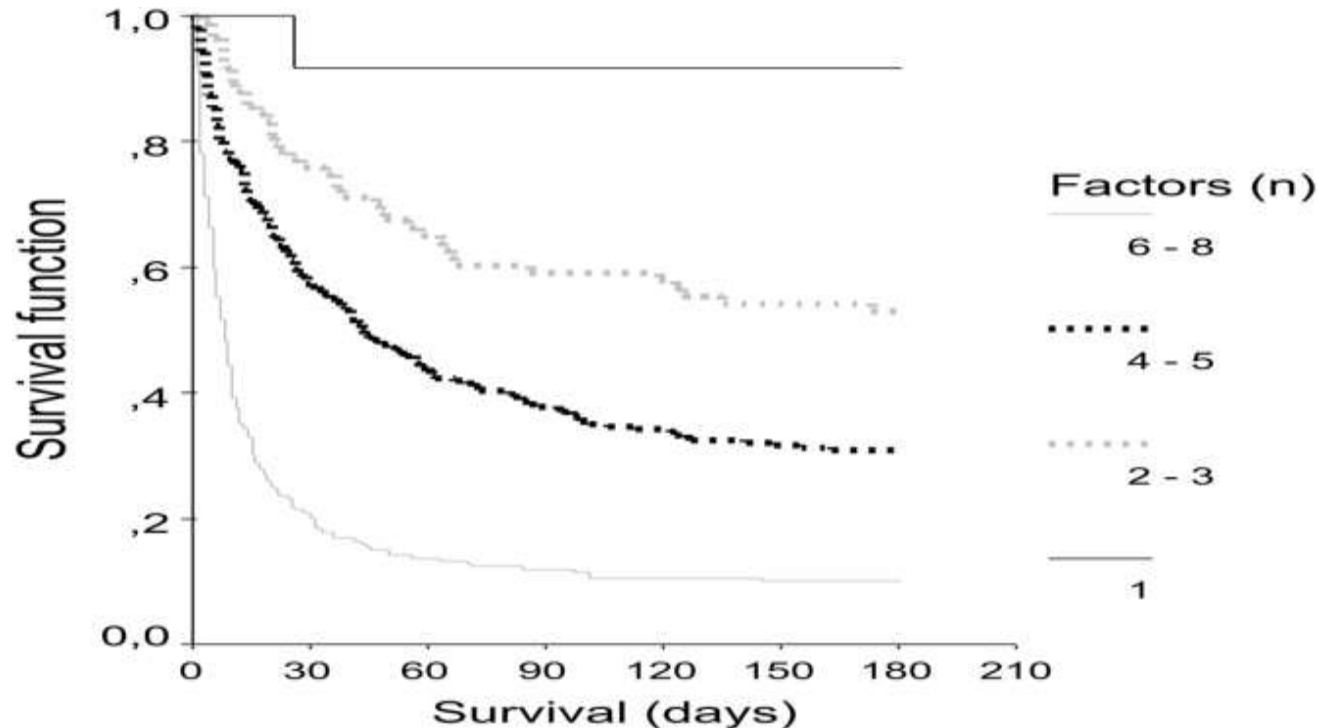


Fig. 1 Survival curves for patients with cancer and sepsis according to the number of independent risk factors (log-rank test, 143.26; $P < .001$). Patients were classified into categories of risk for death: low (1 factor), moderate (2 or 3 factors), high (4 or 5 factors), and very high risk (6-8 factors).

Conclusion:

- Le sepsis est une complication fréquente chez les malades ayant un cancer
- associé à une mortalité élevée.
- Aide les réanimateurs et les oncologistes à une meilleure décision thérapeutique
- Discussion avec les patients et leurs familles
- Contribue à une meilleure stratification du risque
- Besoins d'autres études

Facteurs de risque de SDRA chez les malades d'oncohématologie en neutropénie fébrile

Available online <http://ccforum.com/content/13/6/R173>

Research

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Risk factors for acute respiratory distress syndrome during neutropenia recovery in patients with hematologic malignancies

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Introduction

- La neutropénie peut être associée à une détérioration de l'oxygénation et une exacerbation d'une pathologie pulmonaire préexistante
- FDR de SDRA chez les malades d'OH n'ont pas été étudiés

Objectifs:

- Description des ces malades
- Déterminer les FDR de SDRA

Patients et méthodes:

- Etude de cohorte, Mars 2002-Aout 2008
- Tous les malades d'OH hospitalisés en réa avec une neutropénie fébrile

Résultats: Caractéristiques de la population

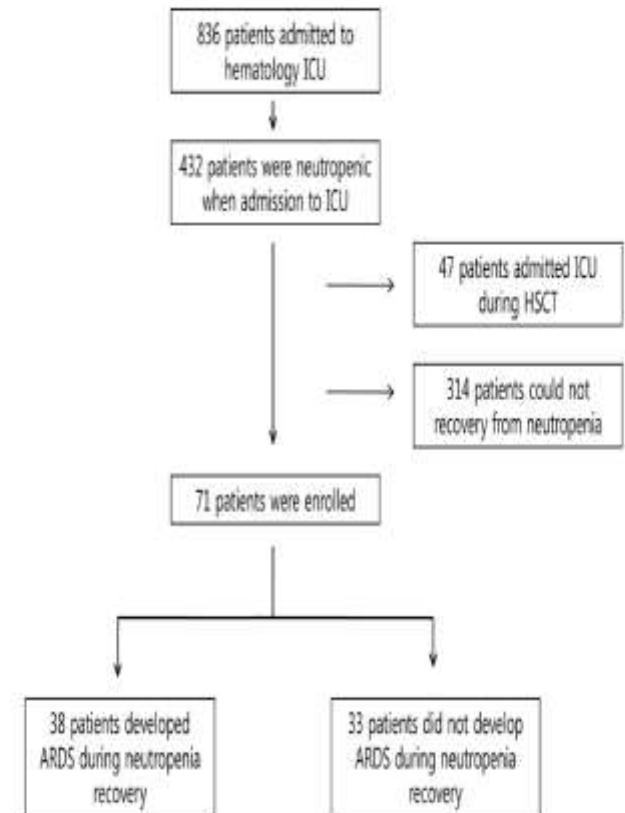
Table 1

Characteristics of the 71 patients with hematologic malignancies who recovered from neutropenia during intensive care unit stay

Characteristic	No (%) or mean $\pm$ SEM
Age, year	45.71 $\pm$ 1.50
Male	33 (46.5%)
AML	35 (49.3%)
ALL	22 (31.0%)
Lymphoma	8 (11.3%)
Multiple myeloma	4 (5.6%)
Relapsed malignancy	17 (23.9%)
History of HSCT	13 (18.3%)
Neutropenia duration, days	22.54 $\pm$ 1.65
ARDS during neutropenia recovery	38 (53.5%)
Pneumonia during neutropenia	45 (63.4%)
Mechanical ventilation	53 (74.6%)
Renal replacement therapy	21 (29.6%)
In-ICU mortality	50 (70.4%)

AML = acute myeloid leukemia; ALL = acute lymphoblastic leukemia; ARDS = acute respiratory distress syndrome; HSCT = hematopoietic stem cell transplantation; ICU = intensive care unit; SEM = standard error of the mean.

Figure 1



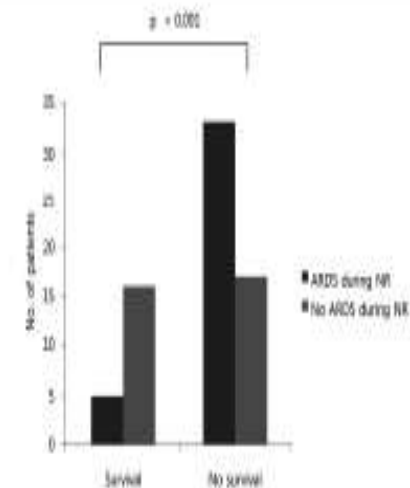
Study subjects. ARDS = acute respiratory distress syndrome; ICU = intensive care unit; HSCT = hematopoietic stem cell transplantation.

Analyse univariée:

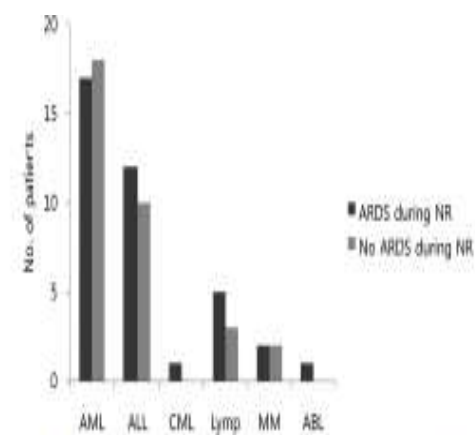
Comparison of patients with and without ARDS during neutropenia recovery

	ARDS during NR	No ARDS during NR	Odds ratio	95% CI	P value
Age	45.99 ± 1.79	45.38 ± 2.51			0.84
Male	20 (52.6%)	13 (39.4%)	1.71	0.66-4.40	0.27
AML	17 (44.7%)	18 (54.5%)	0.68	0.26-1.72	0.41
ALL	12 (31.6%)	10 (30.3%)	1.06	0.39-2.91	0.91
Relapsed malignancy	10 (26.3%)	7 (21.2%)	1.33	0.44-3.99	0.62
Time from diagnosis, days	244 ± 98.7	140 ± 56.8			0.38
Anthracycline treatment	20 (52.6%)	18 (54.5%)	0.93	0.36-2.36	0.87
Cyclophosphamide treatment	9 (23.7%)	5 (15.2%)	1.74	0.52-5.83	0.37
Methotrexate treatment	6 (15.8%)	3 (9.1%)	1.88	0.43-8.12	0.49
Total chemotherapy times	2.79 ± 0.27	2.36 ± 0.25			0.26
Total chemotherapy times ≥3 times	20 (52.6%)	14 (42.4%)	1.51	0.59-3.86	0.39
Previous history of HSCT	8 (21.1%)	5 (15.2%)	1.88	0.43-8.18	0.52
Time between chemotherapy and onset of neutropenia	3.45 ± 0.69	2.18 ± 0.65			0.19
Time between chemotherapy and onset of neutropenia >10 days	3 (7.9%)	1 (3.0%)	2.74	0.27-27.73	0.62
Neutropenia duration, days	22.4 ± 2.3	22.7 ± 2.4			0.94
Neutropenia duration >10 days	34 (89.5%)	26 (78.8%)	2.29	0.61-8.66	0.22
Platelet transfusion during neutropenia	38 (100%)	32 (97%)			0.47
Plasma transfusion during neutropenia	33 (86.8%)	26 (78.8%)	1.78	0.51-6.25	0.37
Pneumonia during neutropenia	32 (84.2%)	13 (39.4%)	8.21	2.68-25.07	< 0.001
RRT during neutropenia	13 (34.2%)	8 (24.2%)	1.63	0.57-4.60	0.36
ICU admission for respiratory failure	25 (65.8%)	10 (30.3%)	4.42	1.63-12.03	0.003
Mechanical ventilator therapy	34 (89.5%)	19 (57.6%)	6.26	1.80-21.75	0.002
In-ICU mortality	33 (86.8%)	17 (51.5%)			< 0.001

Figure 2



Number of patients who survived or died during intensive care unit stay. ARDS = acute respiratory distress syndrome; NR = neutropenia recovery.



Number of patients according to underlying disease. ABL = acute biphenotypic leukemia; AML = acute myeloid leukemia; ALL = acute lymphoblastic leukemia; ARDS = acute respiratory distress syndrome; CML = chronic myeloid leukemia; Lymph = lymphoma; MM = multiple myeloma; NR = neutropenia recovery.

Analyse multivariée des caractéristiques des malades:

over, neutrophil function is defective because many neutrophils are dysfunctional during neutropenia during HSCT. In HSCT, more intense chemotherapy is

Table 3

Multivariate analysis of patient characteristics

	Odds ratio	95% CI	P value
Mechanical ventilator	3.68	0.91-14.82	0.067
Pneumonia during neutropenia	4.76	1.41-16.00	0.012
ICU admission for respiratory failure	2.44	0.79-7.57	0.121

ICU = intensive care unit. Goodness of fit (Hosmer-Lemeshow) chi-squared *P* value = 0.904.

- La survenue d'une pneumonie augmente le risque de SDRA chez les malades en neutropénie fébrile

Conclusion:

- La survenue d'une pneumonie chez les POH en neutropénie fébrile augmente le risque de SDRA

Merci