Baseline metabolic tumor volume Prognosis value in Hodgkin lymphoma

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Background

- CT Tumor volume influences the outcome of HL patients (*Willet CG, JCO 1988; Gobbi P, JCO 2001*): difficult to implement in routine clinical practice
- Few studies have evaluated the prognosis impact of the total baseline metabolic volume (TMTV0) in Hodgkin lymphoma

(Song MK, Cancer Sci 2013; Tseng D, Radiat Oncol2012)

- Various methodologies
- Contradictory results

Study design

- Retrospective single center study
- **59 consecutive patients** with a first diagnosis of HL between January 2007 and January 2010
- PET performed at baseline (PET0) and after 2 cycles of chemotherapy (PET2)
- No treatment change on the basis of PET2 results
- Assessment of:
 - Total Metabolic Tumor Volume at baseline (MTV0)
 - Tumor bulk (>10 cm) at baseline (CT scan)
 - ΔSUVmaxPET 0-2

TMTV0 Assessment

- A region of interest (ROI) was drawn around each foci FDG uptake.
- In each ROI, voxels presenting a threshold of 41% SUVmax were incorporated to define tumor volumes (*Meignan M et al, EJNM 2014;41:1113-22*)
- Extranodal involvement :
 - the liver, lung and bone marrow were considered involved only if there was focal uptake,
 - Spleen involvement was considered if there was focal uptake or diffuse uptake >150 % of the liver background.
- All tumors volume were added to assess the TMTV0
- All of the images were reviewed by 2 nuclear medicine physicians blinded to the patients' outcomes





Patients characteristics

- Median age 36 y (16 76)
- Histological type: NS = 76%, MC 12%
- Stage III/IV = 63%, Bulk>10cm = 15%
- IPS>3 = 61%
 - ABVD = 85%, Radiotherapy = 23%
 - Median Fu = 50 months :
 - 10 progression/relapse (17%),
 - 5 Death (8%)

TMTV0

- Median (range): 117 ml (4 1611)
- Cut-off value to predict treatment failure: 225 ml



- Reproducibility between the 2 readers:
 - Mean absolute difference = 21 ml
 - TMTV0 <225 or >225 ml: Kappa = 0.9 (very good)

	TMTV0 >225 ml TMTV0 ≤225 ml n = 17 n = 42		р	
	$\frac{11-17}{10}$	n = 42		
Median age at diagnosis (vears)	11 (/0) 31 (17 - 63)	11 (/0) 37 5 (16-76)	NS	
Median age at diagnosis (years)	51 (17 - 05)	57.5 (10-70)		
Gender				
Male	14 (82)	26 (62)	NC	
Female	3 (18)	16 (38)	113	
Histological type				
Lymphocyte rich	1 (6)	4 (10)		
Mixed cellularity	2 (12)	5 (12)	NS	
Nodular sclerosis	12 (71)	33 (79)	NJ	
Unclassified	2 (12)	0		
Ann Arbor Stage				
- 1	1 (6)	4 (10)	NS	
- 11	2 (12)	15 (36)	NS	
- 111	2 (12)	8 (19)	NS	
- IV	12 (71)	15 (36)	<0.025	
Bulky Tumor (mass>10cm)	7 (41)	2 (5)	<0.002	
IPS ≥ 3	14 (82)	22 (52)	0.04	

PFS according to TMTV0



PFS according to Bulk



PFS according to PET2 results

Rossi C et al , JNM 2014, 55:569-73

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∆SUVmaxPET0-2			
>71%	12 (71)	37 (88)	
≤71%	5 (29)	5 (12)	NS

TMTV0 and Δ SUVmaxPET0-2-based prognostic model

- In univariate analysis :
- TMTV0 (≤225cc vs >225 cc): 4y-PFS = 85% vs 42%; p = 0.001
- Bulky tumor (>10cm vs ≤10cm): 4y-PFS = 44% vs 78%; p <0.04</p>
- ΔSUVmaxPET0-2 (≤71% vs >71%): 4y-PFS = 82% vs 30%; p <0.0001</p>
- In multivariate analysis: only ΔSUVmaxPET0-2 (p= 0.0005; RR= 6.4) and MTV0 (p< 0.007; RR= 4.2) remained independent predictors for PFS

PFS according to MTV0 and Δ SUVmaxPET0-2

PFS according to TMTV0 and PET2 Deauville score

Conclusions

- TMTV0 is more relevant that tumor bulk to predict outcome of patients with HL, and adds significant prognosis insights to interim PET response assessment
- The combination of TMTV0 with ∆SUVmaxPET0-2 allows identifying 3 subsets of HL patients with significantly different outcomes that may help clinicians to guide therapeutic strategy
- These results have to be validated in larger series