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Radioembolization-Induced Liver Disease (REILD) in Hepatocellular Carcinoma (HCC) Patients after Treatment with Yttrium-90 (Y90) - A retrospective cohort study

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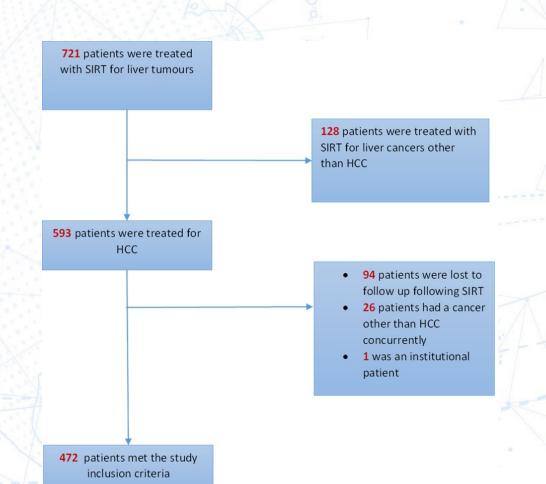
Background + Aim

- Radioembolization-Induced Liver Disease (REILD) is defined as veno-occlusive liver injury caused by radiation treatment to nontumourous parenchyma
- Incidence varied between 0-8% in most reports
- Risk factors remain poorly defined and previous reports were based on relatively small cohorts
- Aim: Determine the risk factors for REILD in a cohort of patients treated with Y90 Selective Internal Radiation Therapy (SIRT-Y90) for HCC in a high-volume tertiary institution

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Methods

- This is a retrospective cohort of 472 patients treated with SIRT-Y90 for non-metastatic HCC from 2007 to 2019 at the National Cancer Centre Singapore (NCCS) and Singapore General Hospital (SGH)
- Patients that were treated for liver cancers other than HCC, had a concurrent cancer or lost to follow up were excluded



Methods

- REILD definition: Presence of ascites and jaundice between 4-12 weeks post Y90 in the absence of tumour progression or bile duct obstruction
- Patient demographics, clinical history, pertinent laboratory values and radiological findings were collected pre- and 6 months post-treatment
- Univariate and multivariate analyses of major clinical factors were carried out

Results

- 12 patients (2.54%) developed REILD
- Risk factors:
 - Baseline ALBI score of 2 (P=0.037)

Results

- Risk factors:
 - Bilobar HCC (P=0.012)
 - BCLC C HCC (P=0.002)





Results

- Outcomes:
 - Majority of patients (8/12, 66.7%) developed low WBC count post Y90, all due to a decrease in lymphocytes, whereby lymphocyte count decreased by a mean of 53.4%
 - No patients demised from REILD

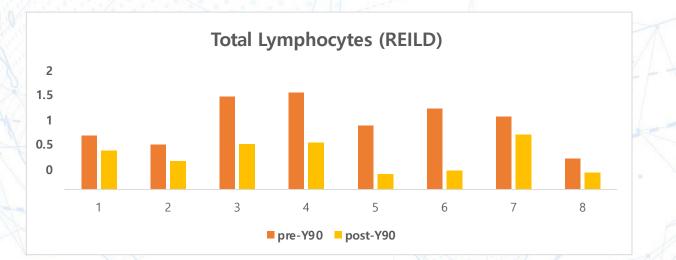


Table 1: Summary of Clinical Findings of REILD Patients Post Y90 SIRT

Table 2: Comparison between REILD and Non-REILD Patients

Baseline		N = 12	Proportions			REILD	Non-REILD	Р
Characteristics				Baseline		N = 12 (2.54%)	N = 460 (97.5%)	
Gender	Male	8	66.7%	Characteristics				
Ethnicity	Female	4	33.3%	Age	Mean (SD)	65.3 (9.41)	64.0 (11.2)	0.664
	Chinese	7	58.3%		Median (IQR)	64 (60.5-69.0)	65 (58.0-71.0)	
	Malay	0	0.00%	Child Pugh	A	8 (66.7%)	391 (85.0%)	0.075
					В	4 (33.3%)	69 (15.0%)	
	Indian	2	16.7%	ALBI Grade	1	0 (0.00%)	130 (28.3%)	0.037
	Others	3	25.0%		2	21 (100%)	297 (64.6%)	
Alcohol	Y	1	11.1%		3	0 (0.00%)	33 (7.17%)	
	N	8	88.9%	BCLC Stage	А	0 (0.00%)	77 (16.7%)	0.002
	NA	3			В	0 (0.00%)	171 (37.2%)	
Change in WBC	Increased	4	33.3%		С	12 (100%)	208 (45.2%)	
	Decreased	8	66.7%		D	0 (0.00%)	4 (0.870%)	
Percentage Decrease	Mean (SD)	27.4 (18.9)		Actiology of	Viral	8 (66.7%)	292 (63.5%)	0.450
in WBC (N = 8)				Hepatitis	(Hep B/C)			
	Median (IQR)	25.1 (12.9-34.2)			Non-Viral	4 (33.3%)	118 (25.7%)	
Change in	Increased	0	0.00%	ester Alexan	Others	0 (0.00%)	50 (10.9%)	
Lymphocytes				Post Y90 AFP	<400	3 (25.0%)	265 (67.3%)	0.002
	Decreased	12	100%		>=400	9 (75.0%)	129 (32.7%)	
Percentage Decrease	Mean (SD)	53.4 (21.7)			NA	0	66	
in Lymphocytes	anna Solini an Solini			Tumour Burden	Solitary	2 (16.7%)	142 (31.0%)	0.203
(N = 12)					2-5 tumours	1 (8.33%)	90 (19.7%)	
	Median (IQR)	51.2 (34.5-76.1)			>5 tumours	9 (75.0%)	226 (49.3%)	
T:N Ratio	Mean (SD)	3.8 (2.08			NA	0	2	
	Median (IQR)	3.46 (2.79-5.37)		Tumour	Unilobar	2 (16.7%)	241 (52.7%)	0.012
Time Between	Mean (SD)	7.92 (7.73)		Location		2 (10.770)		01012
REILD and				Lotation	Bilobar	10 (83.3%)	216 (47.3%)	
Progression of					NA	0	3	
Disease (PD)				Administered	Mean (SD)	1.74 (1.03)	1.67 (0.951)	0.842
(Weeks)				Y90 Dose (Gbg)	(52)			0.0.2
	Median (IQR)	5.86 (1.86-11.7)		270 2000 (300)	Median (IQR)	1.50 (1.33-1.90)	1.40 (0.945-2.20)	
Time Between PD	Mean (SD)	9.89 (6.94)		Predicted Mean	Mean (SD)	162 (62.1)	154 (83.9)	0.808
and Death	Mean (SD)	9.09 (0.94)		Radiation Dose	Moun (OD)	102 (02.1)	104 (00.7)	0.000
(Weeks)				to Tumour (Gy)				
(WCCKS)	Median (IQR)	10.3 (5.57-13)		to runour (SP)	Median (IQR)	138 (121-180)	132 (96.9-191)	
	Median (IQK)	10.5 (5.57-15)			(i Qit)	100 (121 100)		

Conclusions

- REILD development did not correlate with age or administered Y90 dose
- Severity of cirrhosis estimated by the ALBI score is one of the most important risk factors for REILD development post SIRT-Y90
- Lymphocytopenia is a common early sign of the development of REILD

Authors

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