UNIVERSITY of MARYLAND School of Medicine

Introduction

Pancreatic cystic lesions (PCLs) exhibit a wi clinicopathologic behavior. As such, risk these is important to prevent the progression to malignancy. Theoretically PCLs in liver transplant (LT) increased risk of recipients at accelerated are lifelong carcinogenesis Of the setting IN immunosuppressive medications. With improvements in surgical outcomes, LT patients are living lo understanding the incidence and natural c these lesions is paramount.

AIM

To investigate the risk of malignant progress outcomes of PCLs in LT recipients.

METHODS

>Multiple databases were searched (PubMed, Web of Science) to identify studies that used detection from inception until November 2021.

> Primary outcomes incidence of PCLs in LT and progression to malignancy

>Secondary outcomes included those undergoing surgical resection for progression and change in size over time

The Natural History of Pancreatic Cystic Lesions in Liver Transplant Recipients: **A Systematic Review and Meta-Analysis** Andrew Canakis¹, Anusha Vittal,² Smit Deliwala,³ Benjamin Twery,¹ Preet Patel,¹ Justin Canakis,¹ Prabhleen Chahal⁴

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stratifying				
proceion to				

Figure 1: Individual estimates and pooled proportions of progression of pancreatic cystic lesions to malignancy in post liver transplant patients

Study

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onger and course of	Gill 2009 Girometti 2009 Coletti 2014 Ngamuengphon 2014 Mendoza 2015 Ngamuengphon 2015 Trikudanathan 2015 Macinga 2016 Liu 2017 Vidhvarkorn 2017		0.01 [0.00, 0.08] 0.01 [0.00, 0.08] 0.04 [0.00, 0.42] 0.00 [0.00, 0.06] 0.02 [0.00, 0.27] 0.02 [0.00, 0.13] 0.01 [0.00, 0.13] 0.01 [0.00, 0.04]	
Embase and ECS for GC	RE Model	Dreportion (legit coole)	0.01 [0.00, 0.02]	
		Proportion (logit scale)		

Proportion [95% CI]

- PCLs)
- 3.7 ± 1.5 years.
- 93.91, p = 0.01).
- CI, 0.09 0.93, I2 0%, p = 0.04).



RESULTS

> 12 studies met inclusion criteria (17,862 patients with 1411

> The pooled proportion of new PCL development in post-LT patients was 68% (95% CI, 42 – 86, I^2 94%) over a follow up of

> The relative risk of developing PCL in LT recipients compared to non-LT recipients was associated with a significant 67% reduction (RR 0.33; 95% CI, 0.14 – 0.79, l^2

> Among 4158 patients receiving LT, 295 developed PCL and 4 of them progressed to malignancy; the pooled progression to malignancy was 1% (95% CI, 0 – 2, I2 0%) (Figure 1).

> LT was associated with a significant 71% reduction in relative risk of undergoing surgical resection (RR 0.29; 95%)

CONCLUSION

Compared to non-transplant patients, incidental PCLs in LT patients do not carry a higher risk of malignant transformation.

In the setting of immunosuppressive medication, the risk of malignant progression appears to be negligible, and these

patients can be followed like non-LT patients based on guidelines. Furthermore, our findings emphasize that incidental PCLs should not preclude a LT evaluation.