

## Data Sharing Statement

<b>Article Info</b>	<a href="https://dx.doi.org/10.21037/jgo-21-501">https://dx.doi.org/10.21037/jgo-21-501</a>	
<b>Item</b>	<b>Question</b>	<b>Authors' Response (place "-" if not applicable)</b>
1	Would you like to share data collected for your study to others?	We are happy to share our data upon reasonable request.
2	If not, would you like to share the reason for your decision?	-
3	What data in particular will be shared?	Malignancy and clinical outcomes data.
4	Any other documents will be shared? Such as study protocol, statistical analysis plan, informed consent form, clinical study report, analytic code.	Statistical analysis plan, study protocol.
5	When will data availability begin?	From the publication date.
6	When will data availability end?	Two years within the publication date.
7	To whom will you share the data?	Medical oncologists, gastroenterologists, and/or interventional radiologists interested in the study of treatment of symptomatic portal hypertension in the setting of pseudocirrhosis.
8	For what type of analysis or purpose?	For analysis of TIPS for the treatment of non-HCC malignant pseudocirrhosis.
9	How or where can the data/documents be obtained?	Emails can be sent to the corresponding author at: <a href="mailto:lashreve@gmail.com">lashreve@gmail.com</a>
10	Any other restrictions?	We may balance the potential benefits and risks for each request and then provide the data that could be shared. Data sharing may be subject to approval from our institution's IRB.