

ESPS PEER-REVIEW REPORT

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Title: Inflammatory bowel disease in liver transplanted patients

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CLASSIFICATION	LANGUAGE EVALUATION	SCIENTIFIC MISCONDUCT	CONCLUSION
<input checked="" type="checkbox"/> Grade A: Excellent	<input type="checkbox"/> Grade A: Priority publishing	Google Search:	<input type="checkbox"/> Accept
<input type="checkbox"/> Grade B: Very good	<input checked="" type="checkbox"/> Grade B: Minor language polishing	<input type="checkbox"/> The same title	<input type="checkbox"/> High priority for publication
<input type="checkbox"/> Grade C: Good		<input type="checkbox"/> Duplicate publication	
<input type="checkbox"/> Grade D: Fair	<input type="checkbox"/> Grade C: A great deal of language polishing	<input type="checkbox"/> Plagiarism	<input type="checkbox"/> Rejection
<input type="checkbox"/> Grade E: Poor	<input type="checkbox"/> Grade D: Rejected	<input checked="" type="checkbox"/> No	<input type="checkbox"/> Minor revision
		BPG Search:	<input checked="" type="checkbox"/> Major revision
		<input type="checkbox"/> The same title	
		<input type="checkbox"/> Duplicate publication	
		<input type="checkbox"/> Plagiarism	
		<input checked="" type="checkbox"/> No	

COMMENTS TO AUTHORS

MANUSCRIPT TITLE: WJG Manuscript-20170128152907 - Inflammatory bowel disease in liver transplanted patients
 MINOR POINTS: INTRODUCTION: ? This is brief and could include additional information on the following -
 o The indications for OLT in PSC patients should be discussed
 o The differences in prognosis and management in PSC in CD vs UC
 o Small duct vs large duct PSC
 o ? RISK FACTORS ASSOCIATED WITH EXACERBATION OR DE NOVO IBD AFTER LIVER TRANSPLANTATION ? Page 2 - Type - “Tree different patterns of disease.....” ? Is risk of recurrent or de novo IBD after OLT different between CD and UC patients? ? CMV mismatching is only discussed briefly - how important is this in work up for transplanation? ? Is the protective effect of 5-ASA against disease flares in only UC patients - presumably so? ? Regarding post OLT immunosuppressive regimens it is suggested to use cyclosporine instead of tacrolimus and azathioprine instead of mycophenylate - are these recommendations supported by guidelines - eg from ECCO? ? Similarly, in patients on azathioprine (or anti-TNF agents) prior to OLT should these be routinely stopped after OLT?
 TREATMENT OF POST-TRANSPLANTATION INFLAMMATORY BOWEL DISEASE ? These recommendations are not specific to IBD management



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in the post OLT patient ? These two sentences are contradictory - "No significant drug interaction between 5-ASA and immunosuppressant's are observed. In some cases 5-ASA may interact with azathioprine and increase the risk of leukopenia" There are several papers showing the interaction of 5-ASA and thiopurines leading to increased 6TGN levels (Eg. . 5-aminosalicylate therapy is associated with higher 6-thioguanine levels in adults and children with inflammatory bowel disease in remission on 6-mercaptopurine or azathioprine. Hande S, Wilson-Rich N, Bousvaros A, Zholudev A, Maurer R, Banks P, Makrauer F, Reddy S, Burakoff R, Friedman S. *Inflamm Bowel Dis*. 2006 Apr;12(4):251-7). ? Table 1 is a large table that describes only 31 patients on A-TNFs. Of more value would be a larger table showing response rates to all agents used to treat IBD post IBD - it would not need to be so detailed for each agent. ? There should be a separate subheading on management of recurrent PSC post OLT. ? There should be a separate subheading on management of rejection post OLT ? COLORECTAL CARCINOMA (CRC) ? With reference to CRC surveillance the scenic guidelines and chromoendoscopy should be referenced. ? CONCLUSIONS ? The authors suggest "considering" cyclosporine and azathioprine as anti-rejection therapy. This is vague and the readers need stronger recommendations. MAJOR POINTS: ? This manuscript is too brief and inconclusive in its present form. More information is required in several areas as is suggested above. ? Most important is specific data, and therefore recommendations, on IBD management in the post OLT population versus the non-transplant population. ? A table summarizing key recommendations in this population would be useful.