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ESPS Peer-review Report
Name of Journal: World Journal of Gastroenterology
ESPS Manuscript NO: 4946
Title: ERDHEIM CHESTER – a rare syndrome with unique endoscopic features.
Reviewer code: 00239992
Science editor: Zhai, Huan-Huan
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Date reviewed: 2013-08-30 00:26

CLASSIFICATION	LANGUAGE EVALUATION	RECOMMENDATION	CONCLUSION
[ ] Grade A (Excellent)	[ ] Grade A: Priority Publishing	Google Search:	[ ] Accept
[Y] Grade B (Very good)	[Y] Grade B: minor language polishing	[ ] Existed	[ ] High priority for publication
[ ] Grade C (Good)	[ ] Grade C: a great deal of language polishing	[ ] No records	
[ ] Grade D (Fair)		BPG Search:	[ ]Rejection
[ ] Grade E (Poor)	[ ] Grade D: rejected	[ ] Existed	[Y] Minor revision
		[ ] No records	[ ] Major revision

COMMENTS TO AUTHORS

Dear Editor, thank you for giving me the opportunity to review this very interesting case report on a rare disease where many new findings have been made lately. i have nevertheless a few comments: -the manuscript should be carefully re-read by english native speaker: there are many mistakes (such as Langerhans (and not langerhance), Erdheim-Chester disease is the proper term, biopsy-proven (instead of proved)....). Erdheim-Chester disease can also be replaced by ECD as well. - i think the exact number of case reports found in the medical litterature should be more clearly stated by the authors (instead of saying "few reports"). - The iconography is very impressive - The bibliography should be completed by the Blood 2011 series on 53 patients (Arnaud et al.) - Can the authors comment on the choice of treatment by vinblastine + prednisone as it is not at all classical regimen in ECD (but is classical in case of Langerhans cell histiocytosis). Why didn't they consider interferon alpha therapy or alternative therapys ? - Authors should as well comment if possible on the evoluttion under treatment: was the endoscopy controled under treatment ?