Dear editors and reviewers:

Thank you very much for considering our revised manuscript entitled "**Monomorphic Epitheliotropic Intestinal T-cell Lymphoma with bone marrow involved: A case report** " (Manuscript NO: 88793, Case Report) .We appreciate the careful review and constructive comments provided by the reviewers of our manuscript. We have studied the comments carefully and provided answers in a point-by-point manner to each of the reviewers' questions. We had made corrections which we hope are suitable for publication in ***World Journal of Clinical Cases***. Below are our answers for questions raised by the reviewers.

Reviewer #1:

Specific Comments to Authors:

The manuscript topic is actual, as monomorphic epithelial intestinal T-cell lymphoma is a rare intestinal lymphoma.

Response: Thank you for your advice.The manuscript reports a patient diagnosed with monomorphic epithelial intestinal T-cell lymphoma.MEITL is a rare intestinal primary T-cell lymphoma with aggressive behavior, a high risk of severe life-threatening complications, and a poor prognosis. The diagnosis of MEITL should be based on clinical manifestations, pathological features, immunohistochemistry and genetic testing results.Due to the rarity of this disease, more relevant studies and case reports are needed. Herein, we report one rare case of monomorphic epithelial intestinal T-cell lymphoma in a 52-year-old female.Thank you very much for your comments and suggestions.