Reviewer's report

Title: A new family with hereditary lysozyme amyloidosis with gastritis and inflammatory bowel disease as prevailing symptoms.

Version: 4 Date: 26 June 2014

Reviewer: Tadashi Nakamura

Reviewer's report:

The review result is as following.

Major revision

General comments:
The case was very interesting clinically, but rare and unique.
The authors should be suggested to revise the manuscript according to my request.

Specific comments:
1. Tell me the mechanisms why the case showed gastrointestinal symptoms.
2. How do we analyze ALys? Could you check the materials for ALys by sending our samples?
3. How degree do you find ALys among patients with sicca symptoms?
4. In patients with symptoms of alimentary tracts, how do we differentiate AA amyloidosis from ALys?