

## Characterizing helicobacter pylori cagA in Myanmar

Type: Article

Abstract:

**Background/Aims:** Differences in the *Helicobacter pylori* infection rate are not sufficient to clarify the dissimilarity of gastric cancer incidence between Myanmar and its neighboring countries. To better understand this trend, the *H. pylori* virulence gene *cagA* was characterized in Myanmar. **Methods:** Glutamate-proline-isoleucine-Tyrosine-Alanine (EPIYA) patterns and CagA multimerization (CM) motifs of *cagA* genotypes were examined by performing polymerase chain reactions and DNA sequencing. **Results:** Of 69 tested *H. pylori* strains, *cagA*-positive patients had significantly more severe histological scores in their antrum than *cagA*-negative patients. Sequence analysis revealed that 94.1% of strains had Western-Type *cagA* containing an EPIYA motif (92.6%) or EPIYT motif (6.4%). The intestinal metaplasia scores in the antral of patients infected with the ABC and ABCC types of *cagA* were significantly higher than those of patients with AB-Type *cagA*. Interestingly, in patients infected with *H. pylori*, 46.3% of strains with three EPIYA motifs contained two identical Western-Typical CM motifs, and these patients showed significantly higher antrum inflammation scores than patients infected with two identical nontypical-CM motif strains ( $p=0.02$ ). **Conclusions:** In Myanmar strains, Western-Type *cagA* was predominant. The presence of CM motifs and the proportion of multiple EPIYA-C segments might partially explain the intermediate gastric cancer risk found in Myanmar.

Author	a) Myint T. b) Miftahussurur M. c) Vilaichone R.-K. d) Ni N., Aye T.T. e) Subsomwong P. f) Uchida T. g) Mahachai V. h) Yamaoka Y.
Source	PLoS ONE
ISSN	19762283
DOI	10.5009/gnl17053
Volume (Issue)	12 (1)
Page	51-57
Year	2018

Keyword:

CagA, *Helicobacter pylori*, Myanmar

Please Cite As:

Harnanti, D. V., Hidayati, A. N., & Miftahussurur, M. (2018). Concomitant sexually transmitted diseases in patients with diagnosed HIV/AIDS: A retrospective study. *African Journal of Infectious Diseases*, 12(Special Issue 1), 83-89.  
doi:10.2101/Ajid.12v1S.12

URL:

- <https://www.scopus.com/inward/record.uri?eid=2-s2.0-85040525961&doi=10.5009%2fgnl17053&partnerID=40&md5=698e9714c68a0befce2924381fd91cac>
- <https://www.ncbi.nlm.nih.gov/pubmed/29619436>