



Aberrant DNA methylation of *MGMT* and *hMLH1* genes in prediction of gastric cancer

J. Jin^{1,2}, L. Xie², C.H. Xie¹ and Y.F. Zhou¹

¹Department of Radiation & Medical Oncology,
Zhongnan Hospital of Wuhan University,
Clinical Cancer Study Center of Hubei Province,
Key Laboratory of Tumor Biological Behavior of Hubei Province,
Wuhan, China

²Ezhou Central Hospital, Ezhou, Hubei, China

Corresponding author: J. Jun / C.H. Xie
E-mail: jinjun_csh@163.com / chxie_65@hotmail.com

Genet. Mol. Res. 13 (2): 4140-4145 (2014)

Received July 11, 2013

Accepted November 27, 2013

Published May 30, 2014

DOI <http://dx.doi.org/10.4238/2014.May.30.9>

ABSTRACT. We aimed to explore the association between aberrant DNA methylation of the O(6)-methylguanine-DNA methyltransferase (*MGMT*) and human mutL homolog 1 (*hMLH1*) genes with gastric cancer. A total of 283 gastric cancer patients who were confirmed by pathological diagnosis were included in our study. Aberrant DNA methylation of *MGMT* and *hMLH1* were detected. The proportions of DNA hypermethylation in *MGMT* and *hMLH1* in cancer tissues were significantly higher than those in remote normal-appearing tissues. The DNA hypermethylation of *MGMT* was correlated with the tumor-necrosis-metastasis stage in gastric cancer tissues. Results showed that individuals with gastric cancer in the N1 and M1 stages had a significantly higher risk of DNA hypermethylation of *MGMT* in cancer tissues [odds ratio (OR) = 1.97, 95% confidence interval (CI) = 1.15-3.37 for the N1 stage; OR (95%CI) = 5.39 (2.08-14.98) for the M1 stage]. In conclusion, we found that aberrant hypermethylation of *MGMT* could be a predictive biomarker for detecting gastric cancer.

Key words: Aberrant DNA methylation; *MGMT*; *hMLH1*;
Gastric cancer