

[本期目录](#) | [下期目录](#) | [过刊浏览](#) | [高级检索](#)[\[打印本页\]](#) [\[关闭\]](#)**论著****S100A9和NMP238在同时放化疗****敏感性不同宫颈癌中的表达及意义**

朱红1, 曾珊1, 曾亮2, 王学伟1, 肖志强3

1.中南大学湘雅医院肿瘤科, 长沙 410008; 2.湖南省肿瘤医院病理科, 长沙 410013;

3.中南大学湘雅医院卫生部蛋白质组学重点实验室, 长沙 410008

摘要:

目的:初步探讨钙结合蛋白A 9 (S100A9) 和核质蛋白238 (NMP238) 的表达在宫颈癌同时放化疗敏感性中的意义。方法: 收集治疗前的中晚期宫颈癌组织标本, 置-80 °C超低温冰箱中保存, 病理诊断均为中分化鳞癌。在进行同时放化疗后, 根据世界卫生组织实体瘤疗效判断标准, 将收集的宫颈癌组织标本分为高敏感组和低敏感组。提取组织总蛋白, 进行双向凝胶电泳和基质辅助激光解吸电离飞行时间质谱 (MALDI-TOF-MS) 分析鉴定差异表达蛋白。然后分别应用 Western印迹和免疫组织化学方法检测组织中的蛋白表达。结果:筛选得到在同时放化疗敏感性不同宫颈癌组织中差异表达蛋白S100A9和NMP238, S100A9蛋白在高敏感组中高表达, NMP238在低敏感组中高表达。Western印迹检测结果与蛋白质组学结果一致。免疫组织化学结果显示S100A9在高敏感组中的表达强度和表达率 (88.3%) 均显著高于低敏感组 (28.6%), NMP238在低敏感组中的表达强度及表达率 (35.0%) 显著高于高敏感组 (85.7%)。结论: S100A9和NMP238的表达与宫颈癌同时放化疗敏感性有关, 可能作为同时放化疗敏感性预测的标志物。

关键词: 宫颈癌 同时放化疗 蛋白质组学 免疫组织化学

Significance and expression of S100A9 and NMP238 in cervical carcinoma tissues with different concurrent chemoradiotherapy sensitivities

ZHU Hong1, ZENG Shan1, ZENG Liang2, WANG Xuewei1, XIAO Zhiqiang3

1. Department of Oncology, Xiangya Hospital, Central South University, Changsha 410008; 2. Department of Pathology, Hunan Tumor Hospital, Changsha 410013; 3. Key Laboratory of Proteomics, National Ministry of Health, Xiangya Hospital, Central South University, Changsha 410008, China

Abstract:

Objective To determine the significance and expression of S100A9 and NMP238 in cervical carcinoma with different concurrent chemoradiotherapy sensitivities. **Methods** Fresh carcinoma tissues were collected from untreated cervical carcinoma patients and preserved at -80 °C. The tissues were classified into 2 groups: a high sensitivity group (HS) and a low sensitivity group (LS) according to their response to concurrent chemoradiotherapy. Protein was separated by 2-dimensional gel electrophoresis (2-DE). Peptide mass fingerprints (PMF) were acquired by matrix assisted laser desorption/ionization time-of-flight mass spectrometry (MALDI-TOF-MS) and the proteins were identified by data searching in the Mascot-database. Differential expressed proteins were assayed by Western blot and immunohistochemistry. **Results** Most of the gels were clear and were successfully and reproducibly analyzed. Intensity and rate of S100A9 expression were higher in the HS group than in the LS group, and those of NMP238 expression were higher in the LS group than in the HS group. **Conclusion** S100A9 and NMP238 expression is associated with concurrent chemoradiotherapy sensitivity in cervical carcinoma.

Keywords: cervical carcinoma; concurrent chemoradiotherapy; proteomics; immunohistochemistry

收稿日期 2008-12-24 修回日期 网络版发布日期

DOI:

基金项目:

通讯作者:曾珊

作者简介:

作者Email: zengshan2000@yahoo.com

参考文献:

[1] Young T W, Mei F C, Yang G, et al. Activation of antioxidant pathways in ras-mediated oncogenic transformation of human surface ovarian epithelial cells revealed by functional proteomics and mass spectrometry [J]. Cancer Res, 2004, 64(13): 4577-4584.

[2] Imamura T, Kanai F, Kawakami T, et al. Proteomic analysis of the TGF-beta signaling pathway in pancreatic carcinoma cells using stable RNA interference to silence Smad4 expression [J]. Biochem Biophys Res Commun, 2004, 318(1): 289-296.

[3] 王绿化.肿瘤同时放化疗治疗的研究进展 [J].中国癌症杂志, 2006, 16 (6): 405-408.

WANG Luhua. Concurrent chemoradiotherapy in cancer treatment [J]. China Oncology, 2006, 16(6): 405-408.

[4] Lee K H, Yim E K, Kim C J, et al. Proteomic analysis of anti-cancer effects by paclitaxel treatment in cervical cancer cells [J]. Gynecol Oncol, 2005, 98(1):45-53.

[5] Castagna A, Antonioli P, Astner H, et al. A proteomic approach to cisplatin resistance in the cervix squamous cell

扩展功能**本文信息**[Supporting info](#)[PDF\(1927KB\)](#)[\[HTML全文\]](#)[参考文献\[PDF\]](#)[参考文献](#)**服务与反馈**[把本文推荐给朋友](#)[加入我的书架](#)[加入引用管理器](#)[引用本文](#)[Email Alert](#)[文章反馈](#)[浏览反馈信息](#)**本文关键词相关文章**[宫颈癌](#)[同时放化疗](#)[蛋白质组学](#)[免疫组织化学](#)**本文作者相关文章**[PubMed](#)

carcinoma cell line A431 [J]. Proteomics, 2004, 4(10):3246-3267.

[6] Gebhardt C, Breitenbach U, Tuckermann J P, et al. Calgranulins S100A8 and S100A9 are negatively regulated by glucocorticoids in a c-Fos-dependent manner and overexpressed throughout skin carcinogenesis [J]. Oncogene, 2002, 21(27):4266-4276.

[7] Ghavami S, Kerkhoff C, Los M, et al. Mechanism of apoptosis induced by S100A8/A9 in colon cancer cell lines: the role of ROS and the effect of metal ions [J]. J Leukoc Biol, 2004, 76(1):169-175.

[8] Hirano J, Wang X, Kita K, et al. Low levels of NPM gene expression in UV-sensitive human cell lines [J]. Cancer Lett, 2000, 153(12):183-188.

本刊中的类似文章

1. 欧阳红1, 匡韦陆2, 周琴2, 贺理礼2, 周略2, 欧阳淑玉2, 申良方2.腺病毒E1A基因对人宫颈癌细胞体外增殖抑制的实验研究[J]. 中南大学学报(医学版), 2009,34(05): 412-417

2. 曹兰琴 黎欣 张怡 李新国 张瑜 .

改良式宫颈癌广泛性子宫切除加盆腔淋巴结清扫术式探讨

[J]. 中南大学学报(医学版), 2006, 31(04): 588-590

3. 曹兰琴, 黎欣, 李新国, 张怡. 子宫颈癌新辅助化疗的疗效[J]. 中南大学学报(医学版), 2009,34(06): 527-530

4. 郭霞1, 阿布力孜·阿布杜拉1, 武贵臻1, 刘开江2. 维吾尔族妇女宫颈癌患者血清蛋白质组的二维液相色谱分析[J]. 中南大学学报(医学版), 2009,34(07): 624-629

5. 曹兰琴, 林秋华, 黎欣, 张怡, 李萃, 李新国. 紫杉醇对人宫颈癌细胞系HCE1影响的蛋白质组学[J]. 中南大学学报(医学版), 2008,33(06): 512-517

6. 杨丞, 李官成, 李跃辉, 胡锦跃, 肖艳, 张志杰. 太空诱变宫颈癌细胞的生物学研究[J]. 中南大学学报(医学版), 2007,32(03): 380-386

7. 高艳娥1, 惠慧1, 张菊2, 樊江波1, 阎小君2. 高危HPV16 E4基因的表达纯化及临床应用[J]. 中南大学学报(医学版), 2008,33(08): 676-681

8. 李新国1, 吴敏2, 张瑜1. 选择性COX-2抑制剂NS-398对宫颈癌细胞的作用及其作用机制[J]. 中南大学学报(医学版), 2007,32(05): 877-882

9. 符慧群1, 陶光实2, 张隽1, 李征3. 榆皮素对小鼠移植宫颈癌的抑制作用[J]. 中南大学学报(医学版), 2007,32(05): 890-894

10. 曹兰琴*, 黎欣, 吴佳捷, 张瑜. 79例子宫颈癌治疗的临床分析[J]. 中南大学学报(医学版), 0,(): 119-120

11. 曹兰琴*, 黎欣, 吴佳捷, 张瑜. 79例子宫颈癌治疗的临床分析[J]. 中南大学学报(医学版), 2005,30(1): 119-120

12. 邓静1, 谭红专1, 杨土保1, 黄昕1, 周书进2. 宫颈癌扩大筛查计划的成本效益分析[J]. 中南大学学报(医学版), 2010,35(5): 470-

13. 孙阳1, 晋龙2, 刘佳华1, 林赛梅3, 杨茵1, 眭玉霞4, 石红1. 细胞自噬对紫杉醇诱导宫颈癌CaSki细胞死亡的影响[J]. 中南大学学报(医学版), 2010,35(6): 557-

14. 陶光实1, 胡锦跃2, 刘凤英1, 吴宜林1, 刘毅智1, 李官成2. HPV16 E6 DNA疫苗诱导小鼠抗宫颈癌主动免疫[J]. 中南大学学报(医学版), 2004,29(1): 11-14

15. 兰菁, 吴宜林. GM6001对宫颈鳞癌HCE1多细胞球体浸润人脐静脉内皮细胞的影响[J]. 中南大学学报(医学版), 2010,35(8): 868-

Copyright by 中南大学学报(医学版)