miR-506: a regulator of chemo-sensitivity through suppression of the RAD51-homologous recombination axis

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Abstract
Ovarian carcinoma is the most lethal gynecologic malignancy. Resistance to platinum is considered the major problem affecting prognosis. Our recent study established that microRNA-506 (miR-506) expression was closely associated with progression-free survival and overall survival in two independent patient cohorts totaling 598 epithelial ovarian cancer cases. Further functional study demonstrated that miR-506 could augment the response to cisplatin and olaparib through targeting RAD51 and suppressing homologous recombination in a panel of ovarian cancer cell lines. Systemic delivery of miR-506 in an orthotopic ovarian cancer mouse model significantly augmented the cisplatin response, thus recapitulating the clinical observation. Therefore, miR-506 plays a functionally important role in homologous recombination and has important therapeutic value for sensitizing cancer cells to chemotherapy, especially in chemo-resistant patients with attenuated expression of miR-506.

Keywords: MicroRNA-506, Homologous recombination, Drug sensitivity, Synthetic lethality, RAD51

Background
Epithelial ovarian cancer remains the most lethal gynecologic malignancy [1, 2]. The majority of cases are diagnosed at an advanced stage, with a 5-year survival rate of only 30%–40% [3]. Debulking surgery and platinum-based chemotherapy play important roles in the management of epithelial ovarian cancer, whereas acquired resistance to platinum is considered a major factor in disease relapse.

Recently, it has been reported that microRNAs (miRNAs), which are single-stranded, non-coding endogenous RNAs, play a crucial role in regulating drug resistance. Possible mechanisms in regulating drug resistance involve the regulation of cell cycle distribution, drug transport, DNA repair, epithelial-mesenchymal transition (EMT), and apoptosis pathways. Platinum-based drugs can form intra- and inter-strand adducts with DNA, which cause DNA double-strand breaks and trigger DNA damage and repair pathways. Thus, enhancement of the DNA repair system plays a major role in platinum resistance. Homologous recombination (HR) is a critical pathway for DNA double-strand break repair; cells with compromised HR machinery are highly sensitive to DNA-damaging drugs [4].

Our recent investigation of The Cancer Genome Atlas (TCGA) database network for high-grade serous ovarian carcinoma showed that microRNA-506 (miR-506) expression was associated with an increased response to therapy and prolonged progression-free survival (PFS) and overall survival (OS). This result was confirmed in another clinically annotated genomics dataset (Bagnoli) [5]. Our previous studies have demonstrated that miR-506 can inhibit EMT and proliferation by targeting the snail family zinc finger 2 (SNAL2) and cyclin-dependent kinase 4/6-Forkhead box protein M1 (CDK4/6-FOXM1) axes, respectively [6–8]. After overexpressing miR-506 in a panel of ovarian cancer cell lines, we analyzed the down-regulated genes via a microarray and observed a decrease in RAD51 levels. When we used the miRNA target prediction algorithm TargetScan 6.0 (Whitehead Institute for Biomedical Research, Cambridge, MA, USA), we found that RAD51
had one predicted binding site for miR-506. We next performed a set of functional validation experiments using reporter gene assays. We found that miR-506 suppressed the reporter activity linked to the 3′-untranslated region (UTR) of RAD51 that contains the predicted binding site. The use of control miRNA or removal of the binding site did not produce the same regulation exhibited by miR-506. These results showed that RAD51 is a direct target of miR-506. Furthermore, the inverse association between miR-506 and RAD51 expression was confirmed in three different cohorts of clinical samples (the TCGA, Bagnoli, and Tianjin cohorts).

We next sought to understand how important the regulation of RAD51 is to the drug response and survival of serous ovarian cancer patients. RAD51 is known to be a critical component of the HR-mediated double-strand DNA break repair machinery, and during this process, RAD51 assembles onto single-stranded DNA as a nucleoprotein filament and catalyzes the exchange of homologous DNA sequences [9]. Because RAD51 is an integral component of the cellular DNA damage response, RAD51 suppression can sensitize cancer cells to DNA-damaging drugs [10–13]. By performing a homology-directed repair assay, we found that transfection with a miR-506 mimic or treatment with small interfering RNA (siRNA) against RAD51 significantly reduced the HR efficiency. In addition, ectopic overexpression of miR-506 led to higher residual DNA damage (as shown by single-cell gel electrophoresis) and less RAD51 foci as compared with controls after cisplatin treatment, confirming the defect in HR.

HR-deficient cells are sensitive to DNA-damaging drugs and poly(ADP) ribose polymerase (PARP) inhibitors, and this effect has been termed synthetic lethality [14, 15]. Recent studies have demonstrated that breast cancer early onset 2 (BRCA2) mutations, and to a lesser extent breast cancer early onset 1 (BRCA1) mutations/methylation, are associated with improved survival and responses to therapy in serous ovarian cancer patients [16–18]. We next examined the effect of miR-506 on the sensitivity to cisplatin and olaparib (a PARP inhibitor) in ovarian cancer in vitro and in vivo and found that miR-506-transfected cells were more sensitive to cisplatin or olaparib than control cells in both 3-(4,5-dimethyl-2-thiazolyl)-2,5-diphenyl-2-H-tetrazolium bromide (MTT) and clonogenic survival assays. Conversely, anti-miR-506 locked nucleic acid (LNA) transfection enhanced RAD51 expression and induced resistance to cisplatin and olaparib in cells. Moreover, the effect of miR-506 on cisplatin and olaparib sensitivity was rescued by overexpressing RAD51 without its 3′-UTR, suggesting that miR-506-mediated sensitivity to cisplatin and olaparib is the result of the suppression of RAD51 expression. Consistent with the results of in vitro experiments, delivery of miR-506 incorporated in 1,2-dioleoyl-sn-glycerol-3-phosphatidylcholine (DOPC) nanoliposomes effectively enhanced the effect of cisplatin and olaparib in an orthotopic ovarian cancer model.

In summary, our recent investigations have revealed that miR-506 can enhance the sensitivity to cisplatin and PARP inhibitors through suppression of the RAD51-HR axis. Furthermore, nanoparticle delivery of miR-506 enhanced the effects of cisplatin and olaparib in orthotopic ovarian cancer models. Thus, miR-506 not only is a robust clinical marker for the chemotherapy response and survival of serous ovarian cancer patients but also has important therapeutic value in sensitizing cancer cells to chemotherapy. This new discovery shows that miR-506 combined with DNA-damaging agent treatment may benefit patients with high-grade serous ovarian carcinoma. Taken together with the findings of our previous study on this small molecule, we have shown that miR-506 has multiple therapeutic effects on regulating the biological behavior of cancer cells (Fig. 1).
Authors’ contributions
GL, FX, and WZ wrote the manuscript. All authors read and approved the final manuscript.

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Compliance with ethical guidelines

Competing interests
The authors declare that they have no competing interests.

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