

## Secondary *BRCA1* Mutations in *BRCA1*-Mutated Ovarian Carcinomas with Platinum Resistance

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### Abstract

Although ovarian carcinomas with mutated *BRCA1* or *BRCA2* are sensitive to platinum compounds, such carcinomas eventually develop platinum resistance. Previously, we showed that acquired resistance to cisplatin in *BRCA2*-mutated tumors can be mediated by secondary intragenic mutations in *BRCA2* that restore the wild-type *BRCA2* reading frame. Here, we show that secondary mutations of *BRCA1* also occur in *BRCA1*-mutated ovarian cancer with platinum resistance. We evaluated nine recurrent *BRCA1*-mutated ovarian cancers previously treated with platinum compounds, including five with acquired platinum resistance, one with primary platinum resistance, and three with platinum sensitivity. Four of the six recurrent platinum-resistant tumors had developed secondary genetic changes in *BRCA1* that restored the reading frame of the *BRCA1* protein, whereas none of the three platinum-sensitive recurrent tumors developed *BRCA1* sequence alterations. We immunohistochemically confirmed restored expression of *BRCA1* protein in two cases with secondary mutations. Intriguingly, the case with primary platinum resistance showed back mutation of *BRCA1* in the primary tumor and showed another secondary mutation in the recurrent tumor. Our results suggest that secondary mutations in *BRCA1* can mediate resistance to platinum in *BRCA1*-mutated ovarian tumors. [Cancer Res 2008;68(8):2581–6]

### Introduction

DNA cross-linking agents such as cisplatin, carboplatin, melphalan, and cyclophosphamide are widely used anticancer drugs. Resistance to these agents is a major obstacle to effective cancer therapy. Patients with ovarian cancer usually respond well to initial chemotherapy with cisplatin and its derivative carboplatin, but over time the majority of patients become refractory to platinum compounds. Ultimately, progression of chemoresistant disease is the major cause of death in women with ovarian carcinoma (1).

*BRCA1* and *BRCA2* are tumor suppressor genes responsible for familial breast/ovarian cancer. *BRCA1* or *BRCA2* (*BRCA1/2*) mutation carriers have increased risks of developing breast and

ovarian cancer (2). Ten percent to 15% of all epithelial ovarian cancers are associated with inherited mutations in *BRCA1/2* (3, 4). The vast majority of tumors from women with germ-line *BRCA1/2* mutations show loss of the wild-type *BRCA1/2* allele and are considered to be *BRCA1/2* deficient (5–7).

*BRCA1/2*-deficient cancer cells are hypersensitive to DNA cross-linking agents including cisplatin (8–10). Consistently, patients developing *BRCA1/2*-mutated ovarian cancer have a better prognosis compared with noncarriers, if they receive platinum-based therapy (8, 11). However, even *BRCA1/2*-mutated ovarian cancers frequently develop platinum resistance.

Recently, we reported that acquired resistance to cisplatin in *BRCA2*-mutated tumors can be mediated by secondary mutations in *BRCA2* that restore the wild-type *BRCA2* reading frame (12). *BRCA2*-mutated cancer cells selected in the presence of cisplatin acquired secondary genetic changes on the mutated *BRCA2* allele, which canceled the frameshift caused by the inherited mutation and restored expression of functional nearly-full-length *BRCA2* protein. Cells with secondary mutations were resistant to cisplatin (12). However, mechanisms of cisplatin resistance in *BRCA1*-mutated ovarian cancer remain unclear. Because both *BRCA1* and *BRCA2* are involved in homologous recombination DNA repair (13, 14) and are required for cellular resistance to cisplatin (9, 10), we hypothesized that similar secondary mutations of the mutated *BRCA1* gene mediate acquired resistance to cisplatin in *BRCA1*-mutated ovarian cancer. Now we report that secondary mutations of *BRCA1* also occur in *BRCA1*-mutated ovarian cancer with platinum resistance.

### Materials and Methods

**Clinical specimens.** Three DNA samples from recurrent *BRCA1*-mutated ovarian cancer patients were obtained from Cedars-Sinai Medical Center (Los Angeles, CA) through the Pacific Ovarian Cancer Research Consortium. Six DNA samples from *BRCA1*-mutated ovarian cancer patients with recurrent disease were obtained from the tissue bank of University of Washington (Table 1). DNA from UW80, UW40, UWF27, and the recurrent tumor from UW91 were obtained after laser capture microdissection. The study was approved by Institutional Review Boards of Fred Hutchinson Cancer Research Center and University of Washington. Clinical data of UW40 patient and a cancer cell line derived from the same patient's recurrent tumor (UWB1.289) have previously been reported (15). Platinum sensitivity was defined as a complete response to treatment maintained without progression for at least 6 mo after platinum therapy. Tumors that were resistant or refractory had progressive disease on platinum therapy, less than a complete response to platinum therapy or progression within 6 mo of completing platinum therapy.

**Sequencing of *BRCA1*.** PCR products of genomic DNA were analyzed for sequence alterations with BigDye Terminator v3.1 Cycle Sequencing Kit (Applied Biosystems) using ABI 3100 Genetic Analyzer (Applied Biosystems). All detected mutations were confirmed by sequencing of an

**Note:** Supplementary data for this article are available at Cancer Research Online (<http://cancerres.aacrjournals.org/>).

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**Table 1.** Secondary genetic changes of *BRCA1* in *BRCA1*-mutated ovarian cancer treated with platinum

Patient	Inherited mutation	Specimen	Clinical platinum sensitivity	Secondary genetic change
Paired specimens				
CS1	185delAG	Pre-platinum	Sensitive	No
		Post-platinum	Sensitive	
CS4	185delAG	Pre-platinum	Sensitive	No
		Post-platinum	Sensitive	
UW91	185delAG	Pre-platinum	Sensitive	Yes (back mutation)
		Post-platinum	Resistant	
CS14	185delAG	Pre-platinum	Sensitive	No
		Post-platinum	Resistant	
UW40	2594delC	Pre-platinum	Resistant	Yes (back mutation)
		Post-platinum	Resistant	Yes (2606_2628del23)
UW80	185delAG	Pre-platinum	Sensitive	Yes (back mutation)
		Post-platinum	Resistant	
UWF27	185delAG	Pre-platinum	Sensitive	Yes (back mutation)
		Post-platinum	Resistant	
Nonpaired posttreatment specimens				
UW208	3867G>T (E1250X)	Post-platinum	Sensitive	No
UW317	3171insTGAGA	Post-platinum	Resistant	No

independently amplified template. Information of all sequencing and PCR primers for *BRCA1* is available in Supplementary Table S1. All nucleotide numbers refer to the wild-type cDNA human sequence of *BRCA1* (U14680.1) as reported in the GenBank database.

**Immunohistochemical studies.** *BRCA1* protein was detected in formalin-fixed paraffin-embedded sections using the mouse monoclonal antibody MS110 (previously called Ab-1; Oncogene Research Products) as previously described (16). The MS110 antibody recognizes an NH<sub>2</sub>-terminal epitope at *BRCA1* (amino acid residues 89–222; ref. 16). Briefly, paraffin sections were deparaffinized, washed in PBS, and treated with steam heat for 20 min using antigen retrieval solution (DAKO). Endogenous peroxidase activity in paraffin sections was quenched by treatment with 3% H<sub>2</sub>O<sub>2</sub> for 5 min. Sections were washed with PBS and nonspecific binding was blocked by treatment for 2 h in 2% bovine serum albumin in PBS. Primary antibody was applied (MS110 at 1:250 dilution) for 14 to 16 h at 4°C. Secondary antibody and streptavidin biotin-peroxidase were from the Universal Large Volume LSAB+, Peroxidase kit (DAKO) and were each applied for 30 min at room temperature. 3,3'-Diaminobenzidine chromagen (Sigma) was used to visualize antibody complexes and sections were counterstained with hematoxylin. The MS110 antibody can result in nonstaining of tissues under certain conditions and can be variable in its results. Therefore, we used nontumor inflammatory and stromal cells as an internal positive control for *BRCA1* protein staining for every section. Decreased protein was only scored if normal cells on the same section were positive. Primary and recurrent tumors were stained side by side under identical conditions. MS110 is the most reliable antibody for the immunohistochemical detection of *BRCA1* (16). Because the MS110 antibody recognizes an NH<sub>2</sub>-terminal epitope in *BRCA1*, it cannot distinguish between wild-type *BRCA1* protein and truncated nonfunctional protein resulting from frameshift mutations occurring distal to the antibody epitope. Thus, MS110 is not suitable for distinguishing functional versus nonfunctional *BRCA1* protein in UW40, UW208, and UW317. Therefore, we evaluated immunohistochemistry only in those tumors with 185delAG

mutations (proximal to the antibody epitope) for which we had paired primary and recurrent paraffin-embedded tumor samples.

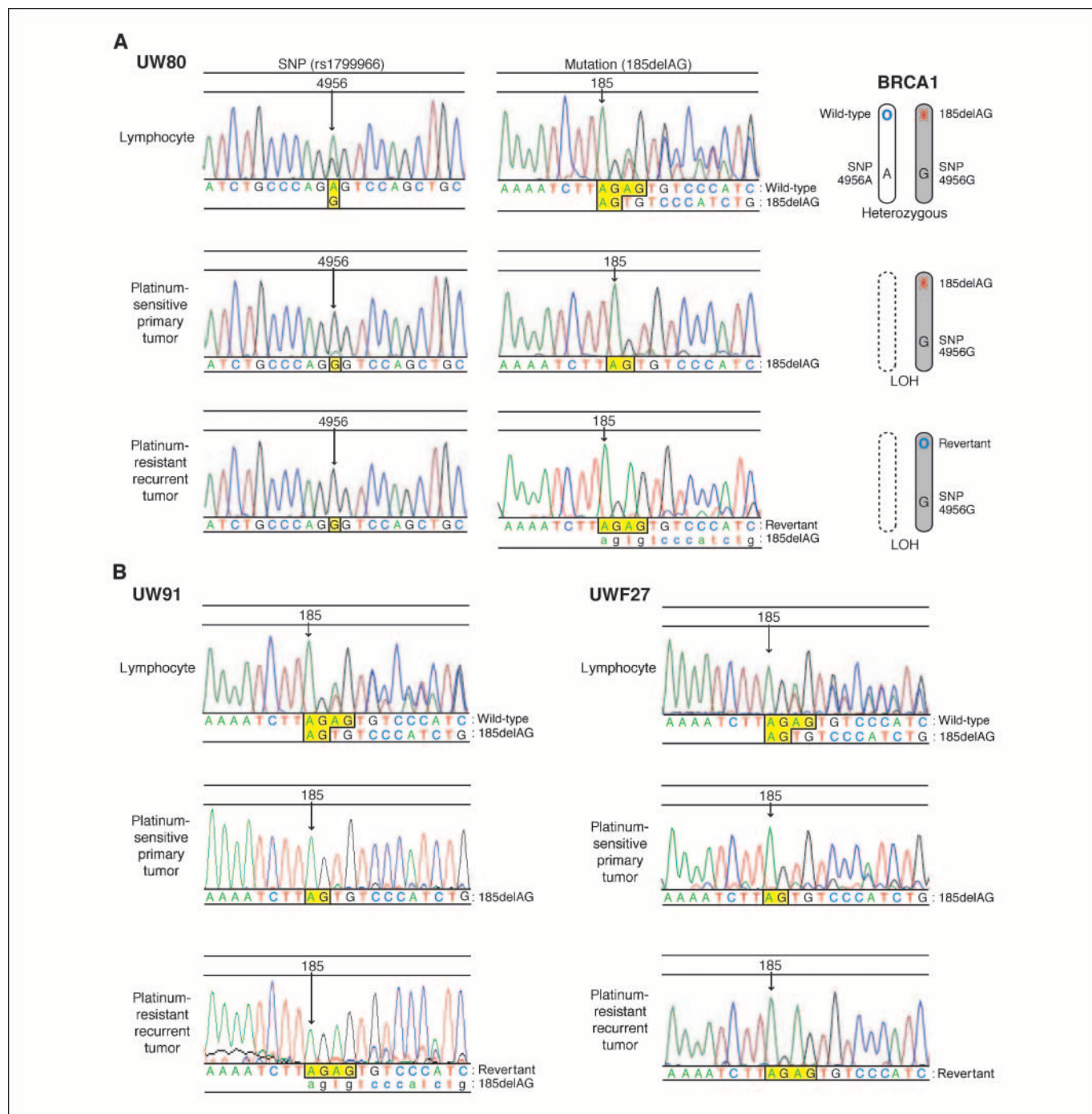
## Results and Discussion

We analyzed nine clinical samples of recurrent *BRCA1*-mutated ovarian cancer previously treated with platinum (Table 1). Six of them were clinically platinum sensitive and three were platinum resistant. One case (UW40) showed primary resistance to platinum, whereas five cases were initially sensitive to platinum and acquired resistance during the disease course. Three of the five tumors with acquired platinum resistance (UW80, UW91, and UWF27) and the primarily resistant tumor (UW40) had secondary genetic changes in *BRCA1*, whereas none of the three cisplatin-sensitive recurrent tumors showed secondary genetic changes in *BRCA1*.

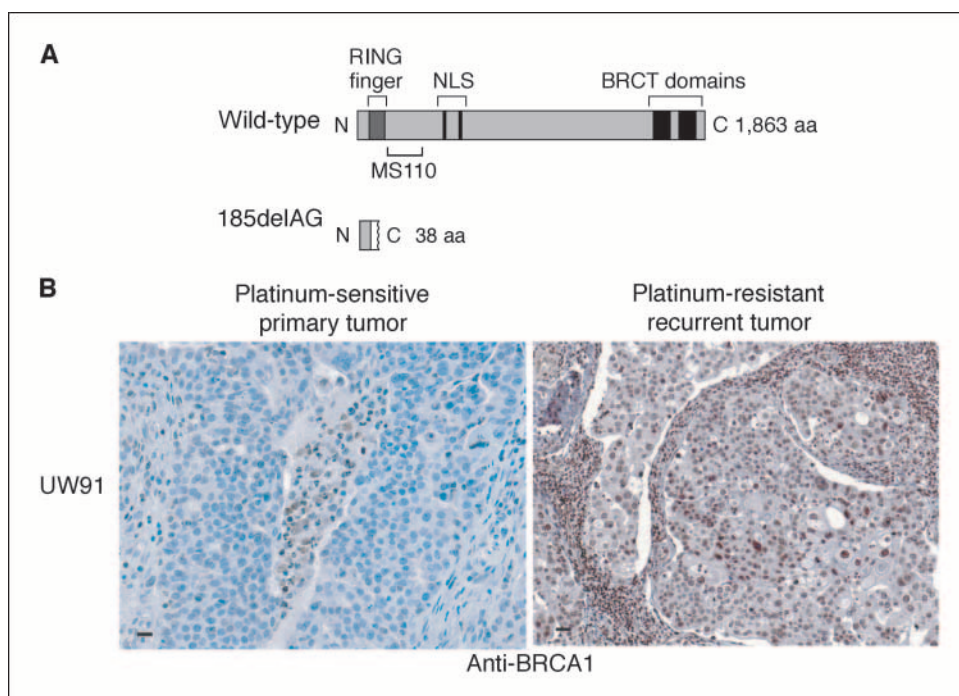
UW80 patient is a heterozygous carrier of a frameshift *BRCA1* mutation, 185delAG, which is common in the Ashkenazi Jewish population (Fig. 1A; ref. 17). In the primary tumor specimen, only *BRCA1* sequence with 185delAG was detected, indicating that the tumor had lost the wild-type allele. Loss of heterozygosity (LOH) of an intragenic *BRCA1* single nucleotide polymorphism (SNP) was also confirmed. In the microdissected recurrent tumor from the same patient after development of resistance to platinum, LOH of the *BRCA1* SNP was confirmed, indicating that the recurrent tumor contained only the same single allele of *BRCA1* as present in the primary tumor and that contamination of nontumor cells was negligible. In this recurrent sample, wild-type *BRCA1* sequence represented >80% of the sequences at the mutation site, suggesting that the recurrent tumor had acquired wild-type *BRCA1* by genetic reversion (back mutation to wild-type).

Similarly, in two more *BRCA1.185delAG* cases (UW91 and UWF27), the primary tumor showed only mutant sequence with loss of the wild-type allele. In the recurrent tumors, we mainly detected wild-type *BRCA1* sequence, suggesting that these two

platinum-resistant tumors acquired wild-type *BRCA1* by genetic reversion (back mutation to wild-type; Fig. 1B). Both cases had confirmation of LOH at intragenic SNPs in the primary and recurrent tumors (Supplementary Fig. S1), indicating the presence



**Figure 1.** Genetic reversion of *BRCA1* mutation in three recurrent *BRCA1*-mutated ovarian cancer cases. **A**, DNA sequences of *BRCA1* in peripheral blood lymphocytes and the primary and recurrent tumors from a patient (UW80) with *BRCA1*-mutated ovarian cancer. In the lymphocytes, a heterozygous SNP of the *BRCA1* locus (4956A/G) was detected, in addition to a heterozygous mutation (185delAG). In the primary tumor, a hemizygous mutation (185delAG) was detected and loss of heterozygosity (LOH) of the SNP was confirmed. In the microdissected recurrent tumor, LOH of the SNP was confirmed, but wild-type sequence was 80% to 90% of the sequences identified at the site of 185delAG. Importantly, the SNP in the recurrent tumor (4956G) is identical to that in the primary tumor (4956G). This indicates that the recurrent tumor had acquired wild-type *BRCA1* by genetic reversion (back mutation to wild-type). Residual 185delAG sequence could represent heterogeneity in the tumor. A speculative model of *BRCA1* alleles in samples from this patient is also depicted. **B**, DNA sequences of *BRCA1* in peripheral blood lymphocytes; the primary and recurrent tumors from two patients (UW91 and UWF27) with *BRCA1*-mutated ovarian cancer. In the lymphocytes, a heterozygous mutation (185delAG) was detected. In the primary tumors, a hemizygous mutation (185delAG) was detected. In the microdissected recurrent tumor specimens, wild-type sequence was detected, suggesting that the recurrent tumors had acquired wild-type *BRCA1* by back mutation. Analyses of intragenic SNPs of these cases are shown in Supplementary Fig. S1.



**Figure 2.** BRCA1 protein expression is restored in recurrent ovarian tumors with genetic reversion of *BRCA1* mutation. **A**, schematic presentation of BRCA1 proteins encoded by 185delAG and wild-type *BRCA1*. Functional domains of BRCA1 protein [RING finger, nuclear localization signals (NLS), and BRCT domains] are depicted. The region that the BRCA1 antibody for immunohistochemical staining recognizes (amino acids 89–222; ref. 16) is also depicted. **B**, immunohistochemical staining of BRCA1 in a primary tumor with *BRCA1.185delAG* mutation and a recurrent tumor with genetic reversion of the *BRCA1* mutation (UW91 patient). The primary tumors are negative for BRCA1 immunostaining, whereas the recurrent tumors are positive for nuclear BRCA1 immunostaining. Bar, 10  $\mu$ m.

of a single *BRCA1* allele that underwent a secondary genetic alteration.

*BRCA1.185delAG* encodes a severely truncated protein with 38 amino acids, and the back mutation should restore full-length BRCA1 protein (Fig. 2A). Consistently, immunohistochemical study revealed that BRCA1 protein expression was absent in the primary tumors of UW80 and UW91, but strongly positive in their recurrent tumors (Fig. 2B and data not shown).

The mechanism for genetic reversion of 185delAG mutations in all three tumors is not clear. At the genomic level, the reinsertion of two missing nucleotides would seem less likely to occur than a secondary downstream deletion that corrects the frameshift and restores the open reading frame. Several possible explanations for this phenomenon are not mutually exclusive. First, we speculate that because 185delAG is in a critical portion of the *BRCA1* gene close to the RING finger (Fig. 2A), a change of one single amino acid here is likely to have deleterious effect on protein function. Similarly, missense mutations in the RING finger such as C61G can be deleterious (18). Therefore, restoration of BRCA1 function may require complete reversion of a mutant in this domain to wild-type. Another possibility is that the particular genomic sequence surrounding 185delAG somehow facilitates the genetic reversion. A third possibility is that other downstream mutations do occur in some tumors but are not detected with our relatively short PCR amplifications. We are unable to do long-range PCR that might identify larger genomic events because of the fragmented nature of our DNA. Thus, it is easier to identify genetic reversion in our samples than larger events that restore the open reading frame.

UW40 patient is a heterozygous carrier of a frameshift *BRCA1* mutation, 2594delC (Fig. 3A and B). The mutant allele encodes an 844-amino-acid protein that lacks BRCT domains (Fig. 3C). The patient had a past history of breast cancer at the age of 42 years and developed ovarian cancer at age 54 years. The primary ovarian carcinoma was clinically resistant to platinum. After six cycles of

platinum and Taxol, the patient was treated with Taxol for three additional cycles with development of progression. She was re-treated with topotecan, then with gemcitabine and Adriamycin to obtain a partial remission. Recurrent tumor was then obtained at the time of surgery for a fistula.

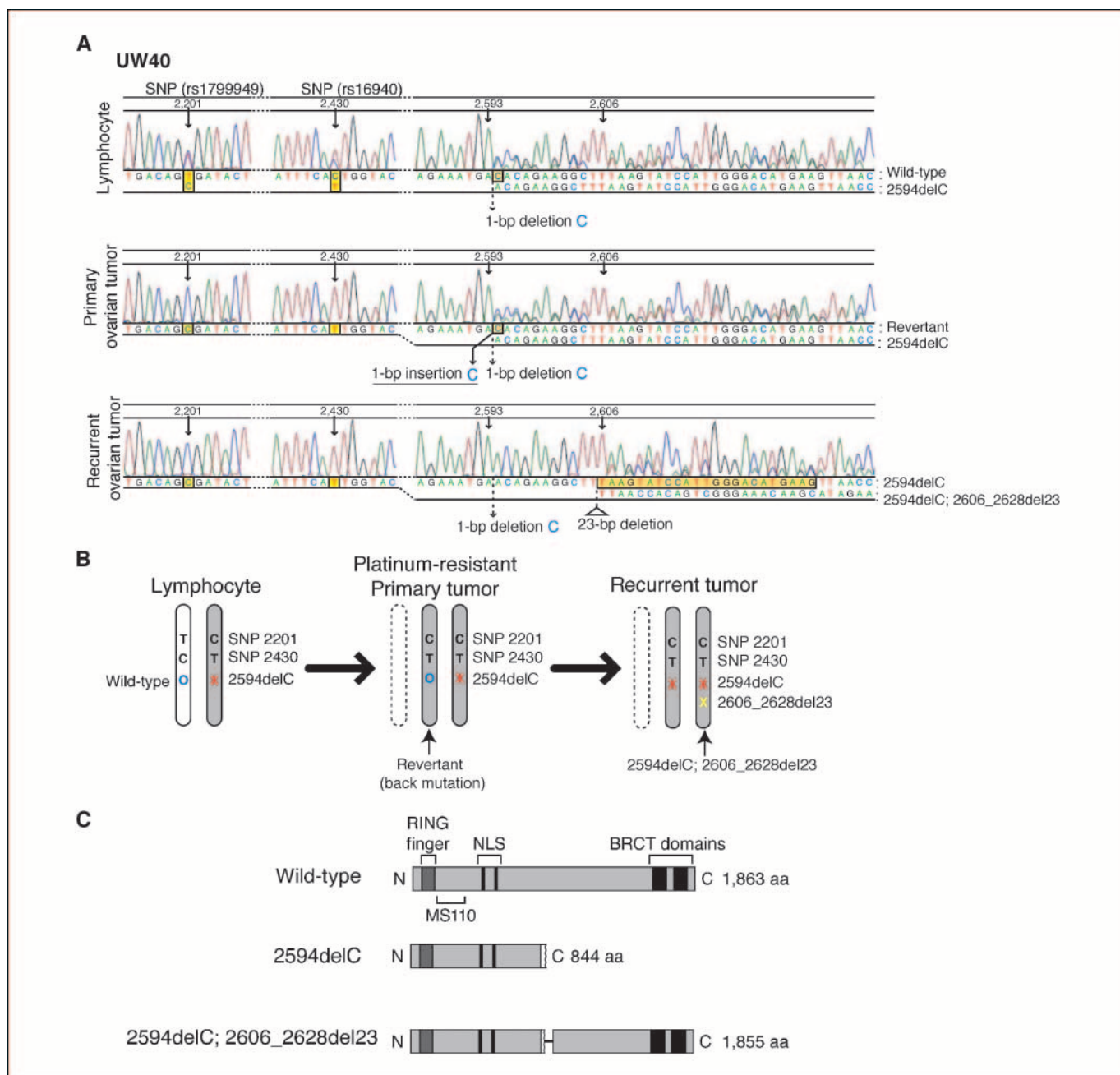
LOH of three intragenic *BRCA1* SNPs (2201C/T, 2430T/C, and 2731C/T) that flank the mutation site was confirmed in both the primary and recurrent tumors (Fig. 3A and B, and data not shown), indicating that contamination by nontumor cells was negligible and that both the primary and recurrent tumors had lost one *BRCA1* allele. Intriguingly, in the primary tumor, both wild-type *BRCA1* sequence and *BRCA1* sequence with 2594delC were detected (Fig. 3A and B). Careful laser microdissection of a separate second sample of this tumor revealed the same result. The presence of both wild-type *BRCA1* sequence and mutant sequence on one allele in the primary tumor suggests that genetic reversion (back mutation to wild-type) occurred on one copy of the mutant allele. We speculate that the presence of the genetically reverted wild-type allele in the primary tumor contributed to the unusual initial platinum resistance of this tumor. The selective pressure for the genetically reverted tumor cells in the primary ovarian tumor is unknown, but may be related to the 11 months of cyclophosphamide, methotrexate, and fluorouracil that the patient previously received for breast cancer. Cyclophosphamide is a DNA cross-linking agent and could exert similar selection pressure for the correction of DNA repair defects as we have observed with exposure to platinum agents.

An alternative explanation for the existence of genetic reversion and other secondary mutations on a mutant allele in this primary tumor is that these events are more common than we appreciate but usually occur in only a small number of primary tumor cells before exposure to chemotherapy. Then, chemotherapy provides the selection pressure for expansion of these subclones in recurrent tumors. Perhaps some reports of retention of the wild-type *BRCA1/2* allele in breast cancers (19) actually represent genetic reversion.

Only careful haplotyping distinguishes tumors with retention of the wild-type allele from those with genetic reversion.

The genetically reverted wild-type allele was then lost at some point during the disease course. In the recurrent tumor, we could no longer detect wild-type sequence, but we found a second site small deletion (2606\_2628del23) in addition to the inherited

mutation (2594delC). The second mutation corrects the frameshift caused by the inherited mutation and the allele *BRCA1*:2594delC;2606\_2628del23 encodes a BRCA1 protein with intact COOH-terminal region, which may be functional (Fig. 3C). The actual genomic DNA sequence of *BRCA1* in the UW40 recurrent tumor is a mixture of 2594delC;2606\_2628del23 and 2594delC without a



**Figure 3.** Genetic reversion of *BRCA1* mutation in a primary ovarian tumor with platinum resistance and another secondary *BRCA1* mutation in a recurrent tumor of the same patient. **A**, DNA sequences of *BRCA1* in peripheral blood lymphocytes and the primary and recurrent tumors from a patient (UW40) with *BRCA1*-mutated ovarian cancer. In the lymphocytes, heterozygous SNPs of the *BRCA1* locus (2201C/T and 2430T/C) were detected, in addition to a heterozygous mutation (2594delC). In the primary tumor, LOH of the SNPs was confirmed, but the primary tumor specimen shows heterozygous sequence of wild-type *BRCA1* and a *BRCA1* mutation (2594delC). In the recurrent tumor specimen, LOH of the SNPs was confirmed and mixed sequences of 2594delC and 2594delC;2606\_2628del23 were detected. **B**, a speculative model of *BRCA1* alleles in samples from this patient. The lymphocytes have both the wild-type and mutant (2594delC) alleles. The primary tumor has lost the wild-type *BRCA1* allele with SNPs 2201T and 2430C, but has retained the allele with the inherited mutation (2594delC) and SNPs 2201C and 2430T. This mutant allele has presumably been duplicated, and one of the mutated alleles is reverted to wild-type by genetic reversion (back mutation to wild-type). In the recurrent tumor, the allele with wild-type *BRCA1* sequence is lost, and the allele with the inherited mutation (2594delC) was again duplicated. One of them obtained a different second mutation (2606\_2628del23), which cancels the frameshift caused by the inherited mutation (2594delC). This model explains why we see mixed sequences of *BRCA1* in the primary and recurrent tumors. **C**, schematic presentation of BRCA1 proteins encoded by 2594delC and 2594delC;2606\_2628del23.

secondary mutation (Fig. 3A). The mixture of sequences on one allele in both the primary and recurrent tumors may have occurred by secondary genetic alterations (reversion to wild-type in the primary tumor and 23-bp deletion in the recurrent tumor) occurring on just one of the duplicated mutant alleles (Fig. 3B), similar to the situation occurring in Capan-1 clones with secondary mutations of *BRCA2* observed in the *in vitro* experiment in the previous report (12). Alternatively, the two sequences could represent tumor heterogeneity, with only a subset of tumor cells acquiring a secondary mutation on the mutant allele. We favor the explanation of tumor heterogeneity in this case, given that a cell line developed from this recurrent tumor contains only mutant sequence without secondary genetic changes (15).

Taken together, secondary mutations that restore the reading frame of mutated *BRCA1* alleles were frequently observed in ovarian tumors with resistance to platinum. These results suggest that secondary mutations of *BRCA1* can be a mechanism of acquired resistance to cisplatin in *BRCA1*-mutated ovarian cancer. In addition, the observation of the back mutation of *BRCA1* in cisplatin-resistant primary tumor of UW40 patient suggests that secondary mutations of *BRCA1* can be a mechanism of primary resistance to cisplatin.

Our present and previous (12) studies emphasize that *BRCA1* and *BRCA2* are not only cancer susceptibility genes but are also critical determinants of clinical sensitivity and resistance to chemotherapy. These studies support the general concept that defective DNA repair leads to chemosensitivity, whereas restoration of functional DNA repair contributes to acquired chemoresistance in tumor cells *in vitro* and *in vivo*.

Testing for secondary mutations in platinum-treated *BRCA1*-mutated cancers may be clinically important because tumors with

secondary mutations are likely to be resistant to platinum and could show cross-resistance to other agents that exploit *BRCA1* defects, such as poly(ADP-ribose) polymerase 1 inhibitors. A larger clinical study is warranted to further determine the prevalence and clinical significance of secondary mutations of *BRCA1* in *BRCA1*-mutated cancer.

Our finding could have implications for sporadic ovarian carcinoma. Although somatic mutation of *BRCA1/2* is rare in sporadic ovarian carcinoma, *BRCA1* is reported to be down-regulated in a subset of sporadic ovarian carcinomas by promoter methylation and other unknown mechanisms (20). *BRCA2* mRNA expression is reported to be undetectable by reverse transcription PCR in 13% of ovarian carcinoma (20). Therefore, it is tempting to hypothesize that down-regulation of *BRCA1/2* causes initial sensitivity to cisplatin, and restoration of functional *BRCA1/2* expression by demethylation of the *BRCA1* promoter or other mechanisms leads to acquired resistance. Testing this hypothesis will be important to elucidate the more general roles of *BRCA1/2* in platinum sensitivity and resistance.

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