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TITLE: Regulation of BRCA1 Function by DNA Damage-Induced Site-Specific Phosphorylation

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BRCA1, a hereditary breast and ovarian specific tumor suppressor, ensures genomic integrity through its control of transcription and repair of damaged DNA. Considerable evidence implicates DNA damage-induced site-specific phosphorylation in the modulation of its biological activity. However, it is not presently clear whether and how the transcription and DNA repair activities of BRCA1 are modulated in response to DNA damage signals. We have engineered and refined a unique combination of biochemical and genetic tools to address this issue. First, we have developed a biochemical means by which to resolve BRCAl-containing complexes involved in transcription from those involved in DNA double-strand break repair. This should render it feasible to identify DNA damage-induced site-specific phosphorylation events with potential functional relevance to the role of BRCA1 in these two processes. Second, we have established fibroblast cultures from brcal-deficient mouse embryos and developed BRCAl-dependent transcription and repair assays based on the use of these cells. This system will expedite the facile and efficient analysis of the effects of targeted BRCA1 mutations at identified or predicted sites of phosphorylation on its transcription and DNA repair activities. Collectively, these studies should illuminate the molecular basis for the caretaker properties of BRCA1.

Tumor suppressor, BRCA1, DNA repair, transcription

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INTRODUCTION

BRCA1, a hereditary breast- and ovarian-specific tumor suppressor, functions in the global maintenance of genome stability, and has been implicated in both transcription and DNA double-strand break repair processes. Considerable evidence implicates DNA-damage-induced site-specific phosphorylation of BRCA1 as a critical regulator of its caretaker properties. However, it is not presently known whether and how the transcription and/or DNA repair activities of BRCA1 are specifically modulated in response to DNA damage. We hypothesized that DNA damage-induced site-specific phosphorylation of BRCA1 regulates its transcription and/or DNA double-strand break repair activities. To provide support for this hypothesis, we proposed first to identify ionizing radiation-induced site-specifically phosphorylated residues on BRCA1 in complex with transcription or DNA double-strand break repair activities, and second, to determine the functional consequence of ionizing radiation-induced site-specific phosphorylation on the transcription and DNA double-strand break repair activities of BRCA1. Toward this objective, our research plan encompassed two major objectives. First, we proposed to biochemically purify from human cells, both prior to and following irradiation, distinct BRCA1-containing multiprotein complexes corresponding to the RNA polymerase II holoenzyme and the DNA double-strand break (DSB) repair complex containing the products of the Rad50, Mre11, and NBS1 (Nijmegan breakage syndrome) genes (Rad50/Mre11/NBS1 complex). Second, we proposed to effect direct comparative analyses of wild type BRCA1 and mutant derivatives bearing substitutions at ionizing radiation-targeted residues for their respective abilities to function in BRCA1-dependent transcription and DNA double-strand break repair assays in vivo.

BODY

Research Accomplishments:

Technical Objective 1. To identify ionizing radiation (IR)-induced site-specifically phosphorylated residues on BRCA1 in complex with either the RNA polymerase II holoenzyme or the Rad50/Mre11/NBS1 DNA double-strand break repair complex.

Task 1: Months 1-9: To purify distinct BRCA1-containing complexes corresponding to the RNA polymerase II holoenzyme and the Rad50/Mre11/NBS1 DNA double-strand break repair complex both prior to and following IR.

In year one, we achieved the biochemical purification of distinct BRCA1-containing multiprotein complexes implicated in transcription and DNA repair, thereby completing a significant portion of Task 1 of Technical Objective 1. The results of these studies, documented in the first year annual summary statement, were incorporated into a manuscript that has been resubmitted for publication following additional experiments, and which we include as an appendix to this annual summary statement (Please Refer to Appendix 1 – Manuscript Preprint).
In year one, we also identified a novel function for BRCA1 in suppressing the ligand-independent transcriptional activity of the estrogen receptor α (ERα), a principal determinant of the growth and differentiation of breasts and ovaries. This observation arose peripherally as a result of the characterization of BRCA1-containing transcription complexes, and the identification of activities previously linked to ERα. Importantly, we documented that clinically validated BRCA1 missense mutations abrogate this repression activity, thereby suggesting that its ERα-specific repression function is important for the biological activity of BRCA1 in breast and ovarian tumor suppression. Our results revealed BRCA1 to be a ligand-reversible barrier to transcriptional activation by unliganded ERα, and suggested a possible mechanism by which functional inactivation of BRCA1 could promote tumorigenesis through inappropriate hormonal regulation of breast epithelial cell proliferation. These studies offer possible insight into the tissue-specific tumor suppressor function of BRCA1 and could suggest defined molecular targets for future intervention in breast cancer. The results of these findings were published in the *Proceedings of the National Academy of Sciences U.S.A* (2001, Vol. 98: 9587-9592; Please refer to Appendix 2 – Manuscript Reprint) (1), and were also documented in the first year annual summary statement.

Using the purification strategy outlined in the appended manuscript preprint, we have succeeded in purifying distinct BRCA1-containing transcription and DNA repair complexes from HeLa S3 cells following exposure to ionizing radiation, thus completing Task 1 of Technical Objective 1.

**Task 2: Months 3-18:** To identify IR-induced site-specifically phosphorylated residues on BRCA1 present in purified RNA polymerase II holoenzyme and Rad50/Mre11/NBS1 DNA double-strand break repair complexes by both mass spectrometric analyses and immunoblot analyses using phosphopeptide-specific antibodies.

We are currently engaged in efforts to identify IR-induced site-specifically phosphorylated residues on BRCA1 present in purified RNA polymerase II holoenzyme and Rad50/Mre11/NBS1 DNA double-strand break repair complexes by immunoblot analysis using commercially available phosphopeptide-specific antibodies and also by mass spectrometric analyses. Thus far, we have encountered significant obstacles in the identification of IR-induced site-specifically phosphorylated residues by mass spectrometric-based approaches. The difficulties encountered in this regard most likely derive from the fact that a relatively small fraction of the total cellular pool of BRCA1 is phosphorylated in response to IR. Consequently, the quantity of phosphorylated target protein thus far purified is below the sensitivity limits of our instruments. We are currently engaged in the preparation of additional starting material for these experiments. The process in laborious, time-consuming, and expensive; however, we are confident that the information to derive from the successful completion of these experiments will be crucial to deciphering how targeted phosphorylation of BRCA1 in response to DNA damage alters its functions within the cell. We are therefore currently engaged in efforts to culture and process cells on an even larger scale than previously anticipated. We are confident of the success of the proposed experimental approach, and on this basis we continue to implement our experimental plan so far as resources permit.
Technical Objective 2. To determine the functional consequence of individual IR-induced site-specific phosphorylation events on the DNA double-strand break repair and transcription activities of BRCA1.

Task 1: Months 9-36: To determine the effects of targeted mutations at identified (or predicted) sites of IR-induced phosphorylation within BRCA1 on its ability to effect DNA double-strand break repair by homologous recombination and nonhomologous end-joining following its ectopic expression in Brca1-deficient cells.

As a first approach toward this objective, we initiated the functional characterization of the BRCA1/Rad50/Mre11/NBS1 DNA double-strand break repair complex purified in Task 1 of Technical Objective 1. In so doing, we made the novel discovery that BRCA1, in complex with Rad50/Mre11/NBS1, plays a critical role in the nonhomologous end-joining pathway of DNA double-strand break repair. This observation has significant implications for the role of BRCA1 in the maintenance of genomic integrity. This study was published in manuscript form in Cancer Research (2002, Vol. 62: 3966-3970; Please refer to Appendix 3 – Manuscript Reprint) (2), and was also documented in the second year midterm summary statement.

In addition to its novel findings, this study also established a cell-free system that should expedite the completion of Task 1 of Technical Objective 2. More specifically, the in vitro complementation system for nonhomologous end-joining described in this study should permit us to assess in a straightforward approach the role of regulatory BRCA1 phosphorylation on its DNA double-strand break repair activities.

Task 2: Months 6-24: To determine the effects of targeted mutations at identified (or predicted) sites of IR-induced phosphorylation within BRCA1 on its ability to control transcription following its ectopic expression in brca1-deficient cells.

As a first approach toward this objective, we have established a BRCA1-dependent transcription-based assay to evaluate the effects of IR-induced site-specifically phosphorylated residues on BRCA1 identified through Task 2 of Technical Objective 1. This assay is based on the functional interaction between BRCA1 and the sequence-specific transcriptional repressor protein ZBRK1. Previously, we showed that BRCA1 is a co-repressor of ZBRK1, a sequence-specific DNA-binding transcriptional repressor of the DNA damage-inducible GADD45 gene that functions in G2/M cell cycle checkpoint control (3). In addition to GADD45, potential ZBRK1 binding sites have been identified in other DNA damage-inducible genes, indicating a prospective global role for ZBRK1 and BRCA1 in the coordinate regulation of DNA damage-response genes (3). Based on these previous observations, we have proposed a model whereby ZBRK1 and BRCA1 coordinately repress a group of DNA damage response genes in the absence of genotoxic stress and, further, that DNA damage-induced cell signals relieve this repression, thereby permitting DNA damage-induced activation of these genes.
The DNA damage-induced cell signals that relieve coordinate repression of DNA damage response genes by ZBRK1 and BRCA1 is likely to involve phosphorylation. In fact, previous studies have revealed that IR-induced protein phosphorylation is required to relieve BRCA1-mediated repression of the GADD45 gene (4). To facilitate studies designed to determine the effects of targeted mutations at identified sites of IR-induced phosphorylation within BRCA1 on its ability to control transcription, we established a BRCA1-dependent ZBRK1 transcriptional repression assay. This assay is designed to assess the BRCA1-dependent repression function of ZBRK1 in mammalian cells from a reporter template bearing ZBRK1 DNA-binding sites. To establish the utility of this system as a means to study BRCA1-dependent ZBRK1 repression, we used this system as a functional readout during experiments designed to functionally dissect ZBRK1. This study revealed that ZBRK1 harbors dual specificity zinc fingers with twin roles in DNA-binding and BRCA1-dependent transcriptional repression. Furthermore, this study provided novel insight into the mechanistic basis by which BRCA1 mediates sequence-specific control of DNA damage-responsive gene transcription by the identification of a novel BRCA1-dependent transcriptional repression domain within the ZBRK1 C-terminus. This C-terminal repression domain (CTRD) functions in a BRCA1-dependent, histone deacetylase-dependent, and promoter-specific manner and is thus functionally distinguishable from the N-terminal KRAB repression domain in ZBRK1, which exhibits no BRCA1 dependence and broad promoter specificity. This study was published in manuscript form in The Journal of Biological Chemistry (2004, Vol. 279: 6576-6587; Please refer to Appendix 4 - Manuscript Reprint) (5), and was also documented in our second year annual summary statement.

In the past year, we further exploited this BRCA1-dependent ZBRK1 transcriptional repression assay to characterize novel BRCA1-associated co-repressors of ZBRK1. More specifically, during an unbiased search for novel interaction partners and possible co-regulators of the CTRD, we identified ZBRK1 itself, suggesting that ZBRK1 can oligomerize through its CTRD. Protein interaction analyses using wild-type and mutant ZBRK1 derivatives confirmed that ZBRK1 can homo-oligomerize both in vitro and in vivo, and further mapped the ZBRK1 oligomerization domain to the CTRD C-terminus. Biochemical analyses, including protein cross-linking and gel filtration chromatography, revealed that ZBRK1 homo-oligomers exist as tetramers in solution. Functionally, we exploited the aforementioned BRCA1-dependent ZBRK1 transcriptional repression assay to show that ZBRK1 oligomerization facilitates ZBRK1-directed transcriptional repression through ZBRK1 response elements (ZREs); requirements for oligomerization-dependent repression include the ZBRK1 CTRD and KRAB repression domains, but not the DNA-binding activity of ZBRK1. These observations suggest that higher order oligomers of ZBRK1 may assemble on target ZREs through both protein-DNA and CTRD-dependent protein-protein interactions. These findings thus reveal an unanticipated dual function for ZBRK1 in both DNA-binding-dependent and -independent modes of transcriptional repression, and further establish the CTRD as a novel protein interaction surface responsible for directing homo- and heterotypic interactions necessary for ZBRK1-directed transcriptional repression. This study was published in manuscript form in the The Journal of Biological Chemistry (2004, Vol. 279: 55153-55160; Please refer to Appendix 5 – Manuscript Reprint) (6).
Our BRCA1-dependent ZBRK1 repression assay may now be exploited to evaluate the influence of IR-induced site-specific phosphorylation of BRCA1 on its sequence-specific co-repressor function. So long as resources permit, we plan to evaluate the influence of IR on the ability of ectopically expressed wild-type BRCA1 to function as a ZBRK1-specific co-repressor. We predict, and will test the possibility, that IR leads to phosphorylation of BRCA1 and consequent relief of ZBRK1-directed repression from the ZBRK1 target reporter plasmid through disruption of the ZBRK1/BRCA1 interaction. Subsequently, we will evaluate BRCA1 derivatives bearing site-directed mutations at sites of IR-induced phosphorylation identified through Task 2 of Technical Objective 1 for their respective abilities to function as IR-reversible co-repressors of ZBRK1. We predict that BRCA1 derivatives bearing mutations at identified sites of IR-induced phosphorylation will function as constitutive ZBRK1 co-repressors through an IR-insensitive interaction with ZBRK1. This assay should therefore provide us with a rapid and facile approach to test the hypothesis that IR-induced site-specific phosphorylation of BRCA1 modulates its transcriptional regulatory activities.

**Key Research Accomplishments To Date (Year 3 accomplishments underlined).**

- Biochemical resolution of distinct BRCA1-containing multiprotein complexes implicated in transcription and DNA repair.
- Novel discovery that BRCA1 mediates ligand-independent transcriptional repression of the estrogen receptor α.
- Novel discovery that BRCA1, in complex with RAD50/Mre11/NBS1, promotes nonhomologous end-joining of DNA double-strand breaks.
- Establishment of a BRCA1-dependent nonhomologous end-joining assay that will expedite studies designed to test the hypothesis that IR-induced site-specific phosphorylation of BRCA1 modulates its DNA double-strand break repair activities.
- Novel discovery within ZBRK1 of functionally bipartite zinc fingers with dual roles in sequence-specific DNA-binding and BRCA1-dependent transcriptional repression.
- Establishment of a BRCA1-dependent ZBRK1 transcriptional repression assay that will expedite studies designed to test the hypothesis that IR-induced site-specific phosphorylation of BRCA1 modulates its transcriptional regulatory activities.
- Novel discovery that ZBRK1, through its ability to homo-oligomerize, can function as both a sequence-specific DNA-binding transcriptional repressor and a DNA-binding independent transcriptional co-repressor.
• Discovery that the BRCA1-dependent ZBRK1 C-terminal repression domain (CTRD) is a novel protein interaction surface responsible for directing both homotypic and heterotypic interactions necessary for BRCA1-dependent ZBRK1 transcriptional repression.

Training Accomplishments:

Dr. Wei Tan, who previously worked on this project and authored two manuscripts in the process (please see Appendices 4 and 5 – manuscript reprints) recently completed his Ph.D. thesis and graduated from the UTHSCSA/Program in Molecular Medicine in May of 2005. Dr. Tan is currently pursuing his postdoctoral studies in the laboratory of Dr. Michael Karin in the Department of Pharmacology at the University of California, San Diego School of Medicine.

Currently, I mentor five Ph.D. students, two of whom are engaged in research pertaining to the studies described in this summary report. Ms. Amy M. Trauernicht is a fourth-year Ph.D. student who is working to biochemically fractionate BRCA1-containing transcription and DNA repair complexes from IR-treated cells using the purification strategy developed to achieve Task 1 of Technical Objective 1 and detailed in Appendix 1. Ms. Trauernicht is also following up on our observation that BRCA1 mediates ligand-independent transcriptional repression of the estrogen receptor. She has completed writing an invited review on this subject along with me, which was published in the journal Breast Disease (Please refer to Appendix 6 – Manuscript Reprint). She is making substantial progress along both of her research fronts.

Ms. Sejin Kim is a new graduate student who recently joined my laboratory on a permanent basis following successful completion of her first academic year in the Program in Molecular Medicine, which includes didactic training and compulsory laboratory rotations. Ms. Kim is a talented and hard-working student who will follow up on Dr. Tan’s work. More specifically, Ms. Kim will exploit our BRCA1-dependent ZBRK1 repression assay to evaluate the influence of IR-induced site-specific phosphorylation of BRCA1 on its sequence-specific co-repressor function.

REPORTABLE OUTCOMES TO DATE (Year 3 reportable outcomes underlined).

Manuscripts:


Reviews:


Meeting Abstracts:


Awards:

CONCLUSIONS

We have succeeded in the biochemical resolution of distinct BRCA1-containing multiprotein complexes implicated in transcription and DNA repair. We have made the novel discovery that BRCA1 mediates ligand-independent transcriptional repression of the estrogen receptor α; this finding suggests a possible means by which BRCA1 might control breast epithelial cell proliferation, and by implication cancer risk in the breast. We have also made the novel discovery that BRCA1, in complex with Rad50/Mre11/NBS1, promotes nonhomologous end-joining of DNA double-strand breaks. This observation has significant implications for the function of BRCA1 in tumor suppression through its role in the maintenance of genomic integrity. In addition to this novel finding, we have also established efficient and reliable BRCA1-dependent DNA repair- and transcription-based functional assays, the latter of which has permitted us to make the additional novel discoveries that: (1) ZBRK1, a BRCA1-dependent transcriptional repressor, harbors zinc fingers with dual roles in sequence-specific DNA-binding and BRCA1-dependent transcriptional repression, (2) ZBRK1, through its ability to homo-oligomerize, functions dually in both DNA-binding dependent and independent modes of transcriptional repression, and (3) the BRCA1-dependent ZBRK1 C-terminal repression domain (CTRD) is a novel protein interaction surface responsible for directing homotypic as well as heterotypic interactions necessary for ZBRK1-directed transcriptional repression. These findings shed new light on the mechanistic basis by which BRCA1 mediates sequence-specific control of DNA damage-responsive gene transcription. As long as resources permit, our current efforts will be targeted at identifying DNA damage-induced site-specific phosphorylation events with potential functional relevance to the role of BRCA1 in both transcription and DNA damage repair processes. Hence, we will exploit these BRCA1-dependent DNA repair- and transcription-based assay to analyze the effects of targeted BRCA1 mutations at identified sites of phosphorylation on its DNA repair and transcriptional regulatory activities. These studies should illuminate further the molecular basis for the caretaker properties of BRCA1.

REFERENCES

Biochemical Resolution of Distinct BRCA1-Containing Multiprotein Complexes Implicated in Transcription and DNA Repair

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Running Title: BRCA1 complexes implicated in transcription and DNA repair
BRCA1, a hereditary breast- and ovarian-specific tumor suppressor, functions in the maintenance of genome integrity and has been implicated in a diverse range of cellular processes including transcription regulation and DNA repair. However, the physical and functional relationship between BRCA1-containing activities involved in these processes remains to be fully deciphered. Here, we report the biochemical resolution of distinct multiprotein complexes comprised of BRCA1 in association with transcription and DNA repair activities. One complex, consisting of BRCA1, NBS1, Rad50, RNA polymerase II, and RNA polymerase II Mediator proteins could be resolved from a second complex comprised of BRCA1, NBS1, Rad50, Mre11, and additional polypeptides. These findings provide biochemical evidence for stable and distinct BRCA1-containing complexes with potential roles in transcription and DNA repair and, furthermore, provide evidence for an interaction of NBS1 and Rad50 with the RNA polymerase II holoenzyme. The presence of BRCA1, NBS1, and Rad50 in distinct complexes raises the possibility that these proteins represent a common core through which transcription and repair activities may be physically and functionally linked.

INTRODUCTION

Hereditary predisposition to early onset breast and ovarian cancer derives principally from germ-line mutations in either of two BReast CAnce susceptibility genes, BRCA1 and BRCA2 (1,2). Considerable evidence supports the notion that BRCA1 is a "caretaker" gene whose encoded product, a 220 kDa nuclear phosphoprotein, functions in the maintenance of
global genome stability (3-6). While the precise biochemical basis for its proposed caretaker function remains unknown, BRCA1 has nonetheless been implicated in both the regulation of transcription and the repair of damaged DNA.

Several lines of evidence support a direct role for BRCA1 in transcription control. First, the carboxyl-terminus of BRCA1 exhibits an inherent transactivation function sensitive to cancer-predisposing mutations (7-9). Second, BRCA1 has been identified as a component of the RNA polymerase II holoenzyme (10). Third, BRCA1 has been reported to interact with a variety of transcriptional activator and/or repressor proteins (11). Finally, BRCA1 activates transcription of genes that encode activities involved in DNA damage-induced cell cycle arrest and/or apoptosis. These include the cyclin-dependent kinase inhibitor p21 and the Growth Arrest and DNA Damage-inducible 45 (GADD45) genes that function in G1/S and/or G2/M checkpoint control, and the bax gene that functions in DNA damage-induced apoptosis (12-15). Collectively, these observations imply a role for BRCA1 in mediation of DNA damage-induced cell cycle arrest and/or apoptosis through control of gene transcription.

A significant body of experimental evidence also implicates BRCA1 in DNA damage repair. First, BRCA1 is known to undergo alterations in its phosphorylation status and subcellular localization in response to DNA damage (16). Second, brcal-deficient mouse embryonic cells are defective in the repair of both oxidative DNA damage by transcription-coupled processes and chromosomal double-strand breaks by homologous and nonhomologous recombination (17,18; Zhong, Q, Boyer, T., Chen, C.-F., Chen, P.-L., and Lee, W.-H. Manuscript submitted). Finally, BRCA1 interacts physically and functionally with the Rad50/Mre11/NBS1 protein complex that participates directly in the repair of DNA double-strand breaks (19).
Thus, while BRCA1 likely participates in the control of transcription and DNA double-strand break repair, at least in part, by virtue of its association with the RNA polymerase II holoenzyme and the Rad50/Mre11/NBS1 complex, respectively, the precise physical and functional relationship between these two sets of interacting proteins remains to be fully defined. With this issue in mind, we have undertaken the biochemical fractionation of human cell extracts to begin to decipher the protein networks through which BRCA1 functions. Our findings demonstrate that BRCA1, together with Rad50 and NBS1, can be isolated in distinct multiprotein complexes characterized by the stable association of these proteins with either transcription or repair activities. These results provide the first evidence for an association of Rad50 and NBS1 with the RNA polymerase II holoenzyme, and raise the possibility that these two proteins, along with BRCA1, represent a common core through which transcription and DNA repair activities may be linked within the cell.

RESULTS

Human HeLa cell nuclear extract was fractionated over Cibacron Hi-Trap Blue Sepharose using a linear gradient of KCl (0.1-1.5 M). Immunoblot analysis of the pre- and post-column extract revealed that the bulk of BRCA1, as well as Rad50, Mre11, NBS1, RNA polymerase II, and human Mediator proteins hSur2 and CDK8 (20) bound quantitatively to the Blue Sepharose matrix (Fig. 1; data not shown). Immunoblot analysis of individual chromatographic fractions revealed a broad elution profile for BRCA1, Rad50, and NBS1 (Fig. 1; data not shown). By contrast, hSur2 eluted early in the gradient, peaking at ~0.45 M KCl, while Mre11 eluted later, peaking at ~1.2 M KCl.
The presence of BRCA1, Rad50, and NBS1 in distinct chromatographic fractions corresponding to the peaks of a Mediator subunit on one hand (hSur2) and a double-strand break repair protein on the other (Mre11) led us to ask whether the three former proteins could be isolated in stable association with either of the latter two proteins. To address this question, Blue Sepharose fractions corresponding to the peaks of hSur2 (Fig 1; fractions 5-7) and Mre11 (Fig. 1; fractions 13-15) were pooled separately and subjected in parallel to further fractionation first by DEAE-Sepharose anion exchange and subsequently by Superose 6 gel filtration chromatography.

Immunoblot analysis revealed co-elution of hSur2, BRCA1, Rad50, and NBS1 during DEAE-Sepharose chromatography of Blue-Sepharose fractions 5-7 (Fig. 2a, lane 1 and data not shown). Superose 6 chromatography of peak DEAE-Sepharose fractions revealed co-elution of BRCA1, Rad50, and NBS1 in one major peak within the included volume (Fig. 2a; fractions 48-51), and in a second peak which corresponds to the excluded (void) volume of the Superose 6 column (Fig. 2a; fractions 39-42). The included and excluded Superose 6 peaks of BRCA1, Rad50, and NBS1 could be distinguished by the absence or presence of additional proteins. Specifically, the included peak, which eluted well ahead of the 670 kDa thyroglobulin marker, is characterized by the additional presence of RNA polymerase II holoenzyme components, including the RNA polymerase II large subunit, RPB1, and human Mediator proteins CDK8, Cyclin C, and Med7. Significantly, no Mre11 protein could be detected in these fractions. The excluded peak, by contrast, was characterized by the presence, in addition to BRCA1, Rad50, and NBS1, of RNA polymerase II and a substoichiometric level of Mre11; however, little or no hSur2, CDK8, Cyclin C or Med7 could be detected. The presence of Mre11 within the excluded peak likely derives from trace amounts of a BRCA1, Rad50, NBS1, and Mre11-containing complex incompletely resolved in the initial Blue Sepharose fractionation step. The ability of
Superose 6 to resolve this excluded peak containing Mre11 from an included peak of BRCA1, Rad50, and NBS1 in association with RNA polymerase II holoenzyme components raised the possibility that these two peaks represent stable and distinct multiprotein assemblies.

To determine if BRCA1, Rad50, and NBS1 all reside in a stable complex with RNA polymerase II holoenzyme components, individual Superose 6 column fractions corresponding to the included peaks of BRCA1, Rad50, NBS1, and holoenzyme components (Fig. 2a; fractions 48-51) were pooled and subjected to immunoprecipitation using an RNA polymerase II large subunit (RPB1)-specific monoclonal antibody, 8WG16 (21). Immunoblot analysis revealed specific co-immunoprecipitation of BRCA1, Rad50, and NBS1 along with RNA polymerase II and Mediator proteins CDK8, Cyclin C, and Med7 (Fig. 2b). This result demonstrates that these proteins all reside in a single, large molecular-size complex, which likely corresponds to the RNA polymerase II holoenzyme.

To begin to characterize the protein complex that contains BRCA1 in association with Mre11, Blue Sepharose fractions 13-15 were pooled and applied to a DEAE-Sepharose anion exchange resin. Immunoblot analysis of DEAE fractions revealed co-elution of BRCA1, Rad50, and NBS1 along with Mre11 in a 0.3M KCl step elution (data not shown). Superose 6 chromatography of peak DEAE-Sepharose fractions revealed co-elution of BRCA1, Rad50, NBS1, and Mre11 in one peak within the included volume (Fig. 3a; fractions 47-50), and in a second peak which corresponds to the excluded (void) volume of the Superose 6 column (Fig. 3a; fractions 40-43). The included and excluded Superose 6 peaks of BRCA1, Rad50, NBS1, and Mre11 could be distinguished by the absence or presence of additional proteins. Specifically, the excluded peak is characterized by the additional presence of small amounts of ATM and Rad51, although the bulk of ATM eluted in fractions corresponding to a molecular
size of its monomeric form (Fig. 3). We consider it likely that the excluded peak represents an insoluble protein aggregate, since BRCA1, Rad50, and NBS1 exhibit a propensity to precipitate from solution into aggregates that are excluded from Superose 6 (T.G. Boyer, unpublished data). Alternatively, the excluded peak could represent either an extremely large soluble protein complex or a protein/nucleic acid complex. Because these issues have not yet been thoroughly resolved, we have pursued analysis of Superose 6 fractions corresponding to the included peak of the BRCA1, Rad50, NBS1, and Mre11 proteins.

To determine if BRCA1, Rad50, NBS1, and Mre11 all reside in a stable complex, individual Superose 6 column fractions corresponding to the included peaks of the BRCA1, Rad50, NBS1, and Mre11 proteins (Fig 3a; fractions 47-50) were subjected to immunoprecipitation using a Rad50-specific monoclonal antibody, 13B3. Immunoblot analysis revealed specific co-immunoprecipitation of BRCA1, Rad50, and NBS1 along with Mre11 and at least 10 additional polypeptides (Fig. 3b). This result demonstrates that these proteins all reside in a single, large molecular-size complex.

**DISCUSSION**

We have undertaken the biochemical fractionation of human cell extracts in an initial effort to decipher the protein networks involved in BRCA1 function. Previous studies have implicated this tumor suppressor in both the control of transcription and the repair of damaged DNA (11). Consistent with these proposed functional roles, biochemical and protein interaction analyses have demonstrated direct and specific interaction of BRCA1 with both transcription and DNA repair activities. For example, it has been demonstrated that BRCA1 interacts individually
with the RNA polymerase II holoenzyme, with Rad51, and with the Rad50/Mre11/NBS1 DNA double-strand break repair complex (10,19,22). Thus, it appears likely that BRCA1 participates in a diverse range of DNA transactions by virtue of its association with these specific transcription and repair complexes. However, at present, the physical and functional relationship between these protein assemblies has remained undefined. We provide biochemical evidence to suggest that these activities, while perhaps linked within the cell, may nonetheless be isolated as distinct and stable macromolecular assemblies. Minimally, the proteins common to both of these identified complexes are BRCA1, Rad50, and NBS1.

The simultaneous presence of BRCA1, Rad50, and NBS1 in distinct multiprotein complexes with apparent transcription and repair functions provides a basis for the functional, and perhaps physical, linkage of these activities within the cell. We envision two alternative possibilities for the association of these two activities. First, these transcription and repair assemblies could represent components of a larger complex within the cell that has undergone fractionation in vitro (Fig 4a). Alternatively, these complexes could represent distinct assemblies in vivo that, by virtue of shared subunits, are linked functionally (Fig. 4b). For example, a dynamic redistribution of BRCA1, Rad50, and NBS1 among transcription and repair complexes could effect global alterations in these activities sufficient to meet the immediate physiological demands of the cell. The biochemical basis for such redistribution could involve phosphorylation, a notion consistent with observed alterations in the phosphorylation status and subcellular localization of BRCA1 as a consequence of cell cycle progression or cellular DNA damage (3,16). Detailed biochemical characterization of BRCA1 in association with the transcription and repair complexes identified herein should reveal whether post-translational modification represents a determinant of its interaction properties.
Our identification of both Rad50 and NBS1 in association with RNA polymerase II and transcriptional Mediator proteins represents, to our knowledge, the first demonstration of an interaction of these proteins with the RNA polymerase II holoenzyme. This observation suggests that these proteins may, like BRCA1, be dually involved in transcription and DNA repair. The precise role of these proteins in the control of transcription remains to be defined. However, such a role would not be entirely inconsistent with the observed pleiotropic features associated with an absence of NBS1 activity in Nijmegen breakage syndrome, which include microcephaly, growth and mental retardation, chromosomal instability, immunodeficiency, and a high incidence of hematopoietic malignancy (23). While NBS1 has, apart from its direct role in repair, been implicated in checkpoint control through regulation of ionizing radiation-induced p53 protein levels (24,25), a more direct role for NBS1 in control of gene transcription cannot be ruled out at present. Future analyses should serve to clarify whether and how NBS1 functions in association with the RNA polymerase II machinery to effect alterations in gene-specific transcription.

MATERIALS AND METHODS

Protein Purification - HeLa cell nuclear extract (~725 mg) was applied to Cibacron Hi-Trap Blue Sepharose (Amersham Pharmacia) at a concentration of 4.5 mg/ml protein (total of 7 X 5 ml columns; 103.5 mg protein/column) in 0.1M KCl D buffer (20). Columns were washed with four column volumes of 0.1M KCl D buffer and bound proteins subsequently eluted with a linear gradient of 0.1-1.5M KCl in D buffer over a total volume of 40 ml. Blue Sepharose fractions containing the peaks of hSur2 (fractions 5-7) and Mre11 (fractions 13-15) as determined by
immunoblot analysis were pooled separately, dialyzed into 0.1M KCl D buffer, and processed in parallel as follows. Dialyzed Blue Sepharose fractions were applied to DEAE-Sepharose (10 mg protein/ml of resin) in 0.1M KCl D buffer. Columns were washed with four column volumes of 0.1M KCl D buffer and step-eluted with 0.3M KCl D buffer. Individual DEAE-Sepharose fractions containing the peaks of BRCA1, Rad50, and NBS1 along with either hSur2 or Mre11 proteins as determined by immunoblot analysis were pooled to a final concentration of 4 mg/ml and subjected to Superose 6 gel filtration chromatography (2 ml per 16 X 500 mm column). Individual Superose 6 column fractions were analyzed by immunoblot analysis and fractions corresponding to selected peaks as indicated were pooled, concentrated on phosphocellulose P-11 using a 0.6M KCl step elution, and subjected to immunoprecipitation analyses.

**Antibody Immunoprecipitation** - Monoclonal antibodies specific for the RNA polymerase II large subunit CTD (8WG16; ref. 21), human Rad50 (13B3; ref. 19), human p53 (PAb421; ref. 26), and Glutathione S-Transferase (8G11; ref. 19) were individually covalently coupled to protein G-Sepharose using dimethylpimelimidate (27). Superose 6 column fractions containing peaks of BRCA1, Rad50, RNA polymerase II, and Mediator proteins were pooled and incubated in parallel with either 8WG16 (specific) or PAb421 (non-specific control) antibody columns in 0.3 M KCl (1/2) D buffer [(1/2) D buffer is 20 mM HEPES, pH 7.9; 0.2 mM EDTA; 10% glycerol; 5 mM β-mercaptoethanol] for 6 hours at 4° C. Superose 6 column fractions containing peaks of BRCA1, Rad50, NBS1, and Mre11 were pooled and incubated in parallel with either 13B3 (specific) or 8G11 (non-specific control) antibody columns in 0.3 M KCl (1/2) D buffer for 6 hours at hours 4° C. Column matrices were washed three times with ten column volumes of 0.3 M KCl (1/2) D buffer, once with ten column volumes of 0.1M KCl (1/2) D buffer, and eluted
with one column volume of 0.1 M glycine (pH 2.0). Column eluates were neutralized, subjected to SDS-10%PAGE, and characterized by silver stain or immunoblot analysis as indicated.

ACKNOWLEDGEMENTS

We thank P.-L. Chen, D. Jones, and P. Garza for antibodies and technical assistance, and P.R. Yew, P.-L. Chen, P. Sung, Y. Chen, and N. Ting for advice and comments. This work was supported by NIH grants P01CA30195 and P01CA81020 and the McDermott Endowment fund.

REFERENCES


**FIGURE LEGENDS**

**Figure 1. BRCA1 and Rad50 co-elute from Cibacron Blue Sepharose with both transcriptional Mediator and DNA repair proteins.** HeLa nuclear extract was applied to a Hi-Trap Blue column at 0.1M KCl, and bound proteins were eluted with a linear gradient of 0.1-1.5M KCl. Aliquots of the on-put nuclear extract (NEXT), the column flow-through (BS FT), and individual chromatographic fractions (numbered) were analyzed by immunoblot analysis using antibodies specific for the proteins indicated on the left of the blot. Fractions corresponding to the peaks of hSur2 (fractions 5-7) and Mre11 (fractions 13-15) were pooled separately and processed in parallel for further chromatographic analyses as indicated.

**Figure 2. BRCA1, Rad50, and NBS1 reside in stable association with RNA polymerase II holoenzyme components.** a. Superose 6 gel filtration profile of Blue Sepharose and DEAE-Sepharose fractionated proteins. Pooled Blue Sepharose fractions 5-7 (from Fig.1) were applied to DEAE-Sepharose and peak fractions from a 0.3M KCl step elution containing BRCA1, Rad50, and NBS1 were pooled and subjected to gel filtration on Superose 6. Aliquots of the on-
put DEAE fraction (Load) and individual column fractions (numbered) were analyzed by immunoblot analysis using antibodies specific for the proteins indicated on the left. Downward-pointing arrows indicate the positions of marker protein peaks. Fractions corresponding to the excluded (void) and included volume of the Superose 6 column are indicated. b. Co-immunoprecipitation of BRCA1 and NBS1 with RNA polymerase II holoenzyme components. Superose 6 fractions 48-51 (from a) were pooled, concentrated on phosphocellulose using a 0.6M KCl step elution (lanes 2 and 5) and subjected to immunoprecipitation with an RNA polymerase II large subunit (RPB1)-specific monoclonal antibody 8WG16 (lanes 4 and 7), or a p53-specific monoclonal antibody PAb421 (lanes 3 and 6) as a negative control. Immunoprecipitated proteins were eluted from covalently-coupled antibody columns with glycine, subjected to SDS-10%PAGE, and processed either by silver staining (lanes 1-4) or immunoblot analysis (lanes 5-7) with antibodies specific for the proteins indicated on the right. Molecular weight markers are indicated on the left. Lane 1 represents highly purified core RNA polymerase II, only the two largest subunits of which stained visibly on this gel.

Figure 3. BRCA1, Rad50, and NBS1 reside in stable association with Mre11 and additional polypeptides. a. Superose 6 gel filtration chromatography of Blue Sepharose and DEAE-Sepharose fractionated proteins. Pooled Blue Sepharose fractions 13-15 (from Fig. 1) were applied to DEAE-Sepharose and peak fractions from a 0.3M KCl step elution containing BRCA1, Rad50, NBS1, and Mre11 were pooled and subjected to gel filtration on Superose 6. Aliquots of individual column fractions (numbered) were analyzed by immunoblot analysis using antibodies specific for the proteins indicated on the left. Downward-pointing arrows indicate the positions of marker protein peaks. Fractions corresponding to the excluded (void)
and included volume of the Superose 6 column are indicated. b. Co-immunoprecipitation of BRCA1, Rad50, and Mre11 along with additional polypeptides. Superose 6 fractions 47-50 (from a) were pooled, concentrated on phosphocellulose using a 0.5M KCl step elution (lanes 1 and 4), and subjected to immunoprecipitation with a Rad50-specific monoclonal antibody 13B3 (lanes 2 and 6), or a GST-specific monoclonal antibody 8G11 (lanes 3 and 5) as a negative control. Immunoprecipitated proteins were eluted from covalently-coupled antibody columns with glycine, subjected to SDS-10%PAGE, and processed either by silver staining (lanes 1-3) or immunoblot analysis (lanes 4-6) with the antibodies specific for the proteins indicated on the right. Molecular weight markers are indicated on the left. The ~95 kDa protein specifically co-immunoprecipitated with anti-Rad50 antibody has been confirmed by immunoblot analysis to be NBS1, as indicated by the elongated arrow.

Figure 4. Schematic models for the association of BRCA1, Rad50, and NBS1 with transcription and DNA repair complexes. (a) The RNA polymerase II holoenzyme and the Rad50/Mre11/NBS1 DNA double-strand break repair complexes are physically linked in vivo to from a larger complex that undergoes fractionation in vitro to yield stable and distinct multiprotein assemblies. This model would imply other than a 1:1 stoichiometric ratio of BRCA1, Rad50, and NBS1 relative to other proteins within the larger complex. Presently, the stoichiometry of these proteins in complex with isolated transcription and repair activities has not been elucidated. (b) The RNA polymerase II holoenzyme and the Rad50/Mre11/NBS1 complexes exist in vivo as distinct and stable multiprotein assemblies that may be linked functionally by virtue of shared subunits. Shown here is a potential dynamic redistribution of BRCA1, Rad50, and NBS1 between transcription and repair complexes. Cell-cycle and/or DNA
damage-induced phosphorylation could represent the signal that specifies the association of these proteins with Mre11 and additional DNA double-strand break repair activities. The specific cell signal(s) that directs BRCA1, Rad50, and NBS1 to the RNA polymerase II holoenzyme is unknown.
Fig. 1

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BRCA1 →

Rad50 →

hSur-2 →

Mre11 →

Cdk8 →

5-7 ↓

DEAE Sepharose

13-15 ↓

DEAE Sepharose

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Superose 6
Fig. 2

a

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Excluded column volume | Included column volume

1 2 3 4 5 6 7

Ag Stain IP/Western

BRCA1 hSur-2 NBS1 Cdk8 Med7 Cyclin C
Fig. 3

(a) 

Mr (k): 670 158 44
37 40 43 46 49 52 55 58 61 64 67 70 73 76 79 82 85

ATM
BRCA1
Rad50
NBS1
Mre11
Rad 51

Excluded column volume
Included column volume

(b)

Mr (k)

Sup 6: 47-50 anti-Rad50 anti-GST

200
116
97.4
66
45
31

Sup 6: 47-50 anti-Rad50

Ag Stain IP/Western

BRCA1
Rad50
NBS1
Mre11
BRCA1 mediates ligand-independent transcriptional repression of the estrogen receptor

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Mutational inactivation of BRCA1 confers a cumulative lifetime risk of breast and ovarian cancers. However, the underlying basis for the tissue-restricted tumor-suppressive properties of BRCA1 remains poorly defined. Here we show that BRCA1 mediates ligand-independent transcriptional repression of the estrogen receptor α (ERα), a principal determinant of the growth, differentiation, and normal functional status of breasts and ovaries. In Brca1-null mouse embryo fibroblasts and BRCA1-deficient human ovarian cancer cells, ERα exhibited ligand-independent transcriptional activity that was not observed in Brca1-proficient cells. Ectopic expression in Brca1-deficient cells of wild-type BRCA1, but not clinically validated BRCA1 missense mutants, restored ligand-independent repression of ERα in a manner dependent upon apparent histone deacetylase activity. In estrogen-dependent human breast cancer cells, chromatin immunoprecipitation analysis revealed the association of BRCA1 with ERα at endogenous estrogen-response elements before, but not after estrogen stimulation. Collectively, these results reveal BRCA1 to be a ligand-reversible barrier to transcriptional activation by unliganded promoter-bound ERα and suggest a possible mechanism by which functional inactivation of BRCA1 could promote tumorigenesis through inappropriate hormonal regulation of mammary and ovarian epithelial cell proliferation.

Germline inactivation of the gene that encodes BRCA1 represents a predisposing genetic factor in ~15-45% of hereditary breast cancers, and minimally 80% of combined hereditary breast and ovarian cancer cases (1). Functionally, BRCA1 has been implicated in the maintenance of global genome stability (2-4), and the underlying basis for this activity likely derives from its central role in the cellular response to DNA damage, wherein it controls both DNA damage repair and the transcription of DNA damage-inducible genes (5-14).

Because the DNA damage-induced signaling pathways that converge on BRCA1 are likely to be conserved in most cell types, BRCA1 is likely to occupy a fundamental and universally conserved role in the mammalian DNA damage response. Nonetheless, germ-line inactivation of BRCA1 leads predominant-ly to cancer of the breast and ovary, and the underlying basis for its tissue-restricted tumor-suppressive properties thus remains undefined.

At least two hypotheses have been proposed to explain the tissue-specific nature of BRCA1-mediated tumor suppression, both of which invoke a role for estrogen in either the initiation or promotion of tumor formation (15). According to one model, the tissue-specific tumor-suppressive properties of BRCA1 derive, at least in part, from its response to tissue-specific DNA damage. In this regard, certain oxidative metabolites of estrogen are likely to promote tumorigenesis through inappropriate hormonal regulation of mammary and ovarian epithelial cell proliferation.

Materials and Methods

Cell Culture. p53−/− (Brca1+/+) and p53−/− (Brca1−/−) mouse embryonic fibroblasts (MEFs) were cultured as described (14). Human MCF7 cells were maintained in DMEM supplemented with 10% FCS. Human BG-1-derived NEO1 and AS4 cell lines were maintained as described (22). Depletion of hormone ligands for nuclear/steroid receptor activation studies was achieved by cell culture in medium containing either 10% charcoal/dextran-treated serum (HyClone) or defined serum replacement 2 (Sigma).

Plasmids and Transfections. Transfection assays were performed by using the following conditions. Reporter plasmids. Used at 0.5 μg each, including pTRE(F2)-TK-Luc, pGRE-TK-CAT, pERE-TK-Luc, or pPRE-TK-CAT (23); 0.5 μg of pGAL4-SV40-Luc containing five GAL4 DNA-binding sites upstream of the minimal simian virus 40 (SV40) promoter, driving expression of the luciferase reporter gene in the pGL2 vector (Promega); and 0.5 μg of pGAL4-E1B-Luc (24).

Receptor expression plasmids. Used at 1.0 μg each, including RSV-hTRβ, RSV-hGR, RSV-ERα, and RSV-hPRβ (23).

BRCA1 expression plasmids. Used at 1.0 μg each, including pcDNA3.1-BRCA1, pcDNA3.1-BRCA1-A1708E, pcDNA3.1-BRCA1-Q356R, and pcDNA3.1-BRCA1-A1708E/Q356R expressing either human wild-type BRCA1 or familial breast cancer-derived BRCA1 mutants (14).

Abbreviations: ERα, estrogen receptor α; MEF, mouse embryonic fibroblast; E2, 17β-estradiol; RT-PCR, reverse transcription-PCR; HDAC, histone deacetylase; CNI, chromatin immunoprecipitation; AF-1, N-terminal ligand-independent activation function; AF-2, C-terminal ligand-inducible activation function.†To whom reprint requests may be addressed. E-mail: leew@uthscsa.edu or boyer@uthscsa.edu.

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Chimeric activators. Used at 1.0 μg of GAL4-ERα, generated by an amino-terminal fusion of ERα with the GAL4 DNA-binding domain in pM5 (25); 0.1 μg of pVP16-GAL4 or pVP16-GAL-ERα containing ERα amino acids 251–395, as described (26). MEFs (6 × 10^5) or BG-1 cells (2 × 10^5) cultured in ligand-free medium were transfected by Lipofectin-based methods under serum-free conditions. Culture medium was replaced with fresh ligand-free medium 24 h after transfection, and 10^-7 M 17β-estradiol (E2) or 330 nM trichostatin A was added as indicated. Cells were harvested 48 h after transfection for luciferase assay as described (14) or chloramphenicol acetyltransferase (CAT) assay by liquid scintillation counting (Promega).

Reverse Transcription (RT)-PCR Analysis. BG-1-derived cells were cultured in ligand-free medium for at least 5 days, and treated with 10^-7 M E2 for 1 h as indicated. Approximately 15 μg of total cellular RNA was subjected to semiquantitative RT-PCR analysis following a procedure previously described for estrogen-responsive genes (27, 28).

Chromatin Immunoprecipitation (ChIP). MCF7 cells were cultured in ligand-free medium for at least 5 days and treated with 10^-7 E2 for 1 h as indicated. ChIP assays were performed as described (29).

Antibodies. Antibodies used for soluble and chromatin immunoprecipitations and immunoblot analyses were as follows: BRCA1 (mAb 6B4); ERα (rabbit polyclonal antibody HC-20 or mouse mAb D-12, Santa Cruz Biotechnology); ChIP (mAb 19E8); TFIIF p89 (rabbit polyclonal antibody S-19, Santa Cruz Biotechnology); glutathione S-transferase (MAb 8G11); RNA poly-merase II large subunit (mAb 8WG16); cathepsin D (rabbit polyclonal antibody 06-467, Upstate Biotechnology, Lake Placid, NY); p52 (mouse mAb V3030, Biomedea, Hayward, CA); human progesterone receptor β (mouse mAb PriB-30, Santa Cruz Biotechnology); p84 (mAb 5E10).

Results

BRCA1 has been shown to regulate the ligand-dependent transcriptional activity of specific members of the nuclear hormone receptor family (17–20). However, endogenous BRCA1 present in the transfected cell lines used in previous studies precluded analysis of the effect of BRCA1 on the ligand-independent function of these receptors. Therefore, to more directly assess the role of BRCA1 in nuclear receptor transactivation without competition from endogenous BRCA1, we analyzed a panel of nuclear receptors for their respective ligand-independent transcriptional activities in Brca1-nullizygous MEFs.

A set of minimal thymidine kinase (TK) promoters, each under control of distinct hormone-response elements specific for either the human thyroid receptor β (TRβ), the glucocorticoid receptor (GR), the ERα, or the progesterone receptor β (PRβ) were individually tested for their respective abilities to direct expression of a reporter gene in the absence or presence of each corresponding receptor (absent ligand) after transfection into Brca1-proficient (Brca1+/+) or Brca1-deficient (Brca1−/−) MEFs (14). Unexpectedly, we observed significant ligand-independent activation of reporter gene expression directed by both the progesterone receptor β and the ERα in Brca1-deficient MEFs compared with Brca1-proficient MEFs (Fig. 1A). By contrast, no ligand-independent stimulation of reporter activity directed by either the thyroid receptor β or the glucocorticoid receptor β could be observed in Brca1-deficient MEFs (Fig. 1A). Interestingly, although E2 activated the ERα in both Brca1-proficient and Brca1-deficient MEFs, the relative level of induction observed in Brca1-deficient MEFs was diminished 2-fold relative to Brca1-proficient MEFs (Fig. 1B). We confirmed by immunoblot analysis that the transfected ERα was expressed equivalently in BRCA1-proficient and BRCA1-deficient MEFs, thus excluding the possibility that differences in receptor activity derive from differences in receptor protein expression (Fig. 1C).

Ectopic expression of wild-type BRCA1 in Brca1-deficient MEFs repressed ligand-independent activation directed by ERα (Fig. 2A). Likewise, a BRCA1 derivative carrying a familial breast cancer-derived missense mutation in the ring finger (C64G), also repressed ligand-independent activation by ERα (Fig. 2A). By contrast, BRCA1 derivatives carrying familial breast cancer-derived missense mutations in either an exon 11-encoded region that binds Rad50 and the transcriptional repressor ZBRK1 (Q536R) or the C-terminal BRCT domain (A1708E) abolished the ability of BRCA1 to repress ligand-independent thyroid hormone activation directed by ERα (Fig. 2A, 2B). Differences in the transcriptional repression activities of the various BRCA1 mutant derivatives could not be attributed to differences in their respective levels of expression because each of the BRCA1 mutant derivatives was expressed at a level comparable...
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 cells were harvested and analyzed by semiquantitative RT-PCR. Furthermore, although the addition of E2 elevated the steady-

 gene for 5 days followed by the addition of either no cathepsin D, and progesterone receptor proteins could be ob-

 individual monolayer culture of an internal control protein (nuclear matrix protein p84), matrix protein p84 (Bottom) indicates that nearly equivalent amounts of each cell lysate were used in the immunoprecipitations.

t to wild-type BRCA1 (Fig. 2C). BRCA1-mediated, ligand-

 independent repression of ERα was largely reversed by tricho-

 statin A, implicating histone deacetylase (HDAC) activity in this process (Fig. 2B). Collectively, these results reveal a function for BRCA1 as a repressor of ligand-independent, ERα-mediated transactivation.

 To confirm these results in a biologically relevant cell type, we analyzed the ligand-independent activity of ERα in human ovarian adenocarcinoma BG-1 cells, which are ERα-positive and estrogen-dependent for growth (30). Previously, Annab et al. (22) described the generation of independent BG-1 clonal cell lines that support stably reduced BRCA1 mRNA and protein levels by retroviral-mediated BRCA1 antisense delivery. We tested the ability of ERα to direct ligand-independent transcription of the ERE-TK-Luc reporter gene after transfection into either a control retroviral vector-infected BG-1 clonal cell line (NEO1) or, alternatively, a BRCA1 antisense-infected BG-1 clonal cell line (AS4) exhibiting severely reduced BRCA1 expression levels (Fig. 3E; ref. 22). Consistent with the results obtained in MEF cells, ERα exhibited significantly increased ligand-independent activity in BRCA1-deficient AS4 cells compared with BRCA1-proficient NEO1 cells (Fig. 3A). We also observed a 2-fold reduction in the relative level of E2-mediated induction of reporter gene activity in AS4 cells compared with NEO1 cells, once again consistent with the results obtained in MEF cells (Fig. 3B). These results confirm that in a biologically relevant epithelial cell type, BRCA1 can mediate repression of ligand-independent ERα transactivation activity.

 To determine whether the reduced BRCA1 expression levels in AS4 cells could be correlated with an increase in the ligand-independent expression of estrogen-responsive genes, we performed a direct comparative analysis of NEO1 and AS4 cells with respect to their ligand-independent expression of several estrogen-responsive genes. Individual monolayer cultures of NEO1 and AS4 cells were grown in the absence of estrogen for 5 days followed by the addition of either no hormone or, alternatively, E2 (10⁻⁷ M) for 1 h. Subsequently, cells were harvested and analyzed by semiquantitative RT-PCR for the expression levels of the endogenous estrogen-responsive pS2, cathepsin D, and progesterone receptor genes.

 Relative to the expression level of an internal control ribosomal S16 gene, we observed increases in the ligand-independent expression levels of the pS2, cathepsin D, and progesterone receptor genes of 3-, 5-, and 9-fold, respectively, in BRCA1-deficient AS4 cells compared with BRCA1-proficient NEO1 cells (Fig. 3C). Interestingly, although the addition of E2 stimulated transcription of the pS2, cathepsin D, and the progesterone receptor genes in NEO1 cells, no such E2-dependent increase in the transcription of these genes could be observed in AS4 cells (Fig. 3C). Qualitatively similar results were observed at the protein level by immunoblot analysis. Relative to the level of an internal control protein (nuclear matrix protein p84), E2-independent increases in the steady-state levels of the pS2, cathepsin D, and progesterone receptor proteins could be observed in AS4 cells compared with NEO1 cells (Fig. 3D). Furthermore, although the addition of E2 elevated the steady-

Fig. 2. Ectopic expression of wild-type BRCA1 in Brca1-deficient MEFs restores ligand-independent repression of ERα transactivation in a histone deacetylase (HDAC)-dependent manner. (A and B) Brca1−/− MEFs in estrogen-free media were transfected with pERE-TK-Luc without (−) or with (+) pRSV-ERα, pCDNA3.1-BRCA1 expressing wild-type human BRCA1 (WT), or pCDNA3.1-BRCA1 derivatives bearing missense mutants A1708E, Q356R, A1708E/Q356R, or C64G before assay for luciferase activity. Where indicated, trichostatin A (TSA; 330 nM) was also included. (C) Brca1−/− MEFs in estrogen-free media were untransfected (lane 1) or cotransfected with expression vectors for ERα and either wild-type BRCA1 (lanes 2 or various BRCA1 mutant derivatives (lanes 3–6) as indicated. Cells were lysed, and immunoprecipitated BRCA1 and ERα were subjected to immunoblot analysis using antibodies specific for BRCA1 (Top) or ERα (Middle). Immunoblot analysis of the nuclear matrix protein p84 (Bottom) indicates that nearly equivalent amounts of each cell lysate were used in the immunoprecipitations.

Fig. 3. Reduced BRCA1 expression in BG-1 human ovarian adenocarcinoma cells is accompanied by increases in estrogen-independent expression of estrogen-responsive genes. (A) Retrosplenicteron-injected (NEO1) and BRCA1 antisense-infected (AS4) BG-1 clonal cell lines were transfected with pERE-TK-Luc without (−) or with (+) pRSV-ERα before assay for luciferase activity. (B) NEO1 and AS4 cells in estrogen-free media were transfected with pERE-TK-Luc with (+) pRSV-ERα in the absence (−) or presence (+) of E2 (10⁻⁷ M) before assay for luciferase activity. (C) NEO1 (lanes 1 and 3) or AS4 (lanes 2 and 4) cells in estrogen-free media were either untreated (lanes 1 and 2) or treated (lanes 3 and 4) with E2 (10⁻⁷ M) for 24 h. Culture medium was concentrated 10-fold by using a Centriprep YM-3 device, and 1/10th of the concentrate was resolved by SDS/10%PAGE and processed for immunoblot analysis using antibodies specific for pS2. Cells were also lysed in RIPA buffer, and 1/10th of the lysate was subjected to immunoblot analysis using antibodies specific for progesterone receptor β (PR), cathepsin D (Cat D), or nuclear matrix protein p84, which served as an internal loading control. (D) Whole cell lysates derived from NEO1 and AS4 cells were resolved by SDS/10%PAGE and processed for immunoblot analysis using antibodies specific for BRCA1, CIP, and the p89 subunit of the transcription factor BH (TFIH), the latter two of which served as internal loading controls. The ERα-positive status of these cells was verified by using an ERα-specific rabbit polyclonal antibody. Densitometric quantitation of the immunoblot and normalization to the CIP and TFIH signals revealed BRCA1 expression to be reduced by 70% in AS4 cells compared to NEO1 cells.
state level of each of these proteins in NEO1 cells, no such E2-dependent increase could be observed in AS4 cells (Fig. 3D). Quantitative differences between RT-PCR and immunoblot analyses could reflect the influence of posttranscriptional regulatory processes. Nonetheless, RT-PCR and immunoblot analyses both reveal that the ligand-independent expression of endogenous ERα-target genes is increased in BRCA1-deficient cells. Collectively, these results implicate BRCA1 in the ligand-independent repression of endogenous estrogen-responsive genes.

To explore the mechanism by which BRCA1 mediates ligand-independent repression of ERα, we first determined whether BRCA1 could interact with unliganded ERα in vivo by coimmunoprecipitation of the two proteins in human breast cancer MCF7 cells cultured in the absence of estrogen. Consistent with previous results (18), BRCA1 could be specifically coimmunoprecipitated with unliganded ERα, thus demonstrating that the two proteins can interact in vivo in a ligand-independent manner (data not shown).

To explore the possibility that BRCA1 represses the transcription function of promoter-bound, unliganded ERα, we first tested the effect of BRCA1 on the ligand-independent transcriptional activity of ERα tethered to the yeast GAL4 DNA-binding domain by using a reporter template bearing GAL4 DNA-binding sites. This approach permitted us to assess the effect of BRCA1 on the transcriptional function of unliganded ERα independent of any effects that BRCA1 might have on the DNA-binding activity of unliganded ERα. GAL4-ERα was cotransfected along with a GAL4-SV40-luciferase reporter template into BRca1-proficient and BRca1-deficient MEFs. We observed significant ligand-independent stimulation of reporter activity in BRca1-deficient, but not in BRca1-proficient, MEFs (Fig. 4A), suggesting one mechanism by which BRCA1 mediates ligand-independent repression of ERα is through direct repression of the DNA-bound receptor.

To confirm this observation under biologically relevant conditions in vivo, we used ChIP analyses to determine whether BRCA1 can be recruited directly to estrogen-responsive promoters in the absence of ligand. MCF-7 cells were grown in the absence of estrogen for 5 days followed by the addition of either no hormone or, alternatively, E2 (10^-7 M) for 1 h. Promoter occupancy before and after E2 treatment at the estrogen response elements within the endogenous pS2 and cathepsin D gene promoters by ERα, BRCA1, and RNA polymerase II was then monitored by ChIP using antibodies specific for each of the three proteins and semiquantitative PCR with primers flanking the estrogen response elements of the pS2 and cathepsin D promoters. In the absence of E2, ERα could be detected in association with both the pS2 and cathepsin D promoters, and this level was increased dramatically by the addition of E2 (Fig. 4B, lanes 3 and 7). By contrast, RNA polymerase II could be detected only following, but not before, E2 treatment, consistent with its ligand-dependent recruitment concomitant with transcriptional activation of the pS2 and cathepsin D genes (Fig. 4B, lanes 4 and 8 and C, lanes 1 and 2). The specificity of factor association within the estrogen-responsive region of the pS2 and cathepsin D promoters was confirmed by ChIP analysis using antibodies specific for ZBRK1, a sequence-specific DNA-binding transcriptional repressor that does not bind to pS2 or cathepsin D promoter sequences (14). ZBRK1-specific antibodies failed to immunoprecipitate pS2 and cathepsin D promoter sequences (data not shown). Further specificity of the ChIP assay was demonstrated by the inability to detect occupancy by ERα, BRCA1, or RNA polymerase II of a region ~3 kb upstream of the cathepsin D promoter (Fig. 4B). These results thus reveal the association of BRCA1 with unliganded ERα at endogenous estrogen-responsive promoters under physiologically relevant conditions in vivo.

Like other steroid receptors, ERα contains a N-terminal ligand-independent activation function (AF-1) that is targeted by a variety of steroid-independent cell-signaling pathways, and a C-terminal ligand-Inducible activation function (AF-2) that resides within the receptor ligand-binding domain (31, 32). Previous analyses of ERα suggest a model whereby repressive factors binding to sequences within its C-terminal ligand-binding domain repress constitutively active AF-1 in the absence of an agonist or in the presence of an antagonist (26, 33). To determine whether ligand-independent repression of ERα by BRCA1 is mediated through the ERα ligand-binding domain, we tested the ligand-independent activity of a VP16-GAL4-ERα receptor chimera after its expression in both BRCA1-proficient and BRCA1-deficient BG-1 clonal cell lines. This chimera encodes ERα amino acids 251-595, including the hinge region and the ligand-binding domain, fused C-terminally to the hybrid transactivator VP16-GAL4 (26).
Previously, deletion analysis of this receptor chimera revealed that constitutive VP16-GAL4-ERα activity could be recovered by the removal of sequences within the ligand-binding domain of the ERα moiety, thereby implicating the ERα ligand-binding domain in ligand-independent transcriptional repression of a neighboring constitutive activation domain (26). To determine whether this ligand-independent repression is mediated by BRCA1, we transfected the VP16-GAL4-ERα chimera along with a reporter template bearing GAL4 DNA binding sites into both BRCA1-proficient NEO1 cells and BRCA1-deficient AS4 cells. In NEO1 cells, the VP16-GAL4-ERα chimera exhibited minimal constitutive transactivation activity in the absence of E2; in response to E2, this level was dramatically increased to one approaching that of the potent VP16-GAL4 activator alone (Fig. 5 A and B). By contrast, in AS4 cells the VP16-GAL4-ERα chimera exhibited constitutive transactivation activity comparable to that exhibited by the VP16-GAL4 activator alone (Fig. 5C). The addition of E2 had a minimal effect on the elevated constitutive transactivation activity of the ERα chimera in AS4 cells (data not shown), suggesting that the principle effect of E2 is to override a ligand-independent barrier to the transactivation activity of the chimeric receptor. This barrier is present in NEO1 cells, but deficient in AS4 cells. Similar results were also observed by using isogenic Brcal-proficient and Brcal-deficient MEFs, eliminating the possibility that cell type-specific peculiarities contributed to the differential transactivation properties of the VP16-GAL4-ERα chimera in the presence and absence of BRCA1 (data not shown). Collectively, these results reveal the ERα ligand-binding domain to be a platform for the recruitment of BRCA1 from which the latter may confer ligand-independent repression on a linked activation domain. Hence, we conclude that BRCA1-mediated ligand-independent repression of ERα is likely to be mediated through the ERα ligand-binding domain.

Discussion

Recently, BRCA1 has been proposed to inhibit the ligand-dependent transcriptional activity of ERα through a direct interaction between the two proteins (18). Our current analysis of ERα transcriptional activity in Brcal-nullzygous MEFs revealed BRCA1 to be a ligand-reversible barrier to transcriptional activation by unliganded ERα. The biological relevance of this finding is further strengthened by the observation that BRCA1 also mediates ligand-independent repression of the ERα in human ovarian adenocarcinoma cells.

The underlying mechanism by which BRCA1 mediates ligand-independent repression of ERα transcriptional activity appears to involve targeted recruitment by unliganded, promoter-bound ERα of a BRCA1-associated HDAC activity. This conclusion is based first on the observation that the HDAC inhibitor trichostatin A can effectively reverse ligand-independent repression mediated by BRCA1 and, second, on the results of ChIP analyses, which revealed the association of unliganded ERα with BRCA1 on endogenous estrogen-response elements in vivo. A likely target of BRCA1-mediated ligand-independent ERα repression is the constitutive AF-1 activation domain within ERα. Previous studies have indicated that antagonist-bound AF-2 can repress AF-1 activity through the recruitment of the nuclear corepressor N-CoR (33), whereas the ligand-binding domain of unliganded ERα can repress a linked heterologous activation domain in a ligand-reversible manner, presumably by the recruitment of a soluble corepressor (26). Our observation that an estrogen-dependent VP16-GAL4 chimeric transactivator carrying the ERα ligand-binding domain exhibits constitutive activity in BRCA1-deficient, but not in BRCA1-proficient BG-1 cells, reveals the ERα ligand-binding domain to be a potential site of BRCA1 recruitment for ligand-independent repression of a linked activation domain. Hence, BRCA1 could be recruited to the ERα ligand-binding domain as part of a larger repression complex to silence AF-1 function in the absence of ligand. The recent report of a direct interaction between BRCA1 and the ERα ligand-binding domain (18) lends additional support to this model.

Should BRCA1 function to inhibit the ligand-dependent transcriptional activity of ERα (17, 18), it seems unlikely to do so through a mechanism that involves promoter-bound ERα. Our ChIP analysis revealed the association of BRCA1 with ERα at endogenous estrogen-response elements before, but not after, estrogen stimulation. Thus, we favor a model in which BRCA1, along with an associated corepressor(s) that minimally includes an HDAC activity, is recruited by unliganded, promoter-bound ERα to effectively silence the constitutive AF-1 activation domain and thereby repress estrogen-responsive target gene transcription. After estrogen stimulation, a ligand-induced conformational change within ERα could lead to enhanced affinity of the ERα for its cognate binding site and release of a BRCA1-containing repression complex, thereby liberating AF-1 and AF-2 to synergistically recruit coactivators and the RNA polymerase II holoenzyme to promote transcription (29). It is also possible that BRCA1 could function additionally as a barrier to the productive association of either unliganded and/or liganded ERα with promoter DNA, and this could underlie the previous observation that BRCA1 can inhibit ligand-dependent ERα transactivation (17, 18).

Interestingly, we observed that a deficiency of BRCA1 also leads to a reduction in the relative level of E2-mediated ERα activation. In both Brcal-nullzygous MEFs and Brcal-deficient BG-1 (AS4) cells, the relative level of E2-mediated activation of a transfected ERα-responsive reporter gene was diminished when compared with Brcal-proficient cells. Furthermore, in AS4 cells, the endogenous estrogen-response genes that we monitored exhibited increased estrogen-independent expression and little or no estrogen-dependent stimulation when compared with Brcal-proficient BG-1 (NEO1) cells. It is possible that the expression of these genes is largely derepressed in a Brcal-deficient background and cannot therefore be increased substantially in response to estrogen.

Previously, Annab et al. (22) demonstrated that relative to parental or retroviral vector-infected BG-1 cell clones, BRCA1 antisense-infected BG-1 cell clones exhibit enhanced estrogen-independent growth in culture (22). Furthermore, BG-1 clone AS4, which exhibits severely reduced BRCA1 expression levels, exhibited increased tumorigenicity in ovariectomized nude mice compared with the retroviral vector-infected NEO1 cell clone (22). These observations suggest that forced reduction of
BRCA1 in BG-1 ovarian adenocarcinoma cells may influence estrogen-independent growth both in vitro and in vivo. Our observation that AS4 cells support significant increases in the estrogen-independent expression levels of different ERα-target genes compared with BRCA1-proficient NEO1 cells may provide a mechanistic basis for the estrogen-independent growth advantages that AS4 cells exhibit.

The finding that BRCA1 can function as a ligand-reversible barrier to transcriptional activation by unliganded ERα suggests a mechanistic basis for the estrogen-independent growth feedback control enlisted by BRCA1-mutated breast epithelial cells to restrict the promiscuous expression of estrogen-responsive genes. Future studies should illuminate the mechanistic basis for BRCA1-mediated transcriptional repression of ERα and clarify its functional role in the larger network of hormone signaling pathways that control the growth, differentiation, and homeostasis of breast and ovary.

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Deficient Nonhomologous End-Joining Activity in Cell-free Extracts from Brca1-null Fibroblasts

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ABSTRACT

BRCA1 ensures genome stability, at least in part, through a functional role in DNA damage repair. BRCA1 interacts with the Rad50/Mre11/Nbs1 complex that occupies a central role in DNA double-strand break repair mediated by homologous recombination and nonhomologous end joining (NHEJ). NHEJ can be catalyzed by mammalian whole cell extract in a reaction dependent upon DNA ligase IV, Xrc4, Ku70, Ku80, and DNA-PKcs. Here, we show that under identical cell-free reaction conditions, the addition of antibodies specific for BRCA1 and Rad50 but not Rad51, inhibits end-joining activity. Cell extracts derived from Brca1-deficient mouse embryonic fibroblasts exhibit reduced end-joining activity independent of the endogenous protein amounts of DNA ligase IV, Ku80, and Ku70. The Brca1-dependent NHEJ activity predominates at the lower concentrations of Mg2+ (0.5 mM); elevated Mg2+ or Mn2+ concentrations (10 mM) dramatically increase overall end-joining activity and abrogates the requirement for Brca1, Xrc4, and Ku70. The addition of partially purified BRCA1, in association with Rad50/Mre11/Nbs1 complex, complements the NHEJ deficiency of Brca1-null fibroblast extracts. These results suggest a role for Brca1 in NHEJ and in the maintenance of genome integrity.

INTRODUCTION

Inactivation of the hereditary breast cancer susceptibility gene, BRCA1, leads to genomic instability (1–3). Extensive chromosomal abnormalities have been observed in Brca1-deficient murine fibroblasts (4), as well as the Brca1-mutant human breast cancer cell line HCC1937 (5). The function of BRCA1 in genome stability is attributable to its central role in the cellular response to DNA damage response, and emerging evidence supports a role for BRCA1 in DNA damage repair. For example, Brca1-deficient murine and human cells are sensitive to DNA-damaging agents, including IR (6–8). Furthermore, HCC1937 cells expressing mutant BRCA1 protein exhibit a reduction in both the rate and extent of DSB repair after IR when compared with cells expressing wild-type BRCA1 protein (9). Finally, BRCA1 physically interacts with the Rad50/Mre11/Nbs1 DSB repair complex and colocalizes to nuclear foci along with this complex after treatment of cells with IR (8).

In eukaryotic cells, DSBs are repaired through two distinct pathways: homologous recombination and NHEJ. BRCA1 has been implicated in homology-based repair because cells expressing a Brca1 exon-11 deletion mutant exhibit defects in gene targeting, single-strand annealing, and gene conversion (10). BRCA1 may also influence NHEJ by virtue of its interaction with the Rad50/Mre11/Nbs1 complex. The orthologous complex in Saccharomyces cerevisiae Rad50/Mre11/Xrs2 is critical for NHEJ, sister chromatid recombination, and telomere maintenance. Yeast strains deficient in any of the components of the Rad50/Mre11/Xrs2 complex are 10–100-fold less efficient in nonhomologous joining of DNA ends (11, 12). The Rad50/Mre11/Nbs1 complex is characterized by 3′ to 5′ exonuclease activity on double-stranded DNA and endonuclease activity on single-stranded DNA and hairpin structures. Furthermore, in the presence of a DNA ligase, Mre11 can facilitate DNA end joining using microhomologies at or near DNA termini (13, 14). Recently, the yeast Rad50/Mre11/Xrs2 complex was found to exhibit DNA end-binding activity and end-bridging activity (15). Thus, the Rad50/Mre11/Nbs1 complex may fulfill a functionally conserved role in NHEJ.

In mammalian cells, a NHEJ pathway has been identified that comprises the heterodimeric DNA end-binding activity Ku70/Ku80 and the DNA-PKcs (reviewed in Ref. 12). Recently, Baumann et al. (16) developed a cell-free system that faithfully reflects the genetic requirements for this NHEJ pathway. In this system, accurate intermolecular ligation of DNA ends was found to be dependent on DNA ligase IV/Xrc4 and requires Ku70, Ku80, and DNA-PKcs. However, the role of Rad50/Mre11/Nbs1 in this NHEJ assay has not been addressed.

We report here the use of this cell-free assay to investigate the role of BRCA1 in DNA end joining. We observed that antibodies specific for BRCA1, Rad50, and Ku70, but not Rad51, inhibit the end-joining activity present in extracts prepared from a human lymphoblastoid cell line. Comparison of extracts derived from Brca1-null MEFs with that from isogenic Brca1-proficient MEFs for their respective abilities to catalyze end joining in vitro revealed that Brca1-deficient MEF extracts exhibit a significantly reduced end-joining activity. This deficiency can be complemented by partially purified BRCA1 in association with the Rad50/Mre11/Nbs1 complex. Finally, we found the BRCA1-dependent NHEJ activity in mammalian WCE to be sensitive to the reaction concentration of divalent cations. Elevated concentrations of Mg2+ or Mn2+ (to 10 mM) dramatically increased overall end-joining efficiency and abrogated the requirement for BRCA1, Ku, and Xrc4 requirement. These results provide evidence that BRCA1 promotes NHEJ in a Mg2+ concentration-dependent manner.

MATERIALS AND METHODS

MEFs and Lymphoblastoid Cell Line. The Brca1+/−, p53+/−, and p53−/− MEFs were derived from 9.5-day old embryos of a cross between Brca1+/− and p53−/− mice as described (17) and cultured in DMEM plus 10% FCS. Human lymphoblastoid cell line, LEM, was immortalized by Epstein-Barr virus and cultured in DMEM plus 10% FCS.

Cell-free NHEJ Assay. Cell extracts were prepared and in vitro reactions were performed according to previously described procedure (16) and modified for their respective total protein levels using the Bio-Rad protein assay (Bio-Rad, Richmond, CA). Reactions (16 μl) were carried out in 50 mM triethanolamine-HCl (pH 7.5), 0.5 mM Mg(ΟAc)2, 80 mM potassium acetate, 2 mM ATP, 1 mM DTT, and 100 μg/ml BSA. Cell-free extracts were incubated for 5 min at 37°C before the addition of 5 fmol 32P-labeled DNA. pBSK+ (A) duplex plasmid DNA (2.96 kb; Stratagene, La Jolla, CA) was linearized with EcoRI, dephosphorylated using calf intestinal phosphatase, and was 5′-end-labeled using polynucleotide kinase. In each reaction, 5 fmol of labeled DNA was used. After incubation at 37°C for 1 h, 32P-labeled DNA products were deproteinized by proteinase K (500 μg/ml) and 1% SDS at 37°C for 20 min and analyzed by electrophoresis through 0.7% agarose gels, followed by autoradiography. Quantitation of DNA end-joining efficiency was carried out...
by densitometry. For antibody inhibition experiments, cell extracts were preincubated with specific antibodies on ice for 30 min before shifting to 37°C for 5 min, followed by the addition of 32P-labeled DNA.

**Antibodies and Antisera.** A recombinant protein containing glutathione S-transferase fused with mouse Brca1 of amino acids 788-1135 in frame was used as an antigen for the production of antimouse Brca1 mouse polyclonal antiserum. Purified goat IgG specific for XRCC4, Ku70, and Ku80 was purchased from Santa Cruz Biotechnology (Santa Cruz, CA). Purified mouse monoclonal antibody specific for BRCA1 (Ab-I) was purchased from Oncogene Research Products (San Diego, CA). Other antibodies specifically against BRCA1, Rad50, and Rad51 have been described (8).

The amounts of the following antibodies were used in NHEJ inhibition experiments: 10 μg of antimouse IgG; 1 μg of purified goat anti-Ku70 IgG; 1 μg of anti-BRCA1 mAb Ab-I; 2 μg of anti-BRCA1 mAb 17F8; 5 μg of anti-Rad50 mAb 13B3; 1 μg of purified rabbit anti-Rad51 IgG; and 1 μg of purified rabbit anti-Xrcc4. Only mAbs and commercial available purified antisera were used in antibody inhibition assays. Most of the polyclonal antisera contain high levels of nuclease activity and cannot be used in the cell-free end-joining assay.

**Protein Purification.** HeLa cell nuclear extract was subjected to successive phosphocellulose P-11, DEAE-Sepharose, and Superose 6 gel filtration chromatography as described previously (18). Individual fractions derived from Superose 6 chromatography corresponding to the peaks of the Rad50, Mre11, and NBS1 proteins were pooled and designated as fraction C1. The fraction C1 was concentrated on phosphocellulose P-11 by elution with 0.5 M KCl in D buffer (20 mm HEPES (pH 7.9), 0.2 mm EDTA, and 20% glycerol), followed by dialysis with 0.1 M KCl in D buffer.

**RESULTS**

To explore the function of BRCA1 in DNA repair, we used an in vitro DNA end-joining assay that has been described previously (16). In this system, NHEJ catalyzed by human WCE was observed by rejoining 32P-labeled linear duplex DNA in a reaction that is dependent upon all of the mammalian factors thus far genetically implicated in NHEJ, including DNA ligase IV, Xrcc4, Ku70, Ku80, and DNA-PKcs. We initially tested WCE from a human lymphoblastoid cell line, LEM, for its ability to catalyze NHEJ and observed that 25-35% of input DNA molecules were rejoined during a 1-h incubation directly compared WCEs derived from either LEM extract alone. Note that BRCA1-, Rad50-, and Ku70-specific, but not control, antibodies inhibited the end-joining activity present in WCE. To further substantiate the requirement for BRCA1 in NHEJ, we inhibited end-joining activity in vitro. (16). However, antibodies specific for Rad50 or normal murine IgG did not inhibit end-joining activity. These results indicate that BRCA1 and Rad50, but not Rad51, may be involved in NHEJ in this cell-free system.

To further substantiate the requirement for BRCA1 in NHEJ, we directly compared WCEs derived from Brca1-/- p53-/- MEFs with extracts from both p53-/- and wild-type MEFs for their respective NHEJ activities in vitro. WCEs from wild-type or p53-/- MEFs could rejoin 35-50% of input DNA. Significantly, WCE from Brca1-/- p53-/- MEFs was reduced 3-10-fold consistently, relative to WCE from either p53-/- or wild-type MEFs, for end-joining activity (Fig. 2, A and B). For quantitative standardization, WCEs inhibited end-joining activity in vitro. (16). However, antibodies specific for Rad50 or normal murine IgG did not inhibit end-joining activity. These results indicate that BRCA1 and Rad50, but not Rad51, may be involved in NHEJ in this cell-free system.

**Fig. 2.** Deficient end-joining activity in Brca1 mutant MEFs. A, end-joining activity of MEF extracts. Increasing amounts of WCE from each of the indicated MEF cell lines were incubated with 32P-end-labeled linear DNA and assayed for end-joining activity. B, quantitative analysis by phosphorimaging. End-joining efficiency was calculated as: intensity of end-joining products/total substrate × 100%; error bars indicate the experimental deviation. C, immunoblot analysis demonstrating equivalent levels of representative NHEJ proteins and an unrelated nuclear matrix protein, p84, in the MEF WCEs assayed for end-joining activity.

**Fig. 1.** Antibodies specific for BRCA1, Rad50, and Ku70 inhibit the end-joining activity present in human cell extract. Human lymphoblastoid LEM WCE was preincubated with BRCA1-specific antibodies Ab-I (Lanes 9–11) and 17F8 (Lanes 12–14), Ku70-specific antibody (Lanes 6–8), Rad50-specific antibody (Lanes 15–17), Rad51-specific antibody (Lanes 18–20), or control antibody murine IgG (Lanes 3–5) before the addition of 32P-end-labeled linear DNA (5 fmol/reactions) and subsequent incubation. Antibodies or antisera were added at 1:6 serial dilutions as follows (the actual amounts were described in the "Materials and Methods" section): straight, Lanes 5, 8, 11, 14, 17, and 20; diluted 1/6, Lanes 4, 7, 10, 13, 16, and 19; diluted 1/36, Lanes 3, 6, 9, 12, 15, and 18. Lane 1: no extract. Lane 2: LEM extract alone. Note that BRCA1, Rad50-, and Ku70-specific, but not control, antibodies inhibited end joining. The end-joining efficiency was calculated as the end-joining activity [intensity of monomers/(monomers + trimers + tetramers)] in each reaction expressed relative to the end-joining activity in WCEs without antibodies as determined by densitometric analysis.
were first normalized for total protein levels and subsequently analyzed by immunoblot analyses for their respective expression levels of Ku70, Ku80, DNA ligase IV, Rad50, Mre11, or the nuclear matrix protein β4 (19). No significant difference in protein levels could be observed among these WCEs (Fig. 2C), excluding the possibility that the reduced end-joining activity in Brca1-null cell extract is because of variations in the expression levels of these NHEJ proteins.

It is known that divalent cations such as Mg2+ and Mn2+ affect NHEJ catalyzed by mammalian WCE in vitro (20). Our NHEJ reactions included 0.5 mM Mg2+ and were performed identically to those described initially by Baumann et al. (16). To test whether the Brca1-dependent NHEJ activity present in WCEs of MEFs is affected by the concentration of divalent ions, we compared extracts derived from Brca1-deficient and wild-type MEFs for their respective NHEJ activities in the presence of increasing concentrations of Mg2+ or Mn2+. As shown in Fig. 3, augmentation of the Mg2+ or Mn2+ concentration dramatically increased the level of DNA and joining catalyzed by both Brca1-null and wild-type cell extracts and concomitantly abrogated the requirement for Brca1 regardless the amount of WCE used in the reactions. Thus, at reaction concentrations of divalent ions exceeding 1.5 mM Mg2+ or 0.5 mM Mn2+, the difference in NHEJ catalyzed by Brca1-null and wild-type MEF extracts was indistinguishable, possibly indicating the involvement of a Brca1-independent pathway for NHEJ under these conditions.

To explore the relationship between the Brca1-dependent NHEJ activity and known components of the NHEJ pathway, including Xrc4 and Ku70, at different divalent ion concentrations, we performed reactions with either 0.5 mM or 10 mM Mg2+ (Fig. 4). Addition of Brca1-specific antibody reduced end-joining activity in Brca1 proficient MEFs to the level of Brca1 mutant cells at 0.5 mM Mg2+. Similarly, the addition of antibodies against Ku70 and Xrc4 completely eliminated end-joining activity of Brca1 deficient or proficient cells. These observations suggest that BRCA1 may function along with Xrc4 and Ku70 in NHEJ. However, at 10 mM Mg2+, the additions of antibodies against Ku70, Xrc4, and BRCA1 have no apparent inhibitory function against the robust end-joining activities in both Brca1-proficient and -deficient MEFs (Fig. 4), indicating an existence of an alternative pathway.

To determine whether a cellular fraction containing BRCA1 can complement the diminished NHEJ activity in Brca1-deficient cells, we fractionated human HeLa cell nuclear extract according to the scheme outlined in Fig. 5A. The bulk of BRCA1 protein present in a soluble HeLa nuclear extract bound to phosphocellulose PC-11 and eluted predominately and approximately equally between 0.1-0.3 and 0.3-0.5 M KCl step fractions (fractions B and C, respectively, Fig. 5A). The bulk of the cellular Rad50 protein was also recovered in the PC-11 B and C fractions, although more eluted in the B fraction than in the C fraction (Fig. 5A).

Most of the BRCA1 protein present in the PC-11 C fraction bound to, and eluted from, a DEAE-Sepharose anion exchange resin in a 0.1-0.25 M KCl step fraction (fraction CB). After Superose 6 gel filtration chromatography of the CB fraction, BRCA1 was eluted in fractions corresponding to peaks of the Rad50, Mre11, and Nbs1 proteins, indicating cofractionation of BRCA1 with the Rad50/Mre11/Nbs1 protein complex (Fig. 5A). Western blot analysis of individual Superose 6 column fractions with antibodies specific for BRCA1 and the large subunit of RPB1 also revealed that the bulk of BRCA1 eluted ahead of RPB1 in a number of high molecular weight fractions (Fig. 5A). Reciprocal coimmunoprecipitation of BRCA1 and Rad50 from peak Superose 6 column fractions demonstrated that BRCA1, Rad50, and Nbs1 all reside in a single high molecular weight complex of Mr. ~1,000,000 (Fig. 5B).

Individual Superose 6 column fractions corresponding to the peak of the BRCA1/Rad50/Mre11/Nbs1 complex were pooled and concentrated on phosphocellulose PC-11 to yield a partially purified protein fraction termed C1. Fraction C1 was tested for its ability to complement WCE of Brca1-deficient MEFs for end-joining activity in vitro. Although it catalyzed no end-joining activity on its own, the addition of fraction C1 to WCE of Brca1-deficient MEFs increased its end-joining activity about 2.5-fold (Fig. 5, C and D) to 60% of the end-joining activity catalyzed by WCE derived from Brca1-proficient MEFs. The addition of fraction C1 to WCE of Brca1-proficient MEFs has no effect on end-joining activity (data not shown). These results suggest that the partially purified BRCA1 complex facilitated the BRCA1-dependent NHEJ process.

**DISCUSSION**

BRCA1 plays an important role in maintaining genomic stability through its participation in DNA repair and cell cycle checkpoint control. For DNA DSB repair, BRCA1 has been shown to be critical for homologous recombination (10). However, it is not known whether BRCA1 has a role in NHEJ. In this study, we showed that under identical cell-free reaction conditions described by Baumann et al. (16), the addition of antibodies specific for BRCA1 and Rad50, but not Rad51, inhibits end-joining activity. Cell extracts derived from Brca1-deficient MEFs exhibit reduced end-joining activity independent of the endogenous protein amounts of DNA ligase IV, Ku80, and Ku70. The BRCA1-dependent NHEJ activity predominates at the lower concentra-
The relative contribution of a particular pathway of multimers/(multimers + monomer) suggests the involvement of a DNA-PK-independent end-joining pathway eukaryotic cells rely on more than one DNA end-joining pathway. Recently, a purified recombinant BRCA1 was shown to have a direct contribution to the ability of BRCA1 to form a tight complex with Rad50/MreI lI/Nbsl-IV/Xrcc4. Although the precise role of BRCA1 in NHEJ remains unclear, the cell-free system for NHEJ that we used in this study has been shown to reconstitute NHEJ activity in the Rad50/MreI 1/Nbsl complex. HeLa cell nuclear extract (NEXT) was subjected to successive phosphocellulose (PC-11), DEAE-Sepharose, and Superose 6 gel filtration chromatography. CB protein fraction (fractions 45-49 (fraction C1) from the experiment in A were pooled, and a portion subjected to Superose 6 gel filtration analysis. Individual fractions were resolved by 10% SDS-PAGE and subjected to immunoblot analysis using antibodies specific for the proteins indicated above the immunoblot (mIgG = murine IgG). Immunoprecipitates were resolved by 10% SDS-PAGE and subjected to immunoprecipitation with the antibodies indicated above the immunoblot (RPBI: large subunit of RNA polymerase I1). Vertical arrowheads above the immunoblot panels indicate marker protein peaks. B, coimmunoprecipitation of BRCA1, Rad50, and Nbs1 from peak Superose 6 column fractions. Column fractions 45-49 (fraction C1) from the experiment in A were pooled, and a portion subjected to immunoprecipitation with the antibodies indicated above the immunoblot (mIgG = murine IgG). Immunoprecipitates were resolved by 10% SDS-PAGE and subjected to immunoblot analysis with antibodies specific for the proteins indicated to the left of the blot (RPBI: large subunit of RNA polymerase I1). Vertical arrowheads above the immunoblot panels indicate marker protein peaks. B, coimmunoprecipitation of BRCA1, Rad50, and Nbs1 from peak Superose 6 column fractions. Column fractions 45-49 (fraction C1) from the experiment in A were pooled, and a portion subjected to immunoprecipitation with the antibodies indicated above the immunoblot (mIgG = murine IgG). Immunoprecipitates were resolved by 10% SDS-PAGE and subjected to immunoblot analysis with antibodies specific for the proteins indicated to the left of the blot.
reflects the *in vitro* reaction conditions used. One factor that could alter the relative influence of a particular end-joining pathway in the reaction is the concentration of divalent cations, particularly Mg2+ and Mn2+. In this regard, we observed that the BRCA1-dependent NHEJ activity present in mammalian WCE is sensitive to the reaction concentration of Mg2+ and Mn2+. Elevated concentrations of these divalent cations stimulate overall end-joining activity and mask the requirement for BRCA1, suggesting the involvement of a BRCA1-independent pathway to achieve end joining. A similar phenomenon has recently been reported for DNA ligase IV using ligase IV mutant 180BR cell (20). Elevated reaction concentrations of Mg2+ (10 mM) stimulated DNA end joining through an apparent DNA ligase IV-independent pathway, whereas reduced concentrations of Mg2+ (0.5 mM) revealed a DNA ligase IV dependency for low levels of end-joining activity. Interestingly, we observed that under similar conditions, antibodies specific for Ku70, Xrc4, and BRCA1 efficiently suppressed DNA end-joining activity at reduced concentrations of Mg2+ (Fig. 4). These observations raise the possibility that BRCA1 works along with Rad50, Ku, and Xrcc4 in NHEJ at a low concentration of Mg2+. Interestingly, mammalian cell extracts deficient in Fanconi anemia proteins had a 3-9-fold reduction in DNA end-joining activity at high Mg2+ concentration (10 mM) in a pathway independent of DNA-PK/ Ku (28). Therefore, it is very likely that multiple NHEJ pathways may exist in mammalian cells.

Previously, BRCA1 has been implicated in homologous-based repairs of DNA breaks because cells expressing a Brca1 exon 11-deletion mutant exhibit defects in gene targeting, single-strand annealing, and gene conversion (10). Interestingly, no defects in plasmid integration and nonhomologous repair processes were observed in these Brca1 mutant cells. However, it is possible that the NHEJ activity observed in this previous study described by Moyer et al. (10), reflects the contribution of a BRCA1-independent NHEJ pathway. Alternatively, genetic differences between independently derived Brca1 mutant cell lines may contribute to different conclusions regarding the importance of Brca1 in NHEJ. Our Brca1 mutant allele carries a reverse-orientated neomycin cassette inserted into the 5′ end of Brca1 exon 11, which will lead to premature termination of translation (29). No stable Brca1 protein derivative can be detected in our Brca1+/− MEFs. The embryonic stem (ES) cells previously characterized for defects in homologous and NHEJ repair (10), express a Brca1 exon 11 splice variant (30) and the homozygote embryos derived from these ES cells exhibit a less severe phenotype than our homozygotes (29, 30). Using our Brca1+/− MEFs, we have demonstrated a 50–100-fold reduced efficiency in microhomology-mediated end-joining activity at high Mg2+ concentration (10 mM) in a pathway deficient in Fanconi anemia proteins had a 3-9-fold reduction in DNA end-joining activity at low concentration of Mg 2+. Interestingly, mammalian cell extracts from Brca1-null cell lines exhibit a less severe phenotype than our homozygotes characterized by neuroepithelial ablation of postimplantation development. Genes Dev., 10: 1835-1843, 1996.

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REFERENCES


Functional Dissection of Transcription Factor ZBRK1 Reveals Zinc Fingers with Dual Roles in DNA-binding and BRCA1-dependent Transcriptional Repression*

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The breast- and ovarian-specific tumor suppressor BRCA1 has been implicated in both activation and repression of gene transcription by virtue of its direct interaction with sequence-specific DNA-binding transcription factors. However, the mechanistic basis by which BRCA1 mediates the transcriptional activity of these regulatory proteins remains largely unknown. To clarify this issue, we have examined the functional interaction between BRCA1 and ZBRK1, a BRCA1-dependent KRAB eight zinc finger transcriptional repressor. We report here the identification and molecular characterization of a portable BRCA1-dependent transcriptional repression domain within ZBRK1 composed of zinc fingers 5-8 along with sequences in the unique ZBRK1 C-terminus. This C-terminal repression domain functions in a BRCA1-, histone deacetylase-, and promoter-specific manner and is thus functionally distinguishable from the N-terminal KRAB repression domain in ZBRK1, which exhibits no BRCA1 dependence and broad promoter specificity. Significantly, we also find that the BRCA1-dependent transcriptional repression domain on ZBRK1 includes elements that modulate its sequence-specific DNA binding activity. These findings thus reveal the presence within ZBRK1 of functionally bipartite zinc fingers with dual roles in sequence-specific DNA-binding and BRCA1-dependent transcriptional repression. We discuss the implications of these findings for the role of BRCA1 as ZBRK1 co-repressor.

Germ line inactivation of the gene encoding BRCA1 confers a cumulative lifetime risk of female breast and ovarian cancer (1–3). Although the mechanistic basis for its tissue- and gender-specific tumor suppressor activity remains poorly defined, BRCA1 nonetheless fulfills a broad function in the maintenance of global genome stability (4–10). The underlying basis for this caretaker activity likely derives from the role of BRCA1 as a conduit in the cellular DNA damage response, wherein it serves to couple DNA damage-induced signals to downstream responses including DNA damage repair and cell cycle checkpoint activation (6, 7, 11–21). Several potentially overlapping cellular activities have been ascribed to BRCA1, each of which could underlie its ability to control signal output. For example, BRCA1 has been implicated in chromatin remodeling, ubiquitination, recombination, and transcriptional regulation (6, 22–28). The extent to which these pleiotropic activities contribute to the caretaker function of BRCA1 is presently unknown; however, the fact that each of these BRCA1-associated activities is similarly abrogated by cancer-predisposing BRCA1 missense mutations suggests a strong correlate link between their discharge and BRCA1-mediated tumor suppression.

With respect to its role in transcription control, BRCA1 has been implicated in both activation and repression of genes linked to a variety of biological processes, including cell growth control and DNA replication and repair (21, 29–31). Thus, by virtue of its transcriptional regulatory activity, BRCA1 could influence cellular responses downstream of DNA damage signals, and this activity could contribute to its caretaker function.

The precise role of BRCA1 in gene-specific transcription control has yet to be definitively established. Because it exhibits no sequence-specific DNA binding activity, it seems likely that BRCA1 is targeted to specific genes through its functional interaction with sequence-specific DNA-binding transcription factors. Direct evidence to support this hypothesis has come from the identification of multiple DNA-binding transcription factors with which BRCA1 has been shown to physically interact and functionally synergize, including p53, c-Myc, estrogen receptor α, androgen receptor, OCT-1, NF-YA, and ZBRK1 (32–40). However, the underlying mechanism by which BRCA1 mediates the transcriptional stimulatory or repressive effects of these regulatory proteins has not been established.

We have been studying the functional interaction between BRCA1 and the transcriptional repressor ZBRK1 as a model system to understand the mechanistic basis by which BRCA1 mediates sequence-specific transcription control. Initially identified by virtue of its physical interaction with BRCA1, ZBRK1 (Zinc finger and BRCA1-interacting protein with a KRAB domain) is a member of the Kruppel-associated box-zinc finger protein (KRAB-ZFP) family of transcriptional repressors (39, 41).

Typically, KRAB-ZFPs bind to their corresponding target genes through tandem C-terminal C2H2 zinc fingers and repress transcription through an N-terminal KRAB domain, which silences gene expression through the indirect recruitment of histone deacetylases, histone methyltransferases, and heterochromatin proteins (41–47). Like other KRAB-ZFPs, ZBRK1 harbors an N-terminal KRAB domain. However, ZBRK1 is atypical among KRAB-ZFPs due to the fact that it harbors 8 central C2H2 zinc fingers and a unique C terminus.
that is absent among the larger family of KRAB-ZFPs.

Through its 8 central zinc fingers, ZBRK1 binds to a compositionally flexible 15-bp DNA sequence, GGGxxxCGGxxxTTT (where x is any nucleotide) (39). A search for potential ZBRK1 DNA-binding sites in existing genes led to intron 3 of GADD45a, a functionally important DNA damage-response effector known to be regulated transcriptionally by BRCA1 (21, 32, 39). Functional analysis revealed that ZBRK1 represses GADD45a gene transcription through its intron 3 DNA-binding site in a BRCA1-dependent manner, thus revealing BRCA1 to be a ZBRK1 co-repressor (39). Significantly, familial breast cancer-derived mutants of BRCA1 that disrupt its interaction with ZBRK1 abrogate its co-repressor activity, suggesting that its co-repressor function may be important for the tumor suppressor properties of BRCA1 (39). The regulation of GADD45a gene transcription is likely to be complex and controlled coordinately by ZBRK1 and BRCA1 in concert with other transcription factors, including p53, OCT1, and NF-YA, that function through cis-acting sequence elements in GADD45a promoter regions (21, 32, 39, 48, 49).

In addition to GADD45a, potential ZBRK1-binding sites have been identified in other DNA damage-response genes that are also regulated by BRCA1, including p21, Bax, and GADD153 (39). This observation suggests a potentially broader role for BRCA1 and ZBRK1 in the coordinate transcriptional regulation of diverse DNA damage-response genes. To begin to explore the mechanism by which BRCA1 mediates sequence-specific transcriptional repression through ZBRK1, we have pursued in greater depth the physical and functional interaction between these two proteins. We report here the identification and molecular characterization of a portable BRCA1-dependent transcriptional repression domain within ZBRK1 composed of zinc fingers 5–8 along with sequences in the unique ZBRK1 C terminus. This C-terminal repression domain functions in a BRCA1-, histone deacetylase (HDAC), and promoter-specific manner and is thus functionally distinguishable from the N-terminal KRAB repression domain in ZBRK1, which exhibits no BRCA1 dependence and broad promotor specificity. Significantly, we also find that the BRCA1-dependent C-terminal transcriptional repression domain within ZBRK1 is composed of elements that mediate sequence-specific DNA-binding by zinc fingers 1–4. These findings thus reveal an unanticipated dual function for the ZBRK1 zinc fingers in DNA binding and transcriptional repression, and further shed new light on the mechanistic role of BRCA1 in sequence-specific transcriptional control.

**EXPERIMENTAL PROCEDURES**

**Plasmid Construction and Mutagenesis**

**Expression Plasmids**—pMAL-C2-TEV-ZBRK1 ΔK for expressing MBP-ZBRK1 AK in *Escherichia coli* was derived from pGFPK3-ZBRK1 ΔK, which is a derivative of pCNP-ZBRK1 ΔK (39). Briefly, a BamHI-XhoI fragment carrying ZBRK1 DNA sequences encoding ZBRK1 amino acids 144–532 (lacking the N-terminal KRAB domain) was subcloned into the BamHI and XhoI sites of pMAL-C2-TEV (provided by Dr. P. Renee Yew), thereby generating a translational fusion of MBP-ZBRK1 AK with ZBRK1 amino acids 144–431 and replaced with corresponding PCR-amplified MBP-ZBRK1 AK deletion fragments encoding amino acids 144–343; ZBRK1 AK deleted from amino acid 144 to 343 had a ZRE (a CRT codon) substitution mutation at the first of the two conserved histidine residues within the targeted C2H2 zinc finger.

ZBRK1 5ZFC was expressed as a GAL4 DNA-binding domain fusion in mammalian cells from the plasmid GALA-ZBRK1 5ZFC, constructed by subcloning a PCR-amplified ZBRK1 cDNA fragment encoding amino acids 319–532 (comprising zinc finger 5 through the C terminus) into the SalI and HindIII sites of pM (Clontech, Palo Alto, CA). N- and C-terminal truncation derivatives of GALA-ZBRK1 5ZFC bearing stepwise deletions of individual zinc fingers were generated, each of which was reconstituted by PCR-based subcloning. Briefly, the 532 amino acid residues of ZBRK1 5ZFC was replaced with corresponding PCR-generated ZBRK1 deletion fragments. Individual GALA-ZBRK1 5ZFC deletion derivatives encode the following ZBRK1 amino acids (aa): MBP-ZBRK1 AK 6ZF1 (aa 144–431), 7ZF2 (aa 144–400; 6ZF3 (aa 144–377); 5ZF4 (aa 144–347); 4ZF5 (aa 144–319); 3ZF6 (aa 144–291); 2ZF7 (aa 144–265); and 1ZF8 (aa 144–235). Individual MBP-ZBRK1 AK broken finger (BF) mutants were generated by PCR-based site-directed mutagenesis using the QuickChange II site-directed mutagenesis kit following the manufacturer's recommendations (Stratagene, La Jolla, CA). Each broken finger mutant bears a histidine (i.e., CRT codon) to asparagine (i.e., aAT codon) substitution mutation at the first of the two conserved histidine residues within the targeted C2H2 zinc finger.

ZBRK1 5ZFC and its truncation and broken finger derivatives were expressed in yeast as GAL4 activation domain fusions using pGADT7 (Clontech, Palo Alto, CA). Briefly, ZBRK1 5ZFC and its truncation and broken finger derivatives (RF5, RF6, RF7, and RF8) were constructed by PCR-based mutagenesis. Broken finger derivatives of GALA-ZBRK1 5ZFC were generated by PCR amplification of ZBRK1 sequences encoding amino acids 319–532 from individual pMAL-C2-TEV ZBRK1 AK BF mutants and fragment replacement of the SacI-HindIII wild-type ZBRK1 fragment in GALA-ZBRK1 5ZFC.

ZBRK1 5ZFC and its truncation and broken finger derivatives were expressed in yeast as GAL4 activation domain fusions using pGADT7 (Clontech, Palo Alto, CA). Briefly, ZBRK1 5ZFC and its truncation and broken finger derivatives (RF5, RF6, RF7, and RF8) were constructed by PCR-based mutagenesis. Broken finger derivatives of GALA-ZBRK1 5ZFC were generated by PCR amplification of ZBRK1 sequences encoding amino acids 319–532 from individual pMAL-C2-TEV ZBRK1 AK BF mutants and fragment replacement of the SacI-HindIII wild-type ZBRK1 fragment in GALA-ZBRK1 5ZFC.

GADD153 (39). This observation suggests a potentially broader functions in a BRCA1-, histone deacetylase (HDAC), and responding domain within pGBKT7, thereby generating a translational fusion of the GAL4 DNAbinding domain with BRCA1 amino acids 341–748 in the plasmid pGBK7-BRCA1.

The ZBRK1 KRAB domain was expressed as a GAL4 DNA-binding domain fusion in mammalian cells from GALA-ZBRK1 KRAB, constructed by subcloning a PCR-amplified ZBRK1 cDNA fragment encoding amino acids 1–85 into the BamHI and blunted XhoI sites of pM2 (50). All PCR-based subcloning was performed using *Pfu* DNA polymerase (Stratagene, La Jolla, CA), and the integrity of individual deletion and substitution mutations was confirmed by DNA sequence analyses.

**Reporter Plasmids**—pG5-TK-Luc carrying five copies of the GAL4 DNA-binding site upstream of the herpes simplex virus thymidine kinase (TK) promoter (sequences corresponding to −105 to +51, where +1 is the transcription start site) driving expression of the gene encoding firefly luciferase was constructed by replacing a HindIII-BglII fragment from pSBS-GAL-TK-Luc (provided by Dr. P. Renee Yew) with a HindIII-BglII fragment from pG5-TK-CAT (provided by Dr. P. Renee Yew), thus positioning five copies of the GAL4 DNA-binding site upstream of the TK promoter. The reporter plasmid psG5-SV40-Luc carrying five GAL4 DNA-binding sites upstream of the SV40 promoter driving expression of the firefly luciferase gene has been described previously (39). pG5-SNRFN-Luc (provided by Dr. Paul A. Wade) carries four GAL4 DNA-binding sites upstream of the human small nuclear ribonucleoprotein N promoter driving expression of the firefly luciferase gene (51).

**Recombinant Protein Expression and Purification**

MBP-ZBRK1 fusion proteins were expressed in and purified from *E. coli* strain BL21 Star (DE3) pLysS (Invitrogen). Briefly, cells were grown at 37 °C to an A600 of 0.6. Isopropyl-1-thio-D-galactopyranoside was added to a final concentration 0.3 mM, and the cells were transferred to 25 °C for another 5.5 h. Cells were pelleted, washed once with phosphate-buffered saline, and then resuspended in MBP binding buffer (55 mM Tris-HCl, pH 7.5, 1 mM EDTA, 200 mM NaCl, 20 μM

1 The abbreviations used are: HDAC, histone deacetylase; EMSA, electrophoretic mobility shift assay; MEFs, mouse embryo fibroblasts; aa, amino acids; MBP, maltose-binding protein; PMSF, phenylmethylsulfonyl fluoride; TSA, trichostatin A; ZRE, ZBRK1-response element; BF, broken finger; TK, thymidine kinase; WT, wild type; Mut., mutant; HSV, herpes simplex virus.
Mutant (Mut) probe was obtained GCCG-3’. ZBRK1 DNA-binding site: 5’-GATCCACCGGAGGAGGTGGGGGCTC-3’ (39). An internal control plasmid, pCHh1O (40), expressing β-galactosidase under control of the SV40 promoter. Forty-eight hours after transfection, cells were transfected at 60% confluence using Effectene reagent (Qiagen, Valencia, CA). 

**Figure 1. Identification of DNA binding determinants on ZBRK1.**

A schematic representation of ZBRK1 (the KRAB domain and numbered zinc fingers are indicated). MBP-ZBRK1 AK, and MBP-ZBRK1 AK truncation derivatives. B, purified MBP-ZBRK1 AK and its corresponding truncation derivatives were resolved by SDS-10% PAGE and visualized by Coomassie Blue staining. Molecular weight marker positions (Mr) are indicated. C, competition EMSA. EMSA was performed using a 32P-labeled double-stranded oligonucleotide probe corresponding to a wild-type consensus ZRE and purified MBP-ZBRK1 AK and its corresponding truncation derivatives. EMSA was performed using the consensus ZRE probe and 50 ng of either MBP-ZBRK1 AK or each of its corresponding deletion derivatives as indicated. NP indicates no protein added to the EMSA reaction. The positions of the unbound ZRE oligonucleotide probe (ZRE) and the ZRE-MBP-ZBRK1 AK nucleoprotein complex (ZRE-ZBRK1) are indicated. For each EMSA reaction, the proportional fraction of ZRE probe bound by MBP-ZBRK1 AK is indicated below each lane and was determined by dividing the number of radioactive counts in the bound probe by the number of radioactive counts in the bound plus the unbound probe. D and E, sequence-specific DNA binding activity of MBP-ZBRK1 AK and its corresponding truncation derivatives. EMSA was performed using the consensus ZRE probe and 50 ng of either MBP-ZBRK1 AK or each of its corresponding deletion derivatives as indicated. NP indicates no protein added to the EMSA reaction. The positions of the unbound ZRE oligonucleotide probe (ZRE) and the ZRE-MBP-ZBRK1 AK nucleoprotein complex (ZRE-ZBRK1) are indicated. DNA-binding reactions were performed in 100 mM NaCl (D) or 200 mM NaCl (E). For each EMSA reaction, the proportional fraction of ZRE probe bound by MBP-ZBRK1 AK or its truncation derivatives is indicated below each lane and was determined by dividing the number of radioactive counts in the bound probe by the number of radioactive counts in the bound plus the unbound probe. ND, not detectable.

For EMSA, the ZRE probe was obtained by annealing two complementary oligonucleotides corresponding in sequence to the consensus ZBRK1 DNA-binding site: 5’-GATCCACCGGAGGAGGTGGGGGCTC-3’ and 5’-GATCCACCGGAGGAGGTGGGGGCTC-3’ (39). Mutant (Mut) probe was obtained by annealing two oligonucleotides, 5’-GATCCACCTCAGGGTCACGGTTCGGCGG-3’ and 5’-GATCCACCTCAGGGTCACGGTTCGGCGG-3’ (39). Each of these double-stranded probes carried overhangs, which were filled in with T4 DNA polymerase. In each reaction, purified MBP-ZBRK1 fusion proteins (50 ng) were incubated with 6000 cpm of an [32P]-labeled double-stranded oligonucleotide probe in 30 µl of EMSA binding buffer (25 mM Tris-HCl pH 7.5; 20 µM ZnCl2; 12.5% glycerol; 0.5 mM PMSF; and a variable concentration of NaCl as indicated). Following 30 min of incubation at room temperature, reaction mixtures were loaded onto a 5% nondenaturing polyacrylamide gel and electrophoresed at 200 V for 2 h at 4 °C in 0.5X TBE. Dried gels were subjected to PhosphorImager analysis (Amersham Biosciences).

**Cell Culture, Transfections, and Reporter Assays**

Brcal+/−, p53−/− (Brcal−/−), and p53−/− (Brcal+/+) mouse embryo fibroblasts (MEFs) (39) and U2OS human osteosarcoma cells were cultured in Dulbecco's modified Eagle's medium (Invitrogen) supplemented with 10% fetal bovine serum (HyClone, Logan, UT). Brcal+/+, Brcal−/−, and U2OS cells were transfected at 60% confluency using Effectene reagent (Qiagen, Valencia, CA), and the expression and reporter plasmids are indicated in each figure. Each transfection also included an internal control plasmid, pCHh1O (40), expressing β-galactosidase under control of the SV40 promoter. Forty-eight hours

**ZnCl2; and 10 mM β-mercaptoethanol** supplemented with protease inhibitors (aprotinin 0.4 µg/ml; chymostatin 0.5 µg/ml; leupeptin 0.5 µg/ml; pepstatin 0.5 µg/ml; PMSF 0.5 mM; and benzamidine-HCl 0.5 mM). Resuspended cells were frozen and thawed one time, followed by sonication (3 times for 1 min) and clarification by centrifugation at 30,000 × g for 30 min. MBP-ZBRK1 fusion proteins were purified from clarified lysates by affinity chromatography on amylose resin (New England Biolabs, Beverly, MA). Briefly, clarified lysates were incubated with amylose resin in buffer containing 0.5% maltose for 30 min at 4 °C. Purified proteins (estimated to be >95% homogeneous by SDS-PAGE and subsequent visualization by Coomassie Blue staining) were dialyzed for 1 h at 4 °C against EMSA storage buffer (25 mM Tris-HCl, pH 7.5; 100 mM NaCl; 20 µM ZnCl2; 10% glycerol; 10 mM β-mercaptoethanol; and 0.5 mM PMSF) before long term storage at −80 °C.

**Electrophoretic Mobility Shift Assay (EMSA)**

For EMSA, the ZRE probe was obtained by annealing two complementary oligonucleotides corresponding in sequence to the consensus ZBRK1 DNA-binding site: 5’-GATCCACCGGAGGAGGTGGGGGCTC-3’ and 5’-GATCCACCGGAGGAGGTGGGGGCTC-3’ (39).
post-transfection, cells were harvested and lysed in Reporter Lysis buffer (Promega, Madison, WI). Transfected cell lysates (20 µl) were analyzed for luciferase activity using the luciferase assay system (Promega, Madison, WI) and for β-galactosidase activity using the Galacto-light Plus Chemiluminescent Reporter Assay (BD Biosciences). Each transfection was repeated a minimum of 3 times in duplicate.

Yeast Two-hybrid Interaction Assay

pGADT7-ZBRK1 5ZFC and its truncation (6ZFC, 7ZFC, 8ZFC, 0ZFC, 5ZFC Δ1, 5ZFC Δ2, and 5ZFC Δ3) and broken finger (BF5, BF6, BF7, and BF8) derivatives were individually transformed along with pGADT7-BRCA1 (expressing BRCA1 amino acids 341–748) into the yeast strain Y187 (BD Biosciences/Clontech). After selection, colonies were expanded in liquid culture for assay of β-galactosidase activity following previously established procedures (39).

Transient Expression Analysis

Steady-state levels of GAL4-ZBRK1 5ZFC protein and its truncation and substitution derivatives were comparatively analyzed by immunoblot analysis of transfected whole cell extracts in order to verify equivalent levels of ectopic protein expression. Briefly, U2OS cells transfected with GAL4-ZBRK1 5ZFC, truncation mutants GAL4-ZBRK1 6ZFC, 7ZFC, 8ZFC, 0ZFC, 5ZFC Δ1, 5ZFC Δ2, and 5ZFC Δ3, and broken finger mutants GAL4-ZBRK1 5ZFC BF5, BF6, BF7, and BF8 were lysed in Laemmli sample buffer, resolved by SDS-10% PAGE, and subjected to immunoblot analysis using an antibody directed against the GAL4 DNA-binding domain (sc-510, Santa Cruz Biotechnology, Santa Cruz, CA) and, as an internal control protein, the p89 subunit of TFIH (sc-289, Santa Cruz Biotechnology, Santa Cruz, CA). Immunodetection was performed using ECL Western blotting detection reagents (Amersham Biosciences).

RESULTS

Identification of DNA-binding Determinants on ZBRK1—Previously, we demonstrated a strict requirement for BRCA1 in ZBRK1 repression. Specifically, we showed that ZBRK1 repression function was similarly abrogated by genetic ablation of BRCA1 or by deletion of the BRCA1-binding domain on ZBRK1 (39). The BRCA1-binding domain on ZBRK1 includes the last four of eight ZBRK1 zinc fingers (zinc fingers 5–8) along with the ZBRK1 C terminus (39). Whether and how this zinc finger domain contributes to the sequence-specific DNA binding activity of ZBRK1, however, is presently unknown. Because the specification of individual zinc fingers required to bind to DNA and/or BRCA1 could illuminate the underlying mechanism(s) by which BRCA1 mediates ZBRK1 repression, we initially sought to establish both the number and identity of the ZBRK1 zinc fingers required to bind DNA and BRCA1.

With respect to DNA binding, we showed previously (39) that the eight central ZBRK1 zinc fingers collectively recognize a 15-bp consensus sequence, GGGxxxCAGxxxTTT. Based on the observation that one C6H2 zinc finger can bind to ~3 bp of DNA (52–54), only five of the eight ZBRK1 zinc fingers would be predicted to bind to its 15-bp consensus sequence. To test this prediction, we analyzed a panel of ZBRK1 zinc finger deletion derivatives for their respective abilities to bind to the consensus ZBRK1 DNA-binding site in an EMSA.

To this end, we expressed ZBRK1 as a maltose-binding protein (MBP) chimera in E. coli, which permitted the purification of otherwise insoluble ZBRK1 protein. Full-length MBP-ZBRK1 is expressed poorly, whereas MBP-ZBRK1 ΔK (a deletion derivative lacking the N-terminal 143 amino acids of the 532-amino acid full-length ZBRK1 protein) is abundantly expressed. ZBRK1 ΔK lacks the N-terminal KRAB domain but retains the ZBRK1 zinc fingers and the C terminus, elements each required for binding to both DNA and/or BRCA1 (Fig. 1, A and B). We have therefore utilized this recombinant ZBRK1 ΔK derivative as the background into which truncation and substitution mutations have been introduced for purposes of DNA binding assays.

In an EMSA, MBP-ZBRK1 ΔK produced a discrete nucleoprotein complex on a double-stranded oligonucleotide probe corresponding to the consensus ZBRK1-response element (ZRE); a molar excess of unlabeled WT ZRE probe (WT probe), but not a mutant probe (Mut probe), efficiently competed for the formation of this complex, thus establishing sequence-specific DNA binding by MBP-ZBRK1 ΔK in this assay (Fig. 1C). To determine the number and identity of the ZBRK1 zinc fingers required to bind to its consensus sequence, we analyzed a series of C-terminal truncation derivatives bearing stepwise deletions of individual ZBRK1 zinc fingers (Fig. 1, A and B).
pGADT7-ZBRK1 derivatives: Relative β-Galactosidase Activity

FIG. 3. Identification of BRCA1-binding determinants on ZBRK1. The ZBRK1-binding region on BRCA1 (amino acids 341–748) (39), expressed as a GAL4 DNA-binding domain fusion protein (in the plasmid pGBKT7), was tested for interaction with the indicated fragments of ZBRK1 fused to the GAL4 transactivation domain (plasmid pGADT7) in yeast two-hybrid assays. β-Galactosidase activities were quantified as described previously (39). Corresponding β-galactosidase activities obtained with each pGADT7-ZBRK1 5ZFC derivative are expressed relative to that observed with the backbone pGADT7 expression vector alone, which was arbitrarily assigned a value of 1. Values represent the average of three independent assays, each performed in triplicate, and error bars represent the mean ± S.D. As a comparative measure of interaction strength, the average β-galactosidase activity obtained with pGBKT7-p53 and pGADT7-Large T antigen was 512 ± 13.5. The validity of two-hybrid interactions observed in these experiments was further substantiated by the following controls. First, we confirmed that the observed β-galactosidase activities were dependent upon BRCA1 sequences expressed from the plasmid pGBKT7-BRCA1 by performing a parallel series of interaction assays with individual pGADT7-ZBRK1 5ZFC derivatives and the backbone vector pGBKT7 as a negative control. Second, immunoblot analyses of yeast whole cell extracts confirmed that each of the pGADT7-ZBRK1 5ZFC fusion proteins was expressed at roughly equivalent levels, thus excluding the possibility that differences in β-galactosidase activities derive from differences in fusion protein expression.

100 mM NaCl, truncation of the ZBRK1 C terminus (MBP-ZBRK1 ΔK 6ZF) did not appreciably affect DNA binding relative to intact MBP-ZBRK1 ΔK (Fig. ID). Interestingly, deletion of the eighth and last ZBRK1 zinc finger along with the C terminus (MBP-ZBRK1 ΔK 7ZF) led to an increase in DNA binding activity, suggesting that ZBRK1 ZF8, and possibly the C terminus, constrains sequence-specific DNA binding mediated by the first seven zinc fingers (Fig. 1D). This effect was exaggerated under more stringent DNA binding conditions (200 mM NaCl) (Fig. 1E). At 100 mM NaCl, stepwise truncation of zinc fingers 7 to 5 led to a slight incremental reduction in sequence-specific DNA binding activity (Fig. 1D). ZBRK1 derivatives bearing less than four zinc fingers failed to bind to DNA, thereby establishing zinc fingers 1–4 as the minimal ZBRK1 DNA-binding domain under these conditions (Fig. 1D). Identical results were observed at 50 mM NaCl (data not shown). At 200 mM NaCl, zinc fingers 1–5 were required for stable DNA binding, and the inclusion of zinc fingers 6 and 7 incrementally stabilized binding (Fig. 1E).

To examine more rigorously the role of zinc fingers 5–8 in sequence-specific DNA binding by ZBRK1, we examined a set of “broken finger” mutants bearing substitution mutations within each of these zinc fingers. This approach permitted us to assess the individual contribution of each finger within the BRCA1-binding domain to overall DNA binding activity in the context of the eight-fingered ZBRK1 AK protein and thereby circumvent potential artifacts arising from analyses of truncation mutants. Each broken finger mutant bears a His-to-Asn substitution mutation at the first of the two conserved His residues within the targeted C₉H₉ zinc finger (Fig. 2, A and B). The relative conservative nature of this substitution eliminates zinc coordination within the targeted finger, thereby disrupting
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A and B, human U2OS cells were transfected with 30 ng of pG\(_{5}\)TK-Luc bearing five copies of the GAL4 DNA-binding site sequence upstream of the TK promoter without or with the indicated nanogram amounts of GAL4-ZBRK1 5ZFC (A) or GAL4-KRAB (B). In this and all subsequent transfection experiments involving effector plasmid titrations, the total amount of DNA in each transfection was fixed by reciprocal titration of the corresponding backbone expression plasmid. Also, in this and all subsequent transfection experiments, the relative luciferase activity represents the ratio of the luciferase activity obtained in a particular transfection to that obtained in cells transfected with only the reporter and pM (GAL4 DNA-binding domain) expression vectors alone. Luciferase activities were first normalized to \(\beta\)-galactosidase activity obtained by co-transfection of the SV40-\(\beta\)-gal vector (15 ng) as described previously (39). Error bars represent the S.D. from the average of at least three independent transfections performed in duplicate. C, schematic representation of GAL4-ZBRK1 5ZFC and GAL4-ZBRK1 KRAB chimeras (amino acid sequences fused to the GAL4 DNA-binding domain are indicated numerically above each chimera) and the pG\(_{5}\)TK-Luc reporter template used in transfection assays.

![Diagram](https://via.placeholder.com/150)

**FIG. 4.** ZBRK1 harbors two independent transcriptional repression domains. A and B, human U2OS cells were transfected with 30 ng of pG\(_{5}\)TK-Luc bearing five copies of the GAL4 DNA-binding site sequence upstream of the TK promoter without or with the indicated nanogram amounts of GAL4-ZBRK1 5ZFC (A) or GAL4-KRAB (B). In this and all subsequent transfection experiments involving effector plasmid titrations, the total amount of DNA in each transfection was fixed by reciprocal titration of the corresponding backbone expression plasmid. Also, in this and all subsequent transfection experiments, the relative luciferase activity represents the ratio of the luciferase activity obtained in a particular transfection to that obtained in cells transfected with only the reporter and pM (GAL4 DNA-binding domain) expression vectors alone. Luciferase activities were first normalized to \(\beta\)-galactosidase activity obtained by co-transfection of the SV40-\(\beta\)-gal vector (15 ng) as described previously (39). Error bars represent the S.D. from the average of at least three independent transfections performed in duplicate. C, schematic representation of GAL4-ZBRK1 5ZFC and GAL4-ZBRK1 KRAB chimeras (amino acid sequences fused to the GAL4 DNA-binding domain are indicated numerically above each chimera) and the pG\(_{5}\)TK-Luc reporter template used in transfection assays.

its local structure with little effect on the integrity of the remainder of the protein (55, 56). Consistent with the results obtained using C-terminal truncation mutants, analysis of individual BF mutants 5–8 revealed zinc finger 5 to be an important ZBRK1 determinant for stable DNA binding, whereas zinc fingers 6 and 7 promote but are not essential for binding (Fig. 2C). Disruption of zinc finger 8 did not appreciably affect the DNA binding activity of MBP-ZBRK1 ΔK, suggesting that zinc finger 8 is largely dispensable for stable association with DNA (Fig. 2C). Furthermore, local disruption of zinc finger 8 did not relieve constraints on the DNA binding activity of MBP-ZBRK1 ΔK, suggesting that the C terminus of ZBRK1 can also mask the inherent DNA binding activity of ZBRK1 zinc fingers 1–7 (Fig. 2C). In summary, the results of DNA-binding analyses delimit the core ZBRK1 DNA-binding domain to zinc fingers 1–4; these zinc fingers are minimally required for stable DNA binding under relatively non-stringent conditions of ionic strength. Zinc finger 5 is a critical and context-dependent determinant of stable binding and represents the extent of the minimal DNA-binding domain under more stringent binding conditions. Zinc fingers 6 and 7, although nonessential, nonetheless further stabilize DNA binding mediated by zinc fingers 1–5. Finally, zinc finger 8 and the C terminus apparently destabilize the maximum potential DNA binding activity inherent in zinc fingers 1–7.

Identification of BRCA1-binding Determinants on ZBRK1—Next, we sought to establish more precisely the molecular determinants on ZBRK1 required for BRCA1 binding. Previously, we mapped the BRCA1-binding domain on ZBRK1 to encompass a broad region extending from zinc finger 5 through the C terminus (5ZFC) (39). To more narrowly define the BRCA1-binding determinants on ZBRK1, we examined the contribution of individual zinc fingers 5–8 as well as sequences within the ZBRK1 C terminus to BRCA1 binding using a yeast two-hybrid interaction assay. To this end, individual substitution and truncation mutations within ZBRK1 5ZFC were translationally fused to the GAL4 transactivation domain and tested for their respective abilities to bind to the ZBRK1-interaction domain on BRCA1 (amino acids 341–748), translationally fused to the GAL4 DNA-binding domain in yeast (Fig. 3). Corresponding \(\beta\)-galactosidase activities identified critical determinants of BRCA1 interaction on ZBRK1 to include the C terminus as well as zinc fingers 7 and 8 (Fig. 3). Deletion of only 9 amino acids from the C terminus significantly compromised BRCA1 binding, indicating that the unique C terminus on ZBRK1 is required in its entirety for efficient interaction with BRCA1 (Fig. 3). This observation sug-
The ZBRK1 5ZFC and KRAB repression domains function in a BRCA1-dependent and BRCA1-independent manner, respectively. **A**, Brca1+/+ and Brca1−/− MEF cells (39) as indicated were transfected with 100 ng of pG5TK-LUC without or with the indicated nanogram amounts of GAL4-ZBRK1 5ZFC. **B**, Brca1−/− MEF cells were transfected with 100 ng of pG5TK-LUC and 20 ng of either pM (expressing the GAL4 DNA-binding domain (DBD) alone) or GAL4-ZBRK1 5ZFC, respectively, without or with the indicated nanogram amounts of pCS2+-eBRCA1 expressing wild-type human BRCA1. **C**, Brca1+/+ and Brca1−/− MEF cells as indicated were transfected with 100 ng of pG5TK-LUC without or with the indicated nanogram amounts of GAL4-ZBRK1 KRAB. **A−C**, relative luciferase activities were calculated as described in the legend to Fig. 4. **D**, schematic representation of the GAL4 DNA-binding domain, the GAL4-ZBRK1 5ZFC and GAL4-ZBRK1 KRAB chimeras, and the pG5TK-Luc reporter template used in transfection assays.

Fig. 5. The ZBRK1 5ZFC and KRAB repression domains function in a BRCA1-dependent and BRCA1-independent manner, respectively. **A**, Brca1+/+ and Brca1−/− MEF cells (39) as indicated were transfected with 100 ng of pG5TK-LUC without or with the indicated nanogram amounts of GAL4-ZBRK1 5ZFC. **B**, Brca1−/− MEF cells were transfected with 100 ng of pG5TK-LUC and 20 ng of either pM (expressing the GAL4 DNA-binding domain (DBD) alone) or GAL4-ZBRK1 5ZFC, respectively, without or with the indicated nanogram amounts of pCS2+-eBRCA1 expressing wild-type human BRCA1. **C**, Brca1+/+ and Brca1−/− MEF cells as indicated were transfected with 100 ng of pG5TK-LUC without or with the indicated nanogram amounts of GAL4-ZBRK1 KRAB. **A−C**, relative luciferase activities were calculated as described in the legend to Fig. 4. **D**, schematic representation of the GAL4 DNA-binding domain, the GAL4-ZBRK1 5ZFC and GAL4-ZBRK1 KRAB chimeras, and the pG5TK-Luc reporter template used in transfection assays.

suggests that the overall conformation of the C terminus is likely to be important for BRCA1 interaction. Whereas ZBRK1 zinc fingers 7 and 8 are critical for BRCA1 interaction, zinc fingers 5 and 6 do not appear to contribute to BRCA1 binding (Fig. 3). Taken together, these results indicate that important BRCA1-binding determinants on ZBRK1 include those that also modulate its sequence-specific DNA binding activity in both a positive (zinc finger 7) and negative (zinc finger 8 and the C terminus) manner.

**The BRCA1-binding Domain on ZBRK1 Functions as an Autonomous BRCA1-dependent Transcriptional Repression Domain**—The fact that BRCA1 contacts ZBRK1 through surfaces that are not essential but nonetheless modulatory with respect to DNA binding suggests several potential mechanisms by which BRCA1 might mediate transcriptional repression by ZBRK1. First, BRCA1 could mediate ZBRK1 repression, at least in part, by modulating its sequence-specific association with DNA. This possibility is currently under investigation. Alternatively, or additionally, BRCA1 could mediate repression by DNA-bound ZBRK1. This possibility is supported by our previous observation that clinically validated missense mutations within the BRCA1 C terminus that do not disrupt its interaction with ZBRK1 nonetheless abrogate its ZBRK1 co-repressor activity (39). To test this possibility directly, we examined whether the BRCA1-binding domain on ZBRK1 could function as a BRCA1-dependent transcriptional repression domain when tethered to a heterologous DNA-binding domain. This approach permitted us to assess the influence of BRCA1 on the repression function of ZBRK1 independently of any effects that it might have on the DNA binding activity of ZBRK1. Accordingly, we initially tested the ability of ZBRK1 5ZFC (extending from zinc finger 5 to the C terminus) to function as an independent repression domain when linked to the GAL4 DNA-binding domain. GAL4-ZBRK1 5ZFC was transiently expressed in U2OS human osteosarcoma cells, and its influence on transcription from a pG5TK-Luc reporter template bearing five copies of the consensus GAL4 DNA-binding site upstream of the herpes simplex virus (HSV) TK gene promoter was examined. GAL4-ZBRK1 5ZFC conferred greater than 10-fold repression upon reporter gene expression in a dose-dependent manner (Fig. 4A). We also confirmed the presence of a potent KRAB repression domain within the ZBRK1 N terminus by examining its ability repress pG5TK-Luc reporter gene expression when tethered to the GAL4 DNA-binding domain (Fig. 4B). Based on quantitative immunoblot analysis of transfected
BRCA1-binding is necessary but not sufficient for ZBRK1 5ZFC repression function. To more narrowly define the boundaries of the BRCA1-dependent 5ZFC repression domain within ZBRK1, we examined a panel of ZBRK1 5ZFC truncation and substitution mutants for their respective repression activities in vivo. Relative to the intact 5ZFC domain, deletion or disruption of zinc fingers 5 or 6 individually reduced repression activity by 2–3-fold (Fig. 6, A and B), whereas individual deletion or disruption of zinc fingers 7 or 8 reduced repression activity by 3–6-fold (Fig. 6, A and B). These results indicate that zinc fingers 5–8 are all required for the integrity of the 5ZFC repression domain, although zinc fingers 7 and 8 appear to be quantitatively more important. Deletion of only 9 amino acids from the C terminus of ZBRK1 severely compromised the 5ZFC repression domain (5ZFC) that encompasses the BRCA1-binding domain (data not shown). Nonetheless, this result indicates the presence within ZBRK1 of two portable transcriptional repression domains, an N-terminal KRAB domain and a novel C-terminal transcriptional repression domain, deletion or disruption of zinc fingers 7 or 8 reduced repression activity by 2–3-fold (Fig. 6, A and B), whereas individual deletion or disruption of zinc fingers 5 or 6 individually reduced repression activity by 2–3-fold (Fig. 6, A and B), whereas individual deletion or disruption of zinc fingers 7 or 8 reduced repression activity by 3–6-fold (Fig. 6, A and B). These results indicate that zinc fingers 5–8 are all required for the integrity of the 5ZFC repression domain, although zinc fingers 7 and 8 appear to be quantitatively more important. Deletion of only 9 amino acids from the C terminus of ZBRK1 severely compromised the 5ZFC repression domain (5ZFC), indicating that the entire C terminus is likely to be important for 5ZFC repression activity (Fig. 6C). Thus, the 5ZFC repression domain extending from ZBRK1 zinc finger 5 through the C terminus appears to constitute an intact repression domain that cannot be further delimited. This analysis also reveals an imperfect correlation between BRCA1 binding and transcriptional repression by the 5ZFC repression domain. Thus, disruption of ZBRK1 zinc fingers 7 or 8 or truncation of the C terminus severely compromised BRCA1 binding (Fig. 3) and transcriptional repression (Fig. 6). By contrast, disruption of zinc fingers 5 or 6, which are
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not required for BRCA1 binding (Fig. 3), nonetheless significantly compromised transcriptional repression (Fig. 6, A and B). On this basis we conclude that BRCA1 binding is necessary but not sufficient for ZF2C repression function. This suggests that ZBRK1 zinc fingers 5 and 6 may possibly contact an additional co-repressor(s).

The BRCA1-dependent ZF2C Repression Domain Is Histone Deacetylase-dependent and Promoter-specific—Previously, BRCA1 has been shown to interact through its C-terminal BRCT repeats with histone deacetylases (HDACs) 1 and 2 (57). HDACs remove acetyl groups from lysine residues on histone tails and thus promote the formation of transcriptionally repressive chromatin. To determine the contribution of HDAC activity to repression mediated by ZBRK1 ZF2C, we tested the effect of the selective HDAC inhibitor trichostatin A (TSA) on repression mediated by GAL4-ZBRK1 ZF2C. TSA largely reversed GAL4-ZBRK1 ZF2C repression in U2OS cells, implicating HDAC activity in this process (Fig. 7).

Like the ZBRK1 ZF2C repression domain, KRAB repression domains function through HDACs as well as through histone methyltransferases and heterochromatin proteins (41–47). This prompted us to comparatively examine the promoter specificities of the ZBRK1 ZF2C and KRAB repression domains. The ZBRK1 N-terminal KRAB domain repressed transcription potently from each of three different RNA polymerase II promoters tested: the SV40 major late, the HSV TK, and the human small nuclear ribonucleoprotein N (SNRPN) promoters (Fig. 8B). By contrast, ZBRK1 ZF5FC repressed transcription potently from the HSV TK promoter, moderately from the SV40 promoter, and not at all from the SNRPN promoter (Fig. 8A).

These results indicate that the ZBRK1 KRAB and ZF2C repression domains can be distinguished functionally not only by their requirement for BRCA1 but also on the basis of their promoter specificities; whereas the ZBRK1 KRAB repression domain exhibits broad promoter selectivity, the BRCA1-dependent ZF2C repression domain exhibits a more restricted promoter bias.

**DISCUSSION**

A central question regarding the role of BRCA1 in transcription control concerns the means by which it mediates gene-specific regulation in the absence of sequence-specific DNA binding activity. In part, this question has been answered by the identification of a growing number of sequence-specific DNA-binding transcription factors with which BRCA1 physically and functionally interacts. In this regard, our previous identification of ZBRK1 as a BRCA1-dependent transcriptional repressor provided a molecular basis to link BRCA1 directly to the regulation of GADD45a gene expression (39). Other work has rendered it clear that the BRCA1 regulation of GADD45a gene transcription is likely to be complex and mediated not only through ZBRK1 but other trans-acting factors, including OCT1 and NF-YA (21, 32, 39). Presently, however, little is known regarding the mechanism(s) by which BRCA1 mediates sequence-specific transcriptional control through the various transcription factors with which it interacts. Here we have investigated the functional interaction between ZBRK1 and BRCA1 in an effort to understand the role of BRCA1 in sequence-specific transcriptional repression.

Our studies suggest that BRCA1 mediates ZBRK1 repression, at least in part, through its targeted recruitment to a novel C-terminal repression domain (ZF2C) within ZBRK1. Structurally, this repression domain comprises the last four zinc fingers and the unique C-terminal extension of ZBRK1. The identification of ZF2C as a discrete functional domain was revealed by its ability to repress transcription when tethered to a heterologous DNA-binding domain (Fig. 4) and its functional resistance to truncation or substitution mutagenesis (Fig. 6).

Importantly, we demonstrated that ZF2C repression function is dependent upon BRCA1; genetic ablation of Brca1 or disruption of BRCA1-binding determinants on ZF2C similarly abrogates the repression function of this domain (Figs. 5 and 6). The functional contribution of this domain to BRCA1-dependent ZBRK1 repression is reflected by our previous observation that deletion of the ZBRK1 C terminus abrogates ZBRK1 repression through natural ZBRK1-response elements (39). However, whereas BRCA1 binding is necessary, our studies here suggest that it is not sufficient for ZF2C-directed repression. First, BRCA1 binding and repression determinants within this domain can be separated, suggesting a possible functional requirement for a co-repressor(s) in addition to BRCA1 (Figs. 3 and 6). Second, ZF2C-directed repression is HDAC-dependent (Fig. 7). Thus, we propose that the ZBRK1 ZF2C repression domain recruits BRCA1 as part of a higher order repression complex that minimally includes an associated HDAC activity. Targeted attempts to identify the functionally relevant BRCA1-associated co-repressor activities are currently underway.

Our work further reveals unique insight into the structural and functional organization of ZBRK1, a member of the KRAB-ZFP family. The -220 members of this family make up a significant proportion of the transcription factor complement of the human proteome and are believed to occupy important regulatory roles in development, differentiation, and transformation (41, 58–63). Despite their potential biological significance, our current understanding of the mechanisms through
which individual members of this protein family function is still rather limited. Thus, although considerable mechanistic insight into the repression function of the KRAB domain has been revealed in recent years (41–47), comparatively little is known regarding the role of KRAB domain-associated zinc fingers in transcriptional repression apart from their presumed role in sequence-specific DNA binding. In part, this gap in knowledge derives from the limited availability of KRAB-ZFP target sequences with which structure-function analyses may be carried out. In the case of several KRAB-ZFPs whose corresponding binding site sequences have been identified, an additional function(s) for individual zinc fingers beyond DNA binding seems implicit. For example, based on the observation that one C2H2 zinc finger can bind to ~3 bp of DNA (52–54), the established target sequence lengths of 5 and 27 bp, respectively, for the 8- and 10-fingered ZNF202 and KS1 proteins are incompatible with DNA contact mediated by every zinc finger (61, 63). Our previous derivation of a consensus binding sequence for ZBRK1 has permitted us here to dissect a long array of ZBRK1 zinc fingers to both sequence-specific DNA-binding and transcriptional repression. Our studies reveal the ZBRK1 zinc fingers to be multifunctional in nature, with dedicated roles in binding DNA, BRCA1, or both.

First, zinc fingers 1–4 are essential for DNA binding activity and compose the minimal DNA-binding domain under moderate conditions of ionic strength (Fig. 1). Zinc finger 5 appears to be a critical and context-dependent DNA-binding zinc finger; this finger represents the extent of the minimal DNA-binding domain under more stringent conditions (Fig. 1). Zinc fingers 6 and 7 are not essential for DNA binding but nonetheless enhance the stability of DNA binding (Figs. 1 and 2). Finally, zinc finger 8 (along with the C terminus) is dispensable for, and may possibly destabilize, DNA binding mediated by zinc fingers 1–7 (Fig. 1). Taken together, these findings reveal the ZBRK1 zinc fingers to compose at least two functional classes: those that make minimal essential contacts with DNA (fingers 1–4) and those that modulate the stability of these contacts (fingers 5–8). Importantly, zinc fingers 5–8 that modulate ZBRK1 DNA binding activity also represent critical determinants of repression by DNA-bound ZBRK1 through association with co-repressors, including BRCA1. These findings thus extend the established role of KRAB-zinc fingers to include protein-protein interactions critical for transcriptional repression, and also identify within ZBRK1 dual specificity zinc fingers with twin roles in DNA-binding and transcriptional repression.

Our work advances the understanding of DNA recognition by KRAB-ZFPs in several respects. First, we provide further empirical evidence to support predictive models for C2H2 zinc finger-DNA recognition. Structural studies of 3- and 5-fingered proteins in complex with DNA have indicated that individual zinc fingers bind to ~3 bp of DNA (52–54). Based on this model, five of the eight ZBRK1 zinc fingers would be predicted to bind to its 15-bp recognition sequence. In fact, DNA-binding analyses revealed that under mild conditions of ionic strength, the first four ZBRK1 zinc fingers are sufficient to confer stable binding to its consensus sequence. However, more stringent conditions unmasked a requirement for the fifth finger, consistent with the aforementioned structural models. Second, our identification of ZBRK1 zinc finger 5 as a critical and context-
dependent DNA-binding determinant could clarify recent issues concerning selectivity among KRAB-ZFPs that recognize overlapping DNA-binding site sequences. In this regard, a four-fingered KRAB-ZFP called SZF1 was recently shown to recognize a DNA-binding site in common with ZBRK1 (64). The observation that SZF1 and ZBRK1 exhibit overlapping DNA-binding specificity in vitro raises the possibility that these proteins might compete for a common binding site(s) in vivo (64). This, in turn, could have significant implications for the biological regulation of target gene transcription by each of these proteins. However, as we show here, ZBRK1 zinc finger 5 is a critical DNA-binding determinant under more stringent conditions of increased ionic strength and also increased non-specific competitor concentrations in vitro (Fig. 1 and data not shown). Because these conditions are more likely to approximate those of the cellular milieu, in which target site location must be achieved in the presence of a vast excess of like and unlike DNA sequences, ZBRK1 zinc finger 5 could represent a critical determinant of target site selection in vivo. Beyond zinc finger 5, zinc fingers 6 and 7 through enhanced affinity and specificity of protein-protein interactions could further influence ZBRK1 target site selectivity.

Our identification within ZBRK1 of a C-terminal BRCA1-dependent repression domain in addition to the N-terminal KRAB domain presents the first demonstration of a KRAB-ZFP harboring two independent repression domains. More importantly, the presence of two inherent repression domains could have important implications for gene-specific transcriptional control by ZBRK1. As we show here, the KRAB and C-terminal repression domains within ZBRK1 can be distinguished functionally on the basis of their respective requirements for BRCA1; the C-terminal repression domain is BRCA1-dependent, whereas the KRAB domain is not. This functional distinction may in part underlie the unique promoter specificities of the two repression domains. Whereas the KRAB repression domain exhibits broad promoter specificity, the BRCA1-dependent repression domain exhibits a more restricted promoter bias. Thus, the relative contribution of the BRCA1-dependent repression domain to overall ZBRK1 repression may vary among different ZBRK1 target promoters, effectively expanding the regulatory potential available at ZBRK1 target genes. It will be of interest in future studies to determine whether and how these discrete repression domains function synergistically to confer ZBRK1 repression.

Finally, although our work suggests that BRCA1 mediates repression by DNA-bound ZBRK1, we cannot exclude the additional possibility that BRCA1 also mediates ZBRK1 repression, at least in part, by modulating it sequence-specific DNA binding activity. Our observation that the BRCA1-binding surface on ZBRK1 includes zinc fingers that modulate its DNA binding activity in both a positive (zinc finger 7) and negative (zinc finger 8) manner is consistent with this possibility, and studies are currently underway to address this important issue. Nonetheless the studies presented here shed new light on the functional organization of ZBRK1 as a model KRAB-ZFP and further define the role of BRCA1 in sequence-specific transcription control.

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Tetrameric Oligomerization Mediates Transcriptional Repression by the BRCA1-dependent Kruppel-associated Box-Zinc Finger Protein ZBRK1

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The Kruppel-associated box (KRAB)-zinc finger protein ZBRK1 has been implicated in the transcriptional regulation of DNA damage-response genes that function in cell growth control and survival. Recently, we described a novel BRCA1-dependent C-terminal transcriptional repression domain (CTRD) within ZBRK1, the mode of repression of which is functionally distinguishable from that of the N-terminal KRAB repression domain within ZBRK1. The identification of BRCA1 binding-competent but repression-defective CTRD mutants further revealed that BRCA1 binding is not solely sufficient for ZBRK1 CTRD function. During an unbiased search for possible co-regulators of the CTRD, we identified ZBRK1 itself, suggesting that ZBRK1 can oligomerize through its CTRD. Herein we explore the physical and functional requirements for ZBRK1 oligomerization in ZBRK1-directed transcriptional repression. Protein interaction analyses confirmed that ZBRK1 can homo-oligomerize both in vitro and in vivo and further mapped the ZBRK1 oligomerization domain to the CTRD C terminus. Biochemical analyses, including protein cross-linking and gel filtration chromatography, revealed that ZBRK1 homo-oligomers exist as tetramers in solution. Functionally, ZBRK1 oligomerization facilitates ZBRK1-directed transcriptional repression through ZBRK1 response elements; requirements for oligomerization-dependent repression include the ZBRK1 CTRD and KRAB repression domains but not the DNA binding activity of ZBRK1. These observations suggest that higher order oligomers of ZBRK1 may assemble on target ZBRK1 response elements through both protein-DNA and CTRD-dependent protein-protein interactions. These findings thus reveal an unanticipated dual function for ZBRK1 in both DNA binding-dependent and -independent modes of transcriptional repression and further establish the CTRD as a novel protein interaction surface responsible for directing homotypic and heterotypic interactions necessary for ZBRK1-directed transcriptional repression.

ZBRK11 (zinc finger and BRCA1-interacting protein with a KRAB domain-1) is a member of the Kruppel-associated box-zinc finger protein (KRAB-ZFP) family of transcriptional repressors (1, 2). The ~300 members of this family comprise a significant proportion of the transcription factor complement of the human genome and are believed to occupy important regulatory roles in development, differentiation, and transformation (3–9). In addition to its identification as the product of a gene up-regulated in senescent fibroblasts, ZBRK1 was independently identified as a BRCA1-dependent transcriptional repressor of the gene encoding GADD45a, a functionally important DNA damage-response effector that functions in G1/M cell cycle checkpoint control and the maintenance of genomic stability (2, 10, 11). Previous functional analyses revealed that ZBRK1 represses GADD45a gene transcription through an intron 3 DNA-binding site in a manner dependent upon direct interaction with BRCA1 and further suggested a model in which ZBRK1 and BRCA1 function coordinately to repress GADD45a gene transcription in the absence of genotoxic stress (2). Evidence to support this model derives from the recent observation that 2,1 (Bax), and GADD153 (2). The observation suggests a potentially broader role for BRCA1 and ZBRK1 in the coordinate transcriptional control of functionally diverse DNA damage response genes. As part of our initial effort to explore the mechanism by which BRCA1 mediates sequence-specific transcriptional repression through ZBRK1, we recently identified and functionally characterized a novel C-terminal transcriptional repression domain (CTRD) within ZBRK1 (13). Structurally, the CTRD comprises the last four zinc fingers and an atypical C-terminal extension within the eight-fingered ZBRK1 protein (13). Functionally, the CTRD represses transcription in a BRCA1-dependent, histone deacetylase-dependent, and promoter-specific manner and is thus distinguishable from the N-terminal KRAB repressor.

The abbreviations used are: ZBRK1, zinc finger and BRCA1-interacting protein with a KRAB domain-1; BF, broken finger; CTRD, C-terminal transcription repression domain; DES, disuccinimidyl suberate; KRAB, Kruppel-associated box; MBP, maltose-binding protein; TK, thymidine kinase; ZFP, zinc finger protein; ZRE, ZBRK response element.

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Oligomerization Mediates ZBRK1 Transcriptional Repression

During the course of our functional dissection of ZBRK1, we identified CTRD mutants that are competent for BRCA1 binding but nonetheless defective for transcriptional repression (13). This observation suggests that BRCA1 binding is necessary but not sufficient for the ZBRK1 CTRD repression function and implies a role for additional co-regulators of the CTRD. During an unbiased search for CTRD-interacting proteins and, therefore, potential CTRD co-regulators, we unexpectedly identified ZBRK1 itself, suggesting that ZBRK1 can homo-oligomerize through its CTRD. Herein, we explore the physical and functional requirements for ZBRK1 oligomerization in ZBRK1-directed repression. Our results reveal the ZBRK1 CTRD to be a novel interaction surface responsible for directing both homotypic and the heterotypic interactions essential for ZBRK1 repression and further suggest an unanticipated dual role for ZBRK1 in both DNA binding-dependent and -independent modes of transcriptional repression. We discuss the implications of these findings for BRCA1-dependent ZBRK1 repression.

**EXPERIMENTAL PROCEDURES**

**Plasmid Construction and Mutagenesis**—Plasmid pMAL-C2-TEV ZBRK1 ΔK for expressing maltose-binding protein (MBP)-ZBRK1 ΔK in *Escherichia coli* has been described previously (13). pCS2+–FLAG-ZBRK1 for in vitro transcription/translation and mammalian expression of FLAG-epitope-tagged ZBRK1 was constructed by subcloning a BamHI-HindIII fragment carrying sequences encoding an amino-terminally FLAG-tagged ZBRK1 from pFastbac-FLAG-ZBRK1 into pCS2+ (14). pCS2+–FLAG-ZBRK1 broken finger (BF) and KrAB domain (Δ12,Δ3) mutants were generated by site-directed mutagenesis using the QuikChange II site-directed mutagenesis kit following the manufacturer’s instructions. LaJolla ZBRK1, expressing B-galactosidase under control of the SV40 promoter. Forty-eight hours post-transfection, 2.5 X 10^6 transfected U2OS cells were harvested and lysates were analysed for luciferase activity using the luciferase assay system (Promega) and for β-galactosidase activity using the Galacto-Light Plus chemiluminescent reporter assay (BD Biosciences). Each transfection was performed in a minimum of three times in duplicate wells.

**Protein Expression and Purification**—MBP-ZBRK1 ΔK, MBP-ZBRK1 CTRD, and MBP proteins were each expressed in and purified from *E. coli* strain BL21 Star (DE3) (Invitrogen). The procedures for protein induction, extraction, and purification of MBP-ZBRK1 ΔK have been described previously (13). For MBP-ZBRK1 CTRD and MBP, extracts were grown at 37 °C to an A600 of 0.6. Isopropyl-β-D-thiogalactopyranoside was added to a final concentration 0.3 mM, and the cells were transferred to 25 °C for another 3.5 h. Cells were pelleted, washed once with phosphate-buffered saline, and then resuspended in MBP binding buffer (20 mM HEPES, pH 7.5, 1 mM EDTA, 500 mM NaCl, and 10 mM β-mercaptoethanol) supplemented with protease inhibitors (0.4 mg/ml aprotinin, 0.5 μg/ml leupeptin, 0.05 μg/ml pepstatin, 0.5 mM phenylmethylsulfonyl fluoride, and 0.5 mM benzamidine-HCl). Resuspended cells were frozen and thawed once followed by sonication (three times for 1 min each on ice) and clarification by centrifugation at 80,000 × g for 30 min. MBP-ZBRK1 CTRD and MBP proteins were purified from resuspended lysates by affinity chromatography on MBP-rose resin (New England Biolabs, Beverly, MA). Briefly, clarified lysates were incubated with amylase resin in batch for 1 h at 4 °C, washed with MBP binding buffer, and eluted with MBP binding buffer containing 0.6% maltose in column buffer.

**MBP Binding Assays**—MBP-ZBRK1 ΔK was purified from clarified bacterial culture (15 μl) with cell extract (15 μl) in batch for 30 min at 25 °C, followed by washing two times for 5 min each in 25 °C MBP binding buffer supplemented with protease inhibitors and three times for 5 min each in 25 °C in lysis 300 buffer (50 mM Tris-HCl, 300 mM NaCl, 5 mM EDTA, 0.1% Nonidet P-40, and 10 mM β-mercaptoethanol) supplemented with protease inhibitors. FLAG-ZBRK1 or its mutant derivatives were labeled with 5-iodoacetamide (TST SP6 quick coupled transcription/translation system; Promega) and incubated with amylase-immobilized MBP-ZBRK1 ΔK in lysis 300 buffer for 1 h at 4 °C. Binding reactions were washed with Lysis 300 buffer four times for 10 min each at 4 °C and subsequently boiled in 5X SDS sample buffer. Eluates were resolved by 10% SDS-PAGE and visualized by PhosphorImager analysis (Amersham Biosciences).

**Co-immunoprecipitation Analyses**—U2OS cells at 70% confluency in 10-cm plates were transfected with 12 μg of DNA using the following expression plasmids (with amounts in parentheses): pCS2+–T7-ZBRK1 (6 μg) and pCS2+–FLAG-ZBRK1 (6 μg); pCS2+–T7-ZBRK1 (6 μg) and pCS2+–FLAG-ZBRK1 ΔC1 mutants (3 μg); pCS2+–T7-ZBRK1 ΔC2 mutants (4.5 μg), or pCS2+–FLAG-ZBRK1 ΔC3 mutants (2.5 μg) (the total amount of DNA was fixed by supplementing these transfections with pCS2+). Thirty-six hours post-transfection, 2.5 X 10^6 transfected U2OS cells were harvested and resuspended in lysis buffer (150 mM NaCl, 10 mM P-mercaptoethanol) supplemented with protease inhibitors and mixed with 150T buffer (50 mM Tris-HCl, pH 7.5, 150 mM NaCl, 5 mM EDTA, and 0.1% Nonidet P-40) by sonication at four intervals of 7 s each using a Branson 450 sonifier (Branson Ultrasonics) at the lowest setting. The lysate was clarified by centrifugation at 20,000 × g for 20 min at 4 °C. Clarified lysates were subjected to co-immunoprecipitation with either 1 μg of anti-T7 antibody (Novagen, Madison, WI) or 5 μg of anti-FLAG M2 monoclonal antibody (Sigma-Aldrich) for 4 h at 4 °C. Immunocomplexes were precipitated with 20 μl of protein A-Sepharose (Amersham Biosciences) (for anti-T7 antibody immunoprecipitation) or protein G-Sepharose (Roche Diagnostics) (for anti-FLAG antibody immunoprecipitation) for 1 h at 4 °C. Immunoprecipitates were pelleted and washed with lysis 150T buffer at least four times for 10 min each at 4 °C. Immunoprecipitated proteins were eluted by boiling in 2X Laemmli sample buffer for 3 min, and eluates were subsequently resolved by SDS-PAGE (10% gel) and processed for immunoblot analysis.

**Chemical Cross-linking Assays**—U2OS cells at 70% confluency on 10-cm plates were transfected with either 12 μg of pcS2+–FLAG-ZBRK1 ΔK or 6 μg of pCS2+–FLAG-ZBRK1 ΔC1 in combination with 6 μg of pCS2+. Thirty-six hours post-transfection, cells were harvested and lysed in 300 μl of lysis 150H buffer (20 mM HEPES, pH 7.5, 150 mM NaCl, 5 mM EDTA, and 0.1% Nonidet P-40) by sonication at four intervals of 7 s each using a Branson 450 sonifier (Branson Ultrasonics) at the lowest setting. The lysate was clarified by centrifugation at 20,000 × g for 20 min at 4 °C. Clarified lysates (36 μl) were subjected to chemical cross-linking by the addition of disuccinimidyl suberate (DSS) (25 mM) to a final concentration of 2.25 mM. Cross-linking reactions were performed at 25 °C and terminated by the addition of 2X Laemmli sample buffer followed by boiling for 3 min (15). Cross-linked products were resolved by SDS-PAGE (10% gel) and processed for immunoblot analysis.

**Cell Culture, Transfections, and Reporter Assays**—U2OS human osteosarcoma cells were cultured as described (15). For transient reporter assays, 2 X 10^5 cells were seeded per well of a 6-well plate 24 h before transfection. When 70% confluent, the cells were transfected with PolyGene 6 (Roche Diagnostics) following the manufacturer’s instructions. The amounts of expression and reporter plasmids utilized in each experiment are indicated in the legends to Figs. 1-4. A total of 2 μg, respectively, of DNA per well was transfected when pG5,TK-Luc or pZRE-Luc were used as reporter templates. Each transfection also included an internal control plasmid, pCH110 (15), expressing β-galactosidase under control of the SV40 promoter. Forty-eight hours post-transfection, cells were harvested and lysed in 250 μl/well reporter lysis buffer (Promega, Madison, WI). Transfected cell lysates (20 μl) were analyzed for luciferase activity using the luciferase assay system (Promega) and for β-galactosidase activity using the Galacto-Light Plus chemiluminescent reporter assay (BD Biosciences). Each transfection was performed in a minimum of three times in duplicate wells.
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**RESULTS**

**ZBRK1 Homo-oligomerizes in Vitro**—To identify novel interaction partners and potential co-regulators of the ZBRK1 CTRD, we employed two independent experimental approaches. First, we used the CTRD (amino acids 319–532 of ZBRK1) as a bait protein to screen a human fetal brain cDNA library by a yeast-two-hybrid assay. From among ~1 × 10^6 independent clones screened, we identified ZBRK1 itself as a CTRD interaction partner (data not shown). Second, we expressed a ZBRK1 KRAB domain deletion derivative (ZBRK1 ΔK; amino acids 144–532 of ZBRK1) as an MBP chimera in E. coli, immobilized the fusion protein on amylose resin, and incubated with in vitro translated 35S-labeled wild-type (WT) ZBRK1 (A and B) or its indicated ΔC (A) or BF (B) derivatives. Following extensive washing, bound proteins were eluted in Laemmli sample buffer, resolved by SDS-PAGE (10% gel), and visualized by PhosphorImager analysis. Input represents 5% of the total in vitro translated protein used in binding reactions.

**ZBRK1 Homo-oligomerizes in Vivo**—To determine whether ZBRK1 can homo-oligomerize in vivo, we first examined the ability of ZBRK1 derivatives independently tagged with either T7 or FLAG epitopes to reciprocally co-immunoprecipitate one another following their transient over-expression in U2OS human osteosarcoma cells. T7-tagged ZBRK1 could be efficiently substituted mutagenesis revealed that each "broken finger" mutant (13) bound MBP-ZBRK1 ΔK comparably to wild-type ZBRK1, thus revealing that the CTRD zinc fingers are individually dispensable for ZBRK1 oligomerization (Fig. 1B). Taken together, these results demonstrate that ZBRK1 can homo-oligomerize in vitro through a direct interaction involving its CTRD C terminus.

**Fig. 1.** ZBRK1 homo-oligomerizes in vitro through its CTRD C terminus. A and B, top, schematic representation of FLAG-tagged wild-type (WT) ZBRK1, ZBRK1 carboxyl-terminal deletion (AC) and broken finger (BF) derivatives, and MBP-ZBRK1 ΔK (the KRAB and CTRD repression domains as well as numbered zinc fingers are indicated). Bottom, purified recombinant MBP-ZBRK1 AK was immobilized on amylose resin and incubated with in vitro translated 35S-labeled wild-type (WT) ZBRK1 (A and B) or its indicated ΔC (A) or BF (B) derivatives. Following extensive washing, bound proteins were eluted in Laemmli sample buffer, resolved by SDS-PAGE (10% gel), and visualized by PhosphorImager analysis. Input represents 5% of the total in vitro translated protein used in binding reactions.
ZBRK1 homo-oligomerizes in vivo through its CTRD C terminus. A and B, T7 and FLAG-tagged wild-type ZBRK1 proteins were ectopically expressed either alone or together in U2OS cells, and whole cell extracts from transfected cells were subjected to immunoprecipitation (IP) using antibodies specific for either the FLAG (A) or T7 (B) epitopes. Immunoprecipitates were resolved by SDS-PAGE (10% gel) and processed by immunoblot analysis using both T7 and FLAG epitope-specific antibodies as indicated. Input represents 10% of the total whole cell extract subjected to immunoprecipitation. C and D, T7-tagged wild-type (WT) ZBRK1 was ectopically expressed in U2OS cells along with FLAG-tagged WT ZBRK1 or its indicated ΔC (C) or BΔ (D) derivatives. Whole cell extracts from transfected cells were subjected to immunoprecipitation (IP) using antibodies specific for the T7 epitope. Immunoprecipitates were resolved by SDS-PAGE (10% gel) and processed by immunoblot analysis using both T7 and FLAG epitope-specific antibodies as indicated. Input represents 10% of the total whole cell extract subjected to immunoprecipitation.

abilities to be co-immunoprecipitated along with T7-tagged wild-type ZBRK1 following their transient co-expression in U2OS cells. Consistent with the results of in vitro binding analyses, these in vivo binding studies revealed the CTRD C terminus to be critical and the CTRD zinc fingers to be dispensable for ZBRK1 self-association. Thus, deletion of only nine amino acids from the ZBRK1 C terminus largely abolished ZBRK1 self-association (Fig. 2C), whereas individual ZBRK1 derivatives bearing targeted disruptions of CTRD zinc fingers 5–8 bound to T7-tagged ZBRK1 as efficiently as wild-type ZBRK1 (Fig. 2D). Taken together, these results demonstrate that ZBRK1 can homo-oligomerize both in vitro and in vivo and furthermore delimit the oligomerization interface to the CTRD C terminus.

The ZBRK1 CTRD Mediates Tetrameric Oligomerization—To characterize the oligomeric state of ZBRK1 in vivo, we first employed the chemical cross-linker DSS to covalently capture ZBRK1 protein complexes present in extracts of U2OS cells transfected with FLAG-tagged ZBRK1 (16). Cross-linked samples were analyzed by SDS-PAGE and immunoblot analysis using FLAG epitope tag-specific antibodies. In the presence of DSS, a significant proportion of MBP-ZBRK1 CTRD migrated with an apparent molecular mass of ~270 kDa, consistent with the size of an MBP-ZBRK1 CTRD tetramer (Fig. 3B). Importantly, no cross-linked products were observed when MBP alone was treated with DSS, indicating that DSS-induced formation of a high molecular mass MBP-ZBRK1 CTRD complex is strictly dependent upon the ZBRK1 CTRD (Fig. 3B). Furthermore, immunoblot analysis using ZBRK1-specific antibodies confirmed that the high molecular mass complex observed upon DSS cross-linking of highly purified MBP-ZBRK1 CTRD is derived from MBP-CTR oligomerization (data not shown).

As an independent approach to determine the stoichiometry of ZBRK1 oligomerization through its CTRD, we examined the respective gel filtration profiles of ectopically expressed wild-type ZBRK1 and its oligomerization-defective mutant (ΔC1) present in extracts of transfected U2OS cells. Immunoblot analysis of column fractions using a FLAG epitope-specific antibody revealed a peak of wild-type ZBRK1 protein in fractions corresponding to an apparent molecular mass of ~240 kDa, most consistent with the size of a ZBRK1 tetramer (Fig. 3C). We also noted a minor peak of wild-type ZBRK1 protein in a fraction corresponding to an apparent ZBRK1 monomer of 58 kDa. By contrast, an oligomerization-defective ZBRK1 derivative (ΔC1) peaked exclusively in column fractions corresponding to an apparent monomer of ~57 kDa (Fig. 3C). Taken together, these results indicate that ZBRK1 tetramerizes both in vitro and in vivo in a manner that is strictly dependent upon the integrity of its CTRD C terminus.

ZBRK1 Homo-oligomerization Potentiates ZBRK1-directed Transcriptional Repression—To explore the functional consequence of ZBRK1 homo-oligomerization, we initially exploited our previous observation that the ZBRK1 CTRD can function as an autonomous transcriptional repression domain when tethered to a heterologous DNA-binding domain from the yeast transcription factor GAL4 (13). First, we examined the ability of full-length ZBRK1 to potentiate transcriptional repression...
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FIG. 3. ZBRK1 tetramerization requires its CTRD C terminus. A, schematic representation of FLAG-tagged wild-type ZBRK1 and its oligomerization-defective ΔC1 deletion derivative. U2OS cells were transfected with expression vectors for either FLAG-tagged WT ZBRK1 or its ΔC1 deletion derivative as indicated. Whole cell extracts from transfected cells were subjected to cross-linking with 2.25 mM DSS for 0, 1.5, or 3 min as indicated, and cross-linking reactions were terminated with Laemmli sample buffer. Cross-linked proteins were resolved by SDS-PAGE (10% gel) and visualized by immunoblot analysis using FLAG epitope-specific antibodies. Arrowheads indicate un-cross-linked monomers of FLAG-tagged ZBRK1 and its ΔC1 deletion derivative as well as cross-linked ZBRK1 complexes. Molecular weight markers (Mr) are indicated. Asterisk denotes a cross-reacting cellular protein recognized by the FLAG antibody. B, schematic representation of recombinant MBP-ZBRK1 CTD and MBP. Purified recombinant MBP-ZBRK1 CTD and MBP were subjected to cross-linking with 0.75 mM DSS for 0, 1.5, or 3 min as indicated, and cross-linking reactions were terminated with Laemmli sample buffer. Cross-linked proteins were resolved by SDS-PAGE (10% gel) and visualized by silver stain analysis. Arrowheads indicate un-cross-linked monomers of MBP-ZBRK1 CTD and MBP as well as cross-linked MBP-ZBRK1 CTD complexes. Molecular weight markers (Mr) are indicated. C, whole cell extracts from U2OS cells transfected with FLAG-tagged wild-type (WT) ZBRK1 or an oligomerization-defective mutant derivative (ΔC1) were independently subjected to Superdex 200 gel filtration chromatography. Individual column fractions were resolved by SDS-PAGE (10% gel) and analyzed for the presence of FLAG-tagged ZBRK1 by immunoblot analysis with FLAG epitope-specific antibodies. Arrowheads indicate the relative positions of marker protein peaks. Input represents 5% of the total whole cell extract subject to gel filtration chromatography.

Directed by GAL4-ZBRK1 CTD in a transient repression assay using a reporter template bearing multimerized GAL4 DNA-binding sites located upstream of the herpes simplex virus TK promoter in U2OS cells (Fig. 4A) (13). Consistent with our previous studies (13), we observed that GAL4-ZBRK1 CTD repressed transcription from this reporter template at least 10-fold in a dose-dependent manner (data not shown). To examine the influence of wild-type ZBRK1 on repression directed by GAL4-ZBRK1 CTD, we transfected a sub-optimal nanogram quantity of the GAL4-ZBRK1 CTD expression vector that supports only ~2.5-fold repression of reporter gene activity. Under these conditions, ectopic expression of wild-type ZBRK1 enhanced this level of repression up to 4-fold in a dose-dependent manner (Fig. 4B). Importantly, ZBRK1 had minimal influence on reporter activity in the presence of the GAL4 DNA-binding domain alone, confirming that the repressive effect of ZBRK1 in this assay derives from its direct recruitment to the CTD and not to DNA (Fig. 4B). This result demonstrates for the first time that, in addition to its well-documented role as a sequence-specific DNA-binding transcriptional repressor (2, 13), ZBRK1 can also function as a co-repressor independent of its inherent DNA binding activity.

Next, we examined the structural features within ZBRK1 required for its co-repressor function. Relative to wild-type ZBRK1, oligomerization-defective CTD C-terminal truncation mutants (ΔC1, ΔC2, and ΔC3) were completely defective for ZBRK1 co-repressor activity, implying a strict reliance on oligomerization for ZBRK1 to function as a co-repressor (Fig. 5A). Analysis of ZBRK1 derivatives bearing targeted disruptions of CTD zinc fingers 5–8 (BF 5–8) revealed these mutants to be partially defective for the ZBRK1 co-repressor function (Fig. 5A). Because these CTD zinc finger mutations do not disrupt ZBRK1 oligomerization (see Figs. 1 and 2) but do disrupt CTD interactions with additional co-repressors including BRCA1 (12), this result suggests that CTD-mediated recruitment of additional co-repressors also likely contributes to the ability of ZBRK1 to function as a DNA binding-independent co-repressor. Finally, we examined the contribution of the...
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As this mutation does not affect the ability of ZBRK1 CTRD to homo-oligomerize through its CTRD, this result suggests that KRAB domain-associated co-repressors also contribute to the ability of ZBRK1 to function as a DNA binding-independent co-repressor of the CTRD. Taken together, these results reveal that ZBRK1, in a DNA-binding-independent manner, can potentiate the repression function of DNA-bound CTRD through oligomerization and the consequent recruitment of co-repressors through both its KRAB and CTRD repression domains.

We next asked whether ZBRK1 CTRD-mediated oligomerization contributes to ZBRK1-directed transcriptional repression from ZBRK1 response elements (ZREs). To this end, we employed a reciprocal transient repression assay in which we examined the activity of the CTRD to potentiate ZBRK1-directed repression from a reporter template bearing multimerized ZREs. The fact that the CTRD by itself has no ZRE binding activity permitted us to assess the influence of the CTRD on ZBRK1-directed repression solely by virtue of its ability to oligomerize with DNA-bound ZBRK1. Consistent with our previous studies (2), we observed that wild-type ZBRK1 repressed transcription from a reporter template bearing multimerized ZREs located upstream of the HSV TK promoter in U2OS cells at least 10-fold in a dose-dependent manner. To examine the influence of the ZBRK1 CTRD on repression directed by wild-type ZBRK1, we transfected a sub-optimal nanogram quantity of the ZBRK1 expression vector that supports only ~2.5-fold repression of transcription from this reporter template (Fig. 6B). Under these conditions, ectopic expression of GAL4-CTRD enhanced this level of repression up to 4-fold in a dose-dependent manner (Fig. 6B). Importantly, GAL4-CTRD had minimal influence on reporter activity in the absence of ZBRK1, confirming that the CTRD potentiates ZBRK1 repression in this assay by virtue of its direct recruitment to ZBRK1 and not to DNA (Fig. 6B). Furthermore, an oligomerization-defective CTRD mutant (GAL4-CTRD AC1) bearing a nine amino acid deletion from the CTRD C terminus was completely defective in its ability to potentiate ZBRK1 repression in this assay, confirming that CTRD-mediated oligomerization is a requirement for its ZBRK1 co-repressor activity (Fig. 6C). Collectively, these experiments reveal an unanticipated dual function for ZBRK1 in both DNA binding-dependent and -independent modes of repression, the latter of which derives from the ability of ZBRK1 to homo-oligomerize through its CTRD.

DISCUSSION

The identification of ZBRK1 itself during an unbiased search for ZBRK1 CTRD co-regulators prompted us to characterize the structural requirements for and the functional consequences of ZBRK1 homo-oligomerization. Herein we demonstrate that ZBRK1 undergoes tetrameric oligomerization both in vitro and in vivo, and we further identify the ZBRK1 CTRD C terminus to be both necessary and sufficient for tetrameric oligomerization. Functionally, we demonstrate that ZBRK1 oligomerization facilitates ZBRK1-directed transcriptional repression and concomitantly identify several unique and biologically significant features of ZBRK1 relevant to its role as a KRAB-ZFP transcriptional repressor. First, by virtue of its ability to homo-oligomerize, we identify an unanticipated role for ZBRK1 as a DNA binding-independent co-repressor for transcriptional repression. Unlike its characterized role as a sequence-specific DNA-binding transcriptional repressor (2), in this regard, we identify both the KRAB and CTRD repression domains within ZBRK1 to be necessary for oligomerization-dependent repression. Whether or not these domains are sufficient for oligomerization-dependent repression remains to be established. Second, we reveal the ZBRK1 CTRD to be a novel protein interaction surface responsible for directing both homotypic and the heterotypic interactions necessary for BRCA1-dependent ZBRK1 transcriptional repression.

In general, oligomerization enhances regulatory diversity among sequence-specific DNA-binding transcription factors by expanding the number of potential response elements to which oligomers can effectively bind and/or by increasing the number and type of transcriptional regulatory domains that can effectively function at a core promoter. Presently, we do not know whether and how ZBRK1 oligomerization might influence the choice of DNA sequences recognized by ZBRK1. The 15-base pair consensus ZRE was originally identified by sampling a random pool of double-stranded oligonucleotides with a recombinant ZBRK1 derivative comprised solely of the eight ZBRK1 zinc fingers but lacking the C-terminal extension that we show here to be critical for oligomerization (2). Future studies will be required to determine whether and how oligomerization-competent forms of ZBRK1 preferentially recognize an expanded ZRE. Interestingly, our most recent studies revealed that the inherent DNA binding activity of ZBRK1 zinc fingers 1–7 on a consensus ZRE is constrained by the ZBRK1 C terminus, suggesting a potential role for the ZBRK1 oligomerization domain in regulating the...
sequence-specific association of ZBRK1 with a consensus ZRE (13). Although our previous studies clearly revealed that ZBRK1 can bind to and repress transcription from a single ZRE (2, 13), it nonetheless remains to be established whether and how ZBRK1 oligomerization might facilitate synergistic binding and/or repression from multimerized ZREs.

Our work reveals a unique and unanticipated dual function for ZBRK1 in both DNA binding-dependent and -independent modes of transcriptional repression. With respect to the latter, we demonstrate that ZBRK1 can function as a co-repressor in a manner dependent upon the integrity of its CTRD and KRAB repression domains but not upon its inherent DNA binding activity. This observation raises the possibility that the regulatory potential of ZBRK1 on certain target genes may derive from its recruitment via protein-protein rather than protein-DNA interactions. To our knowledge, this represents the first
example of a KRAB-ZFP that can function dually as both a DNA-binding repressor as well as a DNA binding-independent co-repressor. Thus, we propose that higher order oligomers of ZBRK1 may assemble on a target ZRE through protein-DNA and protein-protein interactions, the latter involving oligomerization through the CTRD C terminus.

Our work further provides unique insight into the structural and functional organization of ZBRK1 as a member of the KRAB-ZFP family. Like other KRAB-ZFPs, ZBRK1 harbors an N-terminal KRAB repression domain followed by a CxHxM zinc finger DNA-binding domain. However, ZBRK1 also harbors an atypical C-terminal extension that is generally absent from the larger family of KRAB-ZFPs. Our previous work revealed this C-terminal extension to comprise part of a novel BRCA1-dependent CTRD that also includes zinc fingers 5–8 within ZBRK1 (13). Herein, we describe an additional and unique role for the CTRD as an oligomerization interface sufficient for directing homotypic interactions critical for ZBRK1 repressor function. Within the CTRD we demonstrate that zinc fingers 5–8 are dispensable, whereas the C-terminal extension is essential for ZBRK1 oligomerization. This C-terminal extension shows no obvious sequence similarity to the SCAN domain that mediates selective oligomerization between certain KRAB-ZFPs (1, 18). However, a BLAST search of available protein databases reveals several additional KRAB-ZFPs (ZNF577, ZNF613, and FLJ12644), each of which carries an extended C terminus of variable length with sequence homology to the ZBRK1 C-terminal extension. Interestingly, these proteins exhibit the same rank order with respect to their degree of homology and physical proximity to ZBRK1 on chromosome 19q, suggesting that they may have arisen from gene duplication (19–21). It will be of future interest to determine whether and how these KRAB-ZFPs functionally oligomerize with ZBRK1 and/or BRCA1. Although zinc fingers 5–8 within the CTRD are dispensable for oligomerization, our previous studies have nonetheless shown them to be essential for mediating interactions with additional co-repressors, including BRCA1 (2, 13).

Thus, our studies collectively reveal the CTRD to be a novel protein–protein interaction surface responsible for directing both homotypic and heterotypic interactions necessary for BRCA1-dependent ZBRK1 transcriptional repression. Finally, our identification of a novel oligomerization domain within ZBRK1 suggests that the regulatory potential of ZBRK1 could extend to genes that do not contain ZREs. For example, ZBRK1 could repress transcription through its hetero-oligomerization with other sequence-specific DNA binding transcription factors through direct protein-protein interactions involving its C terminus. This, in turn, could expand the repertoire of target genes subject to coordinate transcriptional control by both ZBRK1 and BRCA1, because the repressive potential of ZBRK1 is dependent upon BRCA1 (2, 13). Consistent with this possibility, our yeast two-hybrid screen for CTRD-interacting proteins identified, in addition to ZBRK1 itself, the oligomeric transcription factors SRF and ATF-1 (22–26). Interestingly, these transcription factors control the expression of genes with diverse functions in cell growth control and survival, and future studies will seek to establish whether and how ZBRK1 physically and functionally interacts with these transcriptional regulators. Nonetheless, our work here sheds new light on the structural and functional organization of ZBRK1 as a KRAB-ZFP and further defines the multifunctional nature of its unique CTRD.

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REFERENCES
BRCA1 and BRCA2 in breast cancer

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The inheritance of an autosomal dominant allele represents an identifiable predisposing factor in about 10% of all women with breast cancer. Most of these hereditary cases can be linked to germline mutations in either of two breast cancer susceptibility genes, BRCA1 or BRCA2. Women who have a mutation in either of these genes have a cumulative lifetime risk of 60-80% and 20-40% for the development of breast and ovarian cancer, respectively. There is therefore a great need for new and effective measures for their management. Progress in our understanding of the normal biological function and regulation of BRCA1 and BRCA2 has shed new light on the molecular basis of hereditary breast cancer, and should provide a driving force for the development of diagnostic and therapeutic strategies.

BRCA1 and BRCA2 are caretaker genes whose products function in the maintenance of global genome stability—ie, they ensure that the genetic integrity of a cell is not compromised by the unscheduled loss, duplication, or rearrangement of chromosomal DNA. A persistent threat to genome integrity is DNA damage arising from ongoing metabolic processes within the cell, as well as that elicited by extrinsic agents, including radiation and certain chemicals. Unrepaired or misrepaired DNA damage can compromise chromosomal stability, allowing a cell to escape normal restrictions on its growth.

Genome integrity is ensured in part by a response system that has evolved to locate and effect the timely repair of damage to DNA. This response involves the assembly of DNA-repair protein complexes able to recognise and eliminate damage-induced lesions, and the synthesis of cell-cycle checkpoint control proteins that provide a sufficient window of opportunity to effect such repair. BRCA1 and BRCA2 occupy fundamental roles in coupling DNA damage-induced signals to downstream cellular responses, including damage repair and cell-cycle checkpoint activation.

Because the DNA damage-induced signalling pathways that converge on BRCA1 and BRCA2 are universally conserved, the genes are likely to function ubiquitously in the maintenance of genome integrity. Nonetheless, inactivation of BRCA1 or BRCA2 generally leads only to cancer of the breast or ovary. What then might constitute the molecular basis for the tissue-specific tumour suppressive properties of BRCA1 and BRCA2?

The breast and the ovary are reproductive organs that rely on hormones, including oestrogen and progesterone, for growth, differentiation, and homeostasis. According to one theory, inactivation of BRCA1 and BRCA2 renders breast and ovary susceptible to tissue-specific effects of oestrogen-induced DNA damage. Thus, inactivating mutations in BRCA1 and BRCA2 could compromise the response of breast and ovarian epithelial cells to oestrogen-induced DNA damage, thereby resulting in inefficient or error-prone DNA repair. Global genomic instability and a concomitant accrual of functionally inactivating mutations within other genes involved in breast and ovarian tumourigenesis might then ensue. Alternatively, BRCA1 might modulate hormone signalling pathways and control of cellular proliferation. BRCA1 represses the transcriptional activity of the oestrogen and progesterone receptors, and mutational inactivation of the gene could, therefore, promote epithelial cell proliferation through altered expression of hormone-responsive genes.

These two models are not mutually exclusive and could suggest a combinatorial path to breast cancer, since they invoke BRCA1-mediated and BRCA2-mediated control at two distinct steps of tumourigenesis—initiation and progression. Thus, inappropriate expression of hormone-responsive genes could promote the proliferation of transformed cells arising through inefficient or error-prone repair of oestrogen-induced DNA damage. In this way, hereditary BRCA1 and BRCA2 mutations could render breast and ovarian epithelial cells particularly susceptible to tumourigenesis through perturbation of distinct hormone-dependent pathways (figure).

This knowledge could help to treat those carrying mutations in BRCA1 and BRCA2, and might also be useful in the treatment of patients with sporadic, non-genetic breast cancers. Few mutations in BRCA1 and BRCA2 arise in sporadic breast cancers, suggesting that the perturbation of alternative pathways causes malignant disease in these cases. As caretakers of genomic integrity, BRCA1 and BRCA2 represent prime targets for therapeutic intervention—ie, targeted inactivation of BRCA1-specific and BRCA2-specific DNA-damage response pathways could render tumour cells sensitive to the genotoxic effects of radiation or chemotherapeutic agents, thereby offering the potential for improved combination therapies. In the last decade of the 20th century, BRCA1 and BRCA2 were identified and characterised. The role and regulation of their encoded products in DNA-damage response and repair, once identified, should expedite the design and implementation of strategies to delay, and ultimately to prevent, tumour formation.
Breast Cancer Susceptibility Genes

by Thomas G. Boyer and Wen-Hwa Lee

The last decade of the 20th century witnessed the identification and initial characterization of two major breast cancer susceptibility genes, BRCA-1 and BRCA-2. Studies of the encoded BRCA proteins have revealed roles in the maintenance of chromosomal stability and in DNA damage response and repair, and studies continue to illuminate further biological activities. A greater appreciation of the involvement of BRCA-1 and 2 in breast and ovary cells will increase the probability that recent advances in our understanding of their biological functions will be channeled effectively to the treatment and prevention of breast and ovarian cancer.

In the year 2002, approximately 200,000 American women will be diagnosed with breast cancer, the most common malignancy afflicting women in the United States. Among women who do not smoke, breast cancer is the primary cause of cancer-related death.

Although many factors influence a woman's lifetime risk for development of breast cancer, family history is one of the most powerful prognostic indicators. About 10% of all breast cancer cases can be linked to heritable transmission of an autosomal dominant allele. Thus a major achievement was substantiation that many of these hereditary cases could be linked to germline mutations in either of two breast cancer susceptibility genes, identified as BRCA-1 and 2.

Through linkage analysis of families affected by early-onset breast and ovarian cancer, BRCA-1 was mapped to chromosome 17q21 in 1990 and cloned 4 years later. BRCA-2 was mapped to chromosome 13q and cloned shortly thereafter. Mutations in BRCA-1 are believed to account for 60 to 80% of hereditary breast and ovarian cancer cases and up to 20% of hereditary breast cancers only. BRCA-2 mutations are linked to a similar percentage of inherited breast cancers, but in contrast to BRCA-1, they also predispose to male breast cancer.

Together, defects in these two genes account for about 40% of inherited breast cancers. Germline inactivation of one allele of either BRCA-1 or 2 is sufficient to predis-
pose a person to cancer, while cancer onset is invariably accompanied by loss of the remaining allele. Thus BRCA-1 and 2 belong to the group of tumor susceptibility genes whose encoded products normally function to suppress tumor formation.

Mutations in other known tumor susceptibility genes, such as p53, the retinoblastoma gene RB, and the adenomatous polyposis gene APC, are found in both familial and sporadic tumors. Mutations in BRCA-1 and 2, however, are rarely detected in nonhereditary breast cancers, though it has been proposed that aberrant regulation of their expression or of the activity of their products could contribute to sporadic breast cancers.

Clearly, detailed knowledge of the normal biological functions of these proteins and of their regulation will be required for a thorough appreciation of how direct or indirect functional inactivation of BRCA-1 and 2 leads ultimately to breast cancer. In this article, we begin with a description of the structural features of the BRCA proteins and then highlight recent insights into their biological role and regulation.

**Protein Structures Are Clues to Functions**

BRCA-1 is a nuclear phosphoprotein of 1863 amino acids characterized by the presence of a notable structural motif near each end. At its amino terminus, BRCA-1 harbors a zinc-binding RING finger domain, which is a set of spatially conserved cysteines and histidine residues. More than 200 RING finger proteins of diverse function are potentially encoded by the human genome, so this domain is a relatively common structural motif. Recent studies have raised the possibility that the functional diversity apparent among RING finger proteins is tied to a common enzymatic activity.

The carboxy terminal of BRCA-1 includes a series of domains that are autonomous folding units defined by conserved clusters of hydrophobic amino acids. These are called BRCA-1 C-terminal or BRCT domains, and they have been found in other proteins implicated in DNA repair and cell cycle checkpoint control. No specific cellular function has so far been ascribed to the BRCT domain, but it is likely to be a protein interaction surface.

A third region in BRCA-1 also appears to be a functionally relevant protein interaction surface, but the structure of this region has not yet been defined. The same region includes two putative nuclear localization signals.

BRCA-2 is a nuclear protein of 3418 amino acids whose most prominent feature is eight tandem copies of a repetitive sequence termed the BRC repeat. Also notable is a region of about 500 structural and functional domains of BRCA-1 and 2 are named above each of the schematic proteins. Representative proteins that interact with BRCA-1 and 2 are identified beneath them.
BRCA-1 and 2 couple signals of DNA damage to cellular responses, including damage repair and cell cycle checkpoint activities. Though much about this process is not yet understood, it is known that BRCA-1 is phosphorylated by any of several protein kinases, depending on the type of DNA damage. Both BRCA proteins interact physically with DNA repair proteins, and BRCA-1 also participates in transcription control of genes that encode DNA repair and cell cycle checkpoint control proteins.

amino acids that is more highly conserved between human and mouse than the coding sequence as a whole.

**BRCA-1 and 2 Are Caretakers of Genomic Stability**

Insights into the biological functions of BRCA-1 and 2 have come from analyses of cells derived from BRCA-mutant human breast tumors and from embryos of mice carrying targeted deletions of the BRCA genes. Invariably, BRCA-deficient cells exhibit gross chromosomal abnormalities, typified by breaks, aberrant mitotic exchanges, and aneuploidy.

These sorts of DNA damage arising from ongoing metabolic processes within the cell or caused by extrinsic agents, including radiation and certain chemicals, are a persistent threat to genome integrity. A response system has evolved to locate damaged DNA and effect its timely repair. BRCA-1 and 2 are parts of that system, cellular caretakers ensuring that the genetic integrity of a cell is not compromised by the unscheduled loss, duplication, or rearrangement of chromosomal DNA.

The DNA repair response involves the assembly of protein complexes capable of recognizing and eliminating damage-induced lesions, as well as the synthesis of proteins that arrest cell cycle progression while the damage is repaired. Disruption of the damage response system can lead to replication or segregation of damaged chromosomal DNA, and that in turn can permit a cell to escape normal restrictions on its growth, which is practically the definition of cancer.

Evidence to implicate BRCA-1 and 2 in the DNA damage response has come from the observation that cells deficient in either protein are hypersensitive to a variety of DNA-damaging agents. A more specific function was suggested by the finding that cells

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Discovery of the BRCA genes and early work on their protein products was described by Barbara Weber in the January/February 1996 issue of *Science & Medicine.*
deficient in either BRCA-1 or 2 exhibited overt defects in the repair of oxidative DNA damage. Further studies have documented direct interactions between BRCA-1 or BRCA-2 and individual protein components of the DNA repair machinery.

BRCA-1 has been linked to DNA repair through its interaction with a complex of three proteins, RAD-50/MRE-11/NBS-1, that operates in both nonhomologous and homologous recombinational repair of double-strand breaks. The three-protein complex has been proposed to resect DNA ends at the sites of double-strand breaks in order to reveal sequence homologies through which recombination can ensue. What BRCA-1 does in its association with this complex remains to be established.

BRCA-1 is also a resident component of a large multiprotein complex that includes mismatch repair proteins. These and other proteins are involved in replication or in repair of DNA damage that can occur at replication forks. The association of BRCA-1 with these proteins suggests that it participates in resolving aberrant DNA structures that appear during replication or when replication is stalled.

A role for BRCA-2 in DNA damage repair has been suggested by the discovery that it interacts with a recombinase called, in mammals, RAD-51. Mammalian RAD-51 is a homologue of the prokaryotic RecA and yeast Rad51p proteins, the latter a member of the RAD-52 epistasis group. In yeast, RAD-52 epistasis proteins are required for repair of DNA double-strand breaks as well as mitotic and meiotic recombination.

Eukaryotic RAD-51 proteins, like RecA, have intrinsic ATP-dependent DNA binding activity. RAD-51 and single-strand DNA form a nucleoprotein filament that invades and pairs with a homologous DNA duplex, catalyzing homologous DNA pairing and strand exchange. Mouse embryos lacking BRCA-2 exhibit radiation hypersensitivity defects like those seen in mouse embryos lacking RAD-51.

The interaction between BRCA-2 and RAD-51 involves the BRC repeats in BRCA-2. Peptides corresponding to individual BRC repeats can inhibit multimerization of RAD-51 and block nucleoprotein filament formation. Whether this inhibitory activity is a physiological role for BRCA-2 in regulating RAD-51 activity has not been established.

DNA double-strand breaks induced for example by ionizing radiation are repaired by two processes in which the BRCA proteins are involved. In a complex with RAD-50, MRE-11, NBS-1, and other proteins, BRCA-1 takes part in both homologous recombination and nonhomologous end-joining. BRCA-2 complexed with RAD-51 is active in strand exchange during homologous recombination.
In your original drawing, there is an up-arrow just to the right of the DNA below BRCA-1 (above cell cycle). What does this mean, and should it be reinstated?

With associated co-repressor CtIP, BRCA-1 ordinarily represses transcription of cell cycle checkpoint control genes. In response to certain kinds of DNA damage, BRCA-1 and CtIP are phosphorylated by ATM. Phosphorylated CtIP dissociates from BRCA-1, leading to relief of BRCA-1-mediated transcriptional repression and consequent induction of p21 and GADD-45. The protein products of those genes function in G₁-S and G₂-M cell cycle arrest, respectively.

Consistent with that possibility, though, is the observation that formation of RAD-51 protein complexes, normally induced by DNA damage, is diminished in cells either deficient in BRCA-2 or in which the interaction between BRCA-2 and RAD-51 is specifically disrupted. It therefore seems clear that BRCA-2 is necessary for the assembly of RAD-51 complexes.

Processes of DNA repair must be coordinated with regulation of cell cycle transit so that damage is repaired before chromosomal DNA is replicated or segregated. There is considerable evidence that BRCA-1 occupies a central place in activation of cell cycle checkpoints when DNA damage is detected.

First, BRCA-1-mutant cells exhibit defects in DNA damage-induced S and G₂-M cell cycle checkpoints. Second, after DNA damage, BRCA-1 is rapidly phosphorylated by cell cycle checkpoint kinases, suggesting that it functions downstream of DNA damage sensors that trigger cell cycle checkpoints. And third, BRCA-1 has been shown to regulate expression of cell cycle checkpoint control genes, including p21 and GADD-45, which function in G₁-S and G₂-M cell cycle checkpoints, respectively.

The role of BRCA-2 in cell cycle checkpoint control is much less clear. While examined, DNA damage-induced cell cycle checkpoints appear to be largely intact in cells lacking wild-type BRCA-2. Indirect evidence exists to link BRCA-2 to G₂-M control, but it is not entirely clear whether this is an indirect effect secondary to the role of BRCA-2 in DNA damage repair. The fact that specific disruption of the interaction between BRCA-2 and RAD-51 leads to loss of G₂-M checkpoint control suggests that this may be the case. Thus, BRCA-
SOMATIC MUTATION CHECKPOINT INACTIVATION

GERMLINE MUTATION

SOMATIC MUTATION

CHECKPOINT INACTIVATION

CANCER

2 inactivation could trigger existing checkpoints that monitor DNA structure, leading to delays in G2-M progression.

Collectively, the phenotypic characteristics of cells deficient in BRCA-1 or 2 suggest that these proteins are fundamental in the DNA damage response by participating in damage repair, cell cycle checkpoint control, or both.

Chromosomal instability arising from a defective DNA damage repair response has been proposed as the pathogenic basis for tumorigenesis accompanying BRCA deficiency. Paradoxically, chromosomal instability should lead to cell growth arrest or increased cell death, so the question is how BRCA-1 or 2 mutations might lead to the opposite effect.

One answer might lie in the observation that tumor cells deficient in BRCA-1 or 2 frequently harbor other inactivating mutations in cell cycle checkpoint control genes, including p53. Those mutations may circumvent the growth arrest that is normally induced by DNA damage and also inhibit p53-mediated apoptosis, permitting the survival of cells despite severe chromosomal damage.

On the other hand, inactivation of mitotic checkpoint genes could bypass mitotic arrest and permit aberrant chromosomes to segregate into progeny cells. This model is supported by experimental observations and suggests that the genetic instability arising in BRCA-1- or 2-deficient cells is pivotal in tumorigenesis, leading first to compensatory gene mutations that override damage-induced cell cycle arrest and apoptosis and subsequently to the accrual of functionally inactivating mutations of genetic loci involved in breast tumorigenesis.

**BRCA-1 and 2 Regulate Cell Growth and Differentiation**

Emerging evidence suggests important roles for BRCA-1 and 2 in the control of cell growth and differentiation. The clearest example is the observation that homzygous deletion of BRCA-1 in mice results in early embryonic lethality accompanied by developmental retardation and cellular proliferation defects.

This outcome can be explained in part by the involvement of BRCA-1 and 2 in DNA repair,
because targeted deletions in \( p53 \) or its downstream effector \( p21 \) can rescue embryos with homozygous Brca-1 and 2 deficiency. Thus, cumulative DNA damage arising in the absence of Brca-1 and 2 has been hypothesized to trigger \( p53 \)-mediated cell cycle arrest and apoptosis in the developing embryo, while inactivation of \( p53 \) leads to cell cycle checkpoint bypass and survival.

However, inactivation of \( p53 \) only partially rescues these embryos, which survive for only days longer in development. While the delayed embryonic lethality accompanying inactivation of \( p53 \) has been ascribed to the accumulation of gross chromosomal defects that are incompatible with life, the possibility also exists that Brca-1 and 2 are required for transit through a critical point later in embryonic development.

Another line of evidence has come from studies of transgenic mice carrying a Brca-1 allele that can be targeted for conditional inactivation specifically in the mammary glands of female mice. That inactivation elicits defects in ductal morphogenesis and also induces tumors that are associated with genetic instability, aneuploidy, and chromosomal rearrangements. In addition to independently supporting a role for BRCA-1 as a breast tumor suppressor, this mouse model has revealed that BRCA-1 is critical in mammary epithelial development. Conditional inactivation of BRCA-2 specifically in mammary gland has yet to be achieved, so the role of BRCA-2 in mammary gland formation remains to be established.

**BRCA-1 and 2 Regulate Transcription**

In parallel with the genetic studies, biochemical and molecular biological analyses have been carried out to determine how BRCA-1 and 2 execute their functions. The proteins have been linked to a variety of biological activities.

Involvement of BRCA-1 in transcriptional regulation was initially indicated by the identification near its carboxyl terminus of an acidic domain with an inherent transactivation function that is sensitive to cancer-predisposing mutations. This region interacts directly or indirectly with a variety of transcriptional co-activators, including the histone acetyltransferase p300 and hBRG-1, which is the catalytic subunit of a chromatin-remodeling complex called SW-1/SNF.

The same region, interestingly, also interacts with transcriptional co-repressors, including histone deacetylases and the CtIP/CtBP protein complex. BRCA-1 mutations found in familial breast cancer compromise the trans-activation function but also abolish the binding of BRCA-1 to co-repressors. These observations have prompted the speculation that BRCA-1 may function like a nuclear receptor, either activating or repressing transcription depending on associated co-factors.

Gene expression profiling methods have disclosed that ectopic overexpression of BRCA-1 can induce or repress many genes implicated in cell cycle control, cell cycle regulation, and DNA replication and repair. By virtue of this transcriptional regulatory activity, BRCA-1 could influence cellular responses downstream of DNA damage signals, including DNA repair and cell cycle checkpoint activation.

BRCA-1-mediated regulation of GADD-45 transcription illustrates how BRCA-1 might participate in cell cycle checkpoint control and also provides a model for how BRCA-1 can achieve gene-specific transcriptional regulation. GADD-45 is a tumor suppressor gene induced by DNA damage. Its encoded product functions in G2-M cell cycle checkpoint control.

Induction of GADD-45 transcription in response to ultraviolet radiation and radiomimetic agents has been shown to depend on BRCA-1, and evidence exists to
suggested that the same may be true for ionizing radiation. It is also known that BRCA-1 interacts with a co-repressor, CtIP, to repress transcription of GADD-45 and that this interaction is disrupted by DNA damage.

Neither BRCA-1 nor CtIP can bind DNA in a sequence-specific manner, however, so how these proteins are recruited to their target genes was an unresolved question. The answer was recently provided by identification of an intervening protein, named ZBRK-1, that binds to both BRCA-1 and a specific DNA sequence element present in a subset of BRCA-1's target genes, including GADD-45.

In this way, BRCA-1 can be physically tethered and functionally linked to specific regulatory loci. It is ZBRK-1 that actually represses transcription when it is bound to BRCA-1, so that BRCA-1 itself is a co-repressor. Potential ZBRK-1 binding sites have been identified in a large group of genes inducible by DNA damage, so the ZBRK-1/BRCA-1 complex may be a global regulator of DNA damage-responsive genes.

A model has been proposed whereby ZBRK-1, BRCA-1, and CtIP coordinately repress a functionally diverse group of DNA damage-response genes in the absence of genotoxic insult, and that phosphorylation induced by DNA damage disrupts the network of interactions among these proteins, de-repressing transcription.

It must be emphasized that derepression as an operative mechanism in transcriptional control of GADD-45 and other inducible genes in vivo is likely to be coordinated with other mechanisms of gene activation. BRCA-1 has been reported to interact functionally with a variety of sequence-specific DNA-binding transcriptional activators, including the tumor suppressor p53.

Model for sequence-specific transcription control by BRCA-1 through its dual role as a co-repressor and a co-activator: ZBRK-1 is a transcriptional repressor that recruits BRCA-1 to its specific DNA binding sites in target genes, one of which is in intron 3 of GADD-45. BRCA-1 may then (1) recruit CtIP and CBP to reorganize higher chromatin structure, (2) recruit histone deacetylase complexes to effect local gene silencing, or (3) interact with the basal transcription machinery. In response to an appropriate DNA damage signal, BRCA-1-mediated repression of GADD-45 transcription is relieved. That permits BRCA-1 to become a co-activator of, for example, p53, which also binds to intron 3 of GADD-45. BRCA-1 could mediate transcriptional activation by either (1) recruiting chromatin-modifying activities to facilitate transcription complex assembly at the promoter or (2) directly recruiting the RNA polymerase II holoenzyme to the promoter. In this model, damage-induced transcription of GADD-45 results from concerted derepression and activation.
In this regard, p53 appears to be an important link between BRCA-1 and transcriptional activation of DNA damage-inducible genes. It lies at the heart of a cell-signaling pathway that is triggered by genotoxic stresses, including DNA damage. Stress-induced p53-initiated cell cycle arrest or apoptosis ensures the timely repair or elimination of potentially deleterious genetic lesions.

Significantly, p53 and BRCA-1 appear to regulate transcription of an overlapping set of DNA damage-inducible target genes, including GADD-45. This observation initially implied a functional interaction between these two important tumor suppressors, a prediction that has since been borne out experimentally.

BRCA-1 and p53 have been demonstrated to interact physically and to synergize functionally to activate transcription through a p53-binding site in a GADD-45 intron. The ability of BRCA-1 to potentiate p53-dependent transcription without itself binding to DNA has led to the hypothesis that BRCA-1 functions as a p53-specific co-activator, possibly linking the biochemical activities of these two proteins to a common pathway of tumor suppression.

By being both a co-repressor and a co-activator of gene transcription, BRCA-1 appears to function as a link between parallel and perhaps synergistic pathways that lead to induction of DNA damage repair effectors. Before it can be understood how BRCA-1 integrates these dual functions, it will be necessary to decipher the mechanistic basis for its independent activation and repression.

In contrast to BRCA-1, the part that BRCA-2 plays in transcriptional regulation is far less certain. Some evidence implicates BRCA-2 in transcription control, including, again, an inherent trans-activation function within the gene that is sensitive to cancer-predisposing mutations and an association with established transcriptional co-factors and histone acetyltransferases. However, the biological significance of these findings has not been demonstrated.

Most if not all of the cellular pool of BRCA-1 resides in stable complexes with other proteins, so one possibility is that BRCA-1 is a molecular scaffold that facilitates assembly of multiprotein machines. Alternatively, the documented association of BRCA-1 with activities that modify chromatin could point to pleiotropic roles in DNA repair and gene transcription. BRCA-1 could variously promote or disrupt nucleosome-mediated condensation of DNA at gene promoters or DNA damage sites, thus precluding or facilitating access by transcription and repair factors, respectively.

Recent work has uncovered a ubiquitin ligase activity of BRCA-1, which raises the intriguing possibility that the protein's multiple functions could all derive from an ability to selectively mark proteins for destruction. Specifically, BRCA-1 interacts with another RING finger protein named BARD-1 through the respective RING domains of each protein. A heterodimeric complex formed by the isolated RING domains of the two proteins exhibits ubiquitin ligase activity in vitro.

Significantly, cancer-related missense mutations within the BRCA-1 RING finger abrogate this activity, suggesting that ubiquitin ligase activity may be important for the biological function of BRCA-1 in breast and ovarian tumor suppression. Presently, no physiological substrates of BRCA-1/BARD-1-targeted ubiquitination have been identified. But if BRCA-1 is involved in targeting proteins for ubiquitination, its participation in a wide range of cellular processes could be explained to some extent.

BARD-1 also interacts with a polyadenylation factor, CstF-50, which indirectly links BRCA-1 to RNA processing. Whether and how the ubiquitin ligase activity of...
BRCA-1 alone or in association with BARD-1 contributes to the functions of BRCA-1 is an important area for future investigation.

**Tumor Susceptibility Is Tissue-Specific**

DNA damage response pathways that converge on BRCA-1 and 2 are conserved across many cell types, so that BRCA-1 and 2 are likely to function widely in the maintenance of genomic integrity. Nonetheless, mutational inactivation of these genes leads principally to cancer of the breast and ovary. Why?

As reproductive organs, breast and ovary rely on hormones for growth and differentiation. At least two hypotheses invoking the action of hormones have been proposed to explain the tissue-restricted tumor suppressor functions of BRCA-1 and 2. According to one model, mutational inactivation of the BRCA genes renders breast susceptible to the tissue-specific effects of estrogen-induced DNA damage. A major oxidative metabolite of estrogen, 4-hydroxyestradiol, is genotoxic.

The suggestion is that inactivating mutations in *BRCA-1* or 2 could compromise the response of breast epithelial cells in particular to estrogen-induced DNA damage. Inefficient or error-prone DNA repair could then lead to genomic instability and a concomitant accrual of functionally inactivating mutations within other genes involved in breast tumorigenesis. Put another way, *BRCA-1* and 2 mutations might enhance the probability of tumor formation arising from estrogen-induced DNA damage.

A second model proposes that BRCA-1 and 2 modulate hormone signaling pathways that induce cell proliferation. BRCA-1 has been shown to repress the transcriptional activity of the estrogen receptor (ER-α), so mutational inactivation of BRCA-1 could promote epithelial cell proliferation by altering expression of hormone-responsive genes.

The two models are not mutually exclusive and could suggest a combinatorial path to breast cancer, with BRCA-1-mediated control operating at two distinct steps in tumorigenesis, initiation and progression.

ubiquitin is a marker that tags other proteins for destruction. The sequence of events is shown here schematically.

A ubiquitin-activating enzyme E1 (UBA) is charged with ubiquitin, which is then transferred to a ubiquitin-conjugating enzyme E2 (UCB). A ubiquitin ligase E3 presumably functions as a platform for recruitment of both the E2 enzyme and a substrate protein, which is polyubiquitinated and thereby targeted for destruction.

A heterodimer formed by isolated RING domains of BRCA-1 and BARD-1 can function as an E3 ubiquitin ligase in vitro. Remaining surfaces on the two proteins could be involved in substrate recruitment in vivo. The structure of the heterodimer formed by the RING domains has been described.
BRCA-1 is a barrier to transcription of genes that are targets of the estrogen receptor (ER), preventing cell proliferation by repressing unliganded ER bound to the estrogen response element (ERE). BRCA-1-mediated ER suppression additionally involves one or more co-repressors, minimally including a histone deacetylase activity.

In cells deficient in BRCA-1, ERE-bound ER is free to promote transcription of its target genes and cell proliferation independent of estrogen. Such transcription derives from recruitment of co-activators.

Models explaining how BRCA-1 acts through modulation of estrogen receptor function must account for the clinical observation that a significant proportion of BRCA-1-associated breast cancers are negative for ER-α expression. A definitive understanding of this phenomenon is precluded by the fact that it simply is not known how “ER-negative” tumors arise.

It has recently been shown that within the terminal ductal lobular unit, where breast cancers are believed to originate, there are at least three distinct epithelial cell populations: ER-α-positive cells that do not proliferate, ER-α-negative cells that do proliferate, and a small number of ER-α-positive cells that can proliferate as well.

Again, there are two principal models for the genesis of ER-α-negative epithelial-derived tumors, both of which are compatible with a role for BRCA-1 in the control of epithelial cell proliferation through functional interaction with ER-α.

In one model, ER-α-negative breast cancers arise from the loss of ER-α expression during the clinical evolution of cancers that were originally ER-α-positive. In this case, it is possible that the loss of ER-α expression is a relatively late event in breast tumor progression, one that may occur after any proliferative advantages conferred upon transformation-initiated cells by homozygous BRCA-1 mutation have ensued.

Alternatively, it has been proposed that ER-α-negative and ER-α-positive tumors are distinct entities that reflect the receptor status of their clonal origins. Recent data suggest a model in which proliferation of ER-α-negative cells is controlled by paracrine growth factors released from ER-α-positive cells in an estrogen-dependent manner. Here, mutational inactivation of BRCA-1 could promote growth factor-mediated proliferation of ER-α-negative tumors.

Finally, discovery of a second estrogen receptor subtype, ER-β, raises the possibility that this receptor mediates the proliferative response to estrogen in cells that are negative for ER-α expression. ER-β is expressed during the immortalization and transformation of ER-α-negative human breast epithelial cells in vitro.

The functional role of ER-β-mediated estrogen signaling path-
DNA DAMAGE

Role of the BRCA proteins in breast cancer. In normal breast epithelial cells, BRCA-1 and 2 ensure efficient DNA repair, thereby preserving genomic integrity in the face of genotoxic insult, including the action of estrogen metabolites. In addition, BRCA-1 restricts estrogen-independent expression of estrogen-responsive genes by directly inhibiting the unliganded estrogen receptor, thus rendering cells dependent on estrogen for growth. BRCA-deficient breast epithelial cells can develop unstable genomes through inefficient repair of damaged DNA and can become independent of estrogen for growth.

How might the knowledge now at hand concerning the biological functions of BRCA-1 and 2 be exploited to clinical advantage? For women genetically predisposed to BRCA-1 and 2 mutations, restricted exposure to direct or indirect extrinsic sources of DNA damage might be warranted.

In reality, knowledge about BRCA-1 and 2 function might find its most useful applications in the treatment of the 90% of sporadic breast cancers for which no genetic linkage with an identifiable susceptibility locus can be found. In these sporadic cancers, perturbation of other pathways are likely involved in tumorigenesis. Nonetheless, as caretaker genes, BRCA-1 and 2 represent prime targets for therapeutic intervention. For example, targeted inactivation of BRCA-1 and 2-specific DNA damage response pathways could render tumor cells particularly sensitive to the genotoxic effects of radiation or chemotherapeutic agents, offering the potential for improved combination therapies.

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BRCA1 and Estrogen Signaling in Breast Cancer

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Abstract: Since the gene encoding BRCA1 was first cloned in 1994, numerous studies have revealed its role in breast cancer susceptibility. The correlation between genetic variation in BRCA1 and breast cancer risk has been established in several studies. BRCA1 has been associated with increased risk of breast cancer in patients with certain BRCA1 variants. Furthermore, BRCA1 expression levels have been linked to breast cancer risk, with higher expression associated with a higher risk of breast cancer. This association is thought to be mediated by the estrogen receptor (ER) signaling pathway. The ER signaling pathway plays a crucial role in breast cancer biology, and genetic variation in BRCA1 may influence this pathway.

INTRODUCTION

The National Cancer Institute states that 1 in 8 women will develop breast cancer during her lifetime (1). The high incidence of this disease has driven research into identifying risk factors and developing interventions to prevent or delay the onset of breast cancer. BRCA1 is a key gene in the development of breast cancer, and recent studies have shown that BRCA1 polymorphisms are associated with increased breast cancer risk (2). The ER signaling pathway is a critical component of breast cancer biology, and genetic variation in BRCA1 has been linked to altered ER signaling.

Recent studies have suggested that BRCA1 and ER signaling pathways are closely intertwined. Genetic variation in BRCA1 has been shown to affect the expression of estrogen-responsive genes, suggesting that BRCA1 may influence ER signaling. This interaction is thought to play a role in the development of breast cancer.

METHODS

The study was conducted using a case-control design. Breast cancer cases and controls were recruited from a large population-based cohort. DNA was extracted from peripheral blood samples and genotyped for SNPs in the BRCA1 and ER signaling pathway genes. Association analyses were conducted using logistic regression models.

RESULTS

Significant associations were observed between SNPs in BRCA1 and ER signaling pathway genes and breast cancer risk. Specifically, polymorphisms in BRCA1 were found to be associated with increased risk of breast cancer in women with certain ER expression levels. This finding supports the hypothesis that BRCA1 expression may be influenced by the ER signaling pathway.

CONCLUSIONS

Our study provides evidence for a genetic interaction between BRCA1 and ER signaling pathway genes and breast cancer risk. These findings highlight the importance of considering genetic variants in both pathways when assessing breast cancer risk. Further research is needed to understand the mechanisms underlying this interaction and to develop targeted interventions for high-risk populations.