BRCA1-induced large-scale chromatin unfolding and allele-specific effects of cancer-predisposing mutations

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The breast cancer susceptibility gene BRCA1 encodes a protein that has been implicated in multiple nuclear functions, including transcription and DNA repair. The multifunctional nature of BRCA1 has raised the possibility that the polypeptide may regulate various nuclear processes via a common underlying mechanism such as chromatin remodeling. However, to date, no direct evidence exists in mammalian cells for BRCA1-mediated changes in either local or large-scale chromatin structure. Here we show that targeting BRCA1 to an amplified, lac operator–containing chromosome region in the mammalian genome results in large-scale chromatin decondensation. This unfolding activity is independently conferred by three subdomains within the transactivation domain of BRCA1, namely activation domain 1, and the two BRCA1 COOH terminus (BRCT) repeats. In addition, we demonstrate a similar chromatin unfolding activity associated with the transactivation domains of E2F1 and tumor suppressor p53. However, unlike E2F1 and p53, BRCT-mediated chromatin unfolding is not accompanied by histone hyperacetylation. Cancer-predisposing mutations of BRCA1 display an allele-specific effect on chromatin unfolding: 5′ mutations that result in gross truncation of the protein abolish the chromatin unfolding activity, whereas those in the 3′ region of the gene markedly enhance this activity. A novel cofactor of BRCA1 (COBRA1) is recruited to the chromosome site by the first BRCT repeat of BRCA1, and is itself sufficient to induce chromatin unfolding. BRCA1 mutations that enhance chromatin unfolding also increase its affinity for, and recruitment of, COBRA1. These results indicate that reorganization of higher levels of chromatin structure is an important regulated step in BRCA1-mediated nuclear functions.

Introduction

Germ line mutations in breast cancer susceptibility gene 1 (BRCA1)* confer elevated risks in the development of familial breast and ovarian cancers (Rahman and Stratton, 1998). BRCA1 encodes a 1,863–amino acid protein with a highly conserved ring finger motif (RING) at the NH2 terminus, and two BRCA1 COOH terminus (BRCT) repeats at the extreme COOH terminus. Whereas most disease-associated mutations of BRCA1 are predicted to result in gross truncation of the protein, 5–10% of the cancer-predisposing mutations cause single amino acid substitutions, many of which are located in the RING domain or BRCT repeats.

Intense research in the past several years has implicated BRCA1 in the regulation of multiple nuclear processes, including DNA repair and transcription (Zhang et al., 1998b; Scully and Livingston, 2000). For example, BRCA1-deficient mouse and human cells are hypersensitive to ionizing radiation due to defects in transcription-coupled repair of oxidative DNA damage, as well as double-strand break-induced homologous recombination (Gowen et al., 1998; Abbott et al., 1999; Moynahan et al., 1999; Scully et al., 1999; Xu et al., 1999). In addition, BRCA1 associates with several repair and recombination proteins such as RAD51 (Scully et al., 1997b), RAD50/MRE11/NBS1 (Zhong et al., 1999; Wang et al., 2000), and MSH2/MSH6 (Wang et al., 2000). BRCA1 also interacts with and is phosphorylated by protein kinases that are key players in the damage checkpoint control, including ATM, ATR, and CHK2 (Cortez et al., 1999; Lee et al., 2000; Tibbetts et al., 2000). Lastly, it has been shown recently that BRCA1 preferentially binds to branched DNA structures (Paul et al., 2001).

In addition to its potential role in DNA repair, BRCA1 has also been implicated in regulation of transcription...
When tethered to a transcriptional promoter via a heterologous DNA binding domain, the COOH-terminal 304–aminoc acid (aa) region including the BRCT repeats (Fig. 1 B, AD2, amino acids 1560–1863) can act as a transactivation domain (Chapman and Verma, 1996; Monteiro et al., 1996). More recent work has revealed a second transactivation domain of BRCA1 that resides upstream of the BRCT repeats (Hu et al., 2000) (Fig. 1 B, AD1, aa 1293–1559). The two activation domains (ADs), AD1 and AD2, can cooperatively activate transcription in many cell lines tested (Hu et al., 2000). Consistent with its potential role in transcriptional regulation, the BRCA1 polypeptide is associated with the RNA polymerase II holoenzyme via RNA helicase A (Scully et al., 1997; Ouchi et al., 1998, 2000; Zhang et al., 1998a; Fan et al., 1999; Houvras et al., 2000; Zheng et al., 2000).

The multifunctional nature of BRCA1 has raised the possibility that the protein may employ a common mechanism, such as chromatin remodeling, to regulate various chromosomal events. Indeed, the COOH-terminal region of BRCA1 (AD2), which is required for BRCA1 functions in both DNA repair and transcription (Monteiro, 2000; Scully and Livingston, 2000), can induce changes in nucleosome structure when tethered to chromosomal DNA in Saccharomyces cerevisiae (Hu et al., 1999). Furthermore, BRCA1 is associated with histone modifying enzymes (p300 and HDAC) (Neish et al., 1998; Tarden and Brody, 1999; Pao et al., 2000) and an ATP-dependent chromatin remodeling machine (hSNF/SWI) (Bochar et al., 2000). The fact that many cancer-predisposing mutations reduce BRCA1's affinity for these chromatin-modifying proteins suggests that chromatin remodeling may be an important aspect of BRCA1-mediated tumor suppression. However, currently there is no direct evidence in mammalian cells for BRCA1-mediated changes in chromatin structure. This is in part due to the lack of convenient assays for directly monitoring chromatin remodeling at different levels of chromatin structure in mammalian cells.

A lac repressor–based system has allowed direct visualization of large-scale chromatin dynamics in mammalian cells (Belmont, 2001). In this system, multiple copies of the lac operator were engineered into the genome of CHO cells, and together with the surrounding genomic sequences, were amplified to produce a 90-Mb heterochromatic region. By fusing lac repressor with the acidic AD (AAD) of the strong viral transcription factor VP16 and tethering the fusion protein to the heterochromatic chromosome region, this system was used to demonstrate AAD-induced large-scale chromatin decondensation (Tumbar et al., 1999). This large-scale chromatin uncoiling occurred even when RNA pol II–dependent transcription was blocked, suggesting that it was induced through transacting factors recruited by the VP16 AAD, rather than the result of transcription per se. Conceptually, the transacting factors producing this higher order chromatin decondensation could be one of the known chromatin-modifying complexes that modify local nucleosome structure (Peterson and Logie, 2000). Alternatively, AAD-induced chromatin unfolding could involve novel factors acting primarily at the higher levels of chromatin organization.

Although artificial, this lac repressor–tethering system provides a very quick, and therefore powerful, assay to test the possible role of specific proteins in chromatin remodeling and to dissect the protein domains required for the observed large-scale chromatin decondensation. Using this lac repressor–tethering assay, we demonstrate here that BRCA1 induces large-scale chromatin decondensation. We also identify three small subdomains within the transactivation domain of BRCA1 that are capable of independently conferring chromatin unfolding. In addition, cancer-predisposing mutations of BRCA1 display allele-specific effects on the chromatin unfolding activity. Finally, we isolate a novel cofactor of BRCA1 (COBRA1) that binds to one of the chromatin-unfolding domains of BRCA1, and by itself induces large-scale chromatin decondensation. Our results suggest that BRCA1-mediated decondensation of higher levels of chromatin structure may represent a new physiological regulatory pathway related to BRCA1 function. The approach used in the current study also provides a new methodology for identifying novel BRCA1-interacting proteins involved in this regulatory pathway.

**Results**

**BRCA1-mediated large-scale chromatin decondensation in mammalian cells**

To assess the impact of BRCA1 on large-scale chromatin structure in mammalian cells, we made use of a CHO cell line, A03_1, in which multiple copies of the lac operator were engineered to produce a 90-Mb heterochromatic region of the genome (Robinett et al., 1996; Li et al., 1998; Tumbar et al., 1999). The molecular organization of this region consists of ~400-kb repeats of the 14-kb vector transgene that contains the lac operator repeat and dihydrofolate reductase selectable marker. The repeats are separated on average by ~1,000 kb of unknown coamplified genomic DNA. Because other cell clones derived from the same selection procedure contain more open, gene-amplified chromosome regions with comparable or greater content of the vector DNA, the heterochromatic appearance of the A03_1 chromosome region is assumed to be due to properties of the coamplified genomic DNA. In vivo binding of lac repressor or its GFP derivatives to this chromosomal site allows direct visualization of large-scale chromatin dynamics without altering the original chromosome structure.

Consistent with previous findings, lac repressor–expressing cells stained with the corresponding antibody exhibited a compact nuclear dot (Fig. 1 A, a). In contrast, expression of lac repressor fused with the full-length BRCA1 induced an irregularly shaped subnuclear structure in 14% of transfected cells (Fig. 1 A, b). Such a staining pattern was not present in any of the cells expressing lac repressor alone. These results suggest that BRCA1, or a BRCA1-associated protein, can induce large-scale chromatin restructuring. The magnitude of this opening was lower than observed for the VP16 AAD, and was present in a lower percentage of cells (14 vs. 60% for VP16 AAD) (Tumbar et al., 1999 and see Fig. 2). The lack of a response in 100% of cells, even for the
VP16 AAD, is not yet understood. It may represent a combination of several factors, including cell cycle–dependent expression as well as the nature of the qualitative assay employed. Transgene arrays have been shown to display coordinated gene silencing effects that are accompanied by cooperative changes in chromatin structure across the entire array (Pikaart et al., 1998). These changes in gene expression and chromatin structure show a variegating phenotype that is clonally inherited. Therefore, it is possible that the large-scale chromatin decondensation induced by a transcriptional activator in the lac system may also display cooperative and variegating responses.

Deletion analysis showed that chromatin-unfolding activity was conferred by the last 570 aa of BRCA1 (Fig. 1 B, aa 1293–1863). This region of BRCA1, previously designated AD (Hu et al., 1999), consists of two subdomains that act synergistically to stimulate transcription (Fig. 1 B, AD1, aa 1293–1559, and AD2, aa 1560–1863). As illustrated in Fig. 1 B, AD2 contains the two BRCT repeats, BRCT1 and BRCT2. Further domain mapping indicated that AD1, BRCT1, and BRCT2 could independently induce large-scale chromatin unfolding (Fig. 1 A, c, e, and f, and B). It is of note that AD1 often leads to a ball-shaped structure with smooth edges, whereas BRCT1 and BRCT2 tend to give rise to more extended, fiber-like structures with irregular shapes (Fig. 1 A, compare c with e and f). The degree of unfolding by BRCT1 and BRCT2 approached that observed with VP16, with >60% of cells showing this response, whereas the AD1 subdomain showed intermediate unfolding. Interestingly, both the magnitude of this unfolding and the percentage of cells showing unfolding using these subdomains was significantly higher than observed using the full-length BRCA1 fusion protein (Fig. 1 B). Furthermore, AD2, which includes both the BRCT1 and BRCT2 repeats, failed to cause obvious decondensation of high-order chromatin structure (Fig. 1 A, compare d with e and f). As explained below, we interpret this as an indication of a negatively regulated chromatin unfolding activity associated with the full-length BRCA1 and AD2 region.

Further dissection of the BRCT1 domain shows that the 50-aa COOH-terminal half of BRCT1 is sufficient for inducing maximal chromatin unfolding (Fig. 1 B, BRCT1C). In contrast, the NH2-terminal half of BRCT1 (BRCT1N) with a comparable size to BRCT1C, fails to mediate any chromatin decondensation. Furthermore, none of the BRCA1 fragments upstream of AD displayed any activity in chromatin unfolding (Fig. 1 B, 1–324, 260–554, and 554–1293), although they were expressed at similar levels as the chromatin-unfolding domains (unpublished data and see Fig. 5 B). Previous studies have shown that these regions upstream of AD are responsible for BRCA1 interactions with various proteins or protein complexes. For example, the NH2-terminal region
of BRCA1 binds BARD1 (Wu et al., 1996), whereas the central region of the protein mediates BRCA1 interactions with hSNF/SWI (Bochar et al., 2000), RAD50/MRE11/NBS1 (Zhong et al., 1999), and RAD51 (Scully et al., 1997b). The inability of these regions to induce large-scale chromatin decondensation argues that chromatin unfolding is not simply due to recruitment of any large proteins or protein complexes to the lac binding sites. Rather, the chromatin unfolding activity is conferred by three specific subdomains in the transactivation domain of BRCA1, suggesting that chromatin decondensation is related to BRCA1 functions in transcriptional regulation and DNA repair.

**Distinction between BRCT and other well-characterized transactivation domains in large-scale chromatin unfolding**

A previous study has shown that VP16-induced chromatin unfolding is accompanied by recruitment of histone acetyltransferases and local histone hyperacetylation, a property frequently observed for transcriptionally active or competent chromatin (Tumbar et al., 1999) (Fig. 2 A, m–o). Here we extended the previous work by examining the transactivation domains of two cellular transcription factors, E2F1 and p53. Like lac-VP16, lac-E2F1 and lac-p53 also induced significant chromatin unfolding in 60% and 45% of transfected cells, respectively (Fig. 2 A, g and j). Furthermore, the lac-E2F1– and lac-p53–unfolded chromatin regions were enriched with hyperacetylated histone H3 and H4 (Fig. 2 A, g–i and j–l, and unpublished data). Thus, all three well-characterized transactivation domains (VP16, E2F1, and p53) can simultaneously induce large-scale chromatin unfolding and histone hyperacetylation. However, it remains unknown whether the observed histone hyperacetylation is causally related to chromatin unfolding.

The extent of chromatin decondensation induced by a single BRCT repeat is comparable to that exhibited by these potent transcriptional ADs (Fig. 2 A, compare a and d with g, j, and m). However, no obvious histone H3 or H4 hyperacetylation was detected in the BRCT1- or BRCT2-unfolded chromatin regions (Fig. 2 A, a–c and d–f), suggesting that BRCT-mediated chromatin unfolding is a separable event from histone acetylation. Although both BRCT repeats are required for AD2-mediated transcriptional activation, a single repeat does not serve as a strong AD (Chapman and Verma, 1996; Monteiro et al., unpublished data).
Therefore, chromatin unfolding by BRCT may be a necessary, but not sufficient step, in transcriptional activation.

In addition to acetylation, histones are subject to other posttranslational modifications under various physiological conditions. Of particular interest, phosphorylation of H2AX, a histone H2A variant, at serine 139 (γ-H2AX) is rapidly stimulated following ionizing radiation (Rogakou et al., 1999). Before irradiation, a subset of γ-H2AX nuclear foci colocalize with BRCA1 foci (Paull et al., 2000). After DNA damage, the number of both γ-H2AX and BRCA1 nuclear foci increases significantly; furthermore, the majority of BRCA1 foci overlapped γ-H2AX foci (Paull et al., 2000). These observations most likely reflect localized recruitment of the putative H2AX kinase and phosphorylation of H2AX-containing nucleosomes that are already present at these sites, rather than recruitment of new H2AX protein.

Using an antibody that specifically recognizes the phosphorylated form of H2AX (γH2AX), we detected colocalization of the endogenous γ-H2AX with the full-length BRCA1 fusion protein in a sub-population (15%) of the lac–BRCA1-transfected cells (Fig. 2 B, a–c). In contrast, lac-VP16 did not display any colocalization with γ-H2AX (Fig. 2 B, d–f), nor did lac-BRCT1 or lac-BRCT2 (unpublished data). It is not clear whether phosphorylation of H2AX is causally linked to BRCA1-mediated chromatin unfolding, as γ-H2AX colocalization is also observed in lac–BRCA1-expressing cells that do not display chromatin decondensation. Consistent with previous reports (Rogakou et al., 1999; Paull et al., 2000), ionizing radiation significantly increased the number and overall intensity of γ-H2AX foci (unpublished data). However, the strong γ-H2AX signal over the entire nucleus made it difficult to examine the effect of DNA damage on the colocalization between γ-H2AX and lac-BRCA1 at the lac binding sites.

Work by Paull et al. (2000) has shown that H2AX at the damaged sites is rapidly phosphorylated after ionizing radiation, which is followed later by colocalization of BRCA1 and other repair proteins (Paull et al., 2000). It is possible that the putative kinase(s) responsible for H2AX phosphorylation directly binds to the full-length BRCA1. In such an
event, tethering lac–BRCA1 may simply bring the kinase(s) to the tandem array of the lac binding sites, thus causing hyperphosphorylation of H2AX present in the surrounding chromosomal region. Whereas the functional significance of H2AX phosphorylation in chromatin unfolding remains to be explored, our finding is consistent with the previous suggestion of a physical link between γ-H2AX and BRCA1.

Allele-specific effects of cancer-predisposing mutations of BRCA1 on chromatin unfolding

To determine the effect of cancer-associated mutations on the BRCA1-dependent chromatin unfolding, we introduced a series of common cancer-predisposing mutations into either full-length BRCA1 or AD2. Based on their behaviors in the chromatin-unfolding assay, mutations were classified into three phenotypic categories. The first includes nonsense mutations resulting in truncation of the entire COOH terminus (Fig. 3 B, a). According to previous studies, BRCA1 mutants that lack the COOH terminus of the protein are defective in stimulating transcription and DNA repair (Somusundaram et al., 1997; Abbott et al., 1999; Scully et al., 1999; Jin et al., 2000). As shown in group a of Fig. 3 B, these COOH-terminal truncation mutants also failed to induce chromatin unfolding. The second group of mutants include missense mutations that are located upstream of AD2 (group b, i.e., C61G, S1040N, and R1347G). None of the mutants in this group significantly affects BRCA1-mediated chromatin unfolding.

Contrary to the behaviors of first two groups, mutations in group c markedly enhanced the ability of lac-BRCA1 to induce chromatin unfolding (Fig. 3 B). For example, A1708E, M1775R, and Y1853X led to a pronounced enlargement of the unfolded chromatin structure (Fig. 3 A, compare b with c and d, and e with f–h). The same mutations also significantly increased the percentage of transfected cells that showed chromatin unfolding (Fig. 3 B). For instance, 79% of the cells that expressed the M1775R mutant displayed significant chromatin unfolding, compared with 14% for the wild-type full-length protein. This is an even higher percentage than that previously observed for the VP16 activator (Tumbar et al., 1999).

Interestingly, all mutations in group c result in single aa substitutions or small deletions within the AD2 region. Many of the mutations in this group have been shown previously to abolish AD2 interactions with other transcription-related proteins, including the RNA pol II holoenzyme (Scully et al., 1997a; Neish et al., 1998; Yarden and Brody, 1999). As discussed below, by retaining the chromatin unfolding activity of BRCA1 but blocking its role in other steps of transcriptional activation, these mutations in group c may lead to accumulation of the highly decondensed chromatin structure as observed in the unfolding assay.

Identification of a novel BRCA1-interacting protein

Application of the chromatin unfolding assay allowed us to identify a large-scale chromatin unfolding activity associated with BRCA1, and to narrow down the chromatin-unfolding region of BRCA1 to small subdomains in the COOH terminus of the protein. To identify cofactors recruited by the BRCT repeats to mediate chromatin unfolding, we used BRCT1 as the bait in a yeast two-hybrid screen. One candidate gene, cofactor COBRA1, was isolated from a human ovary cDNA library. It encodes a novel 580-aa protein rich in leucine residues (17%) (Fig. 4). COBRA1 also contains three repeats of the LXXLL motif, often present in many transcription coactivators and responsible for mediating their ligand-dependent interactions with steroid hormone receptors (Heery et al., 1997). Database searches revealed COBRA1-related hypothetical proteins in mice and flies that share 96 and 51% aa identity with the human protein, respectively (Fig. 4).

To confirm the interaction between BRCA1 and COBRA1, a lysate of human HEK293T cells that ectopically expressed FLAG-tagged COBRA1 was immunoprecipitated with an anti-FLAG antibody, followed by immunoblotting with an anti-BRCA1 antibody. As shown in Fig. 5 A, the endogenous human BRCA1 was coprecipitated in a FLAG-COBRA1–dependent manner (lanes 1 and 2). As a control, addition of an excess of FLAG peptide to the immunoprecipitation reaction abolished the BRCA1 signal in the immunoprecipitate (Fig. 5 A, lane 3).

To further assess the binding specificity of COBRA1 to the BRCT1 region of BRCA1, we cotransfected HEK293T cells with FLAG-COBRA1 and lac repressor fused with various fragments of BRCA1. The cell lysates were then immunoprecipitated with the anti-FLAG antibody and subsequently immunoblotted with the anti-lac antibody. As shown in Fig. 5 B, lac-BRCT1 was capable of binding to the FLAG-COBRA1 (lane 2). Consistent with their activity in

Figure 4. Sequence alignment of human COBRA1 and its homologues from mice and flies. The conserved aa residues are highlighted in black, and the similar residues in gray. The locations of the LXXLL motif are indicated by asterisks.
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Chromatin unfolding, the COOH-, but not the NH$_2$-terminal half of BRCT1 (Fig. 1 B), interacted with COBRA1 (lanes 3 and 4). None of the BRCA1 fragments upstream of the BRCT repeat, including AD1, displayed any significant affinity for COBRA1 (lanes 5–8). Taken together, our data show that the COBRA1 binding correlates with the BRCT1-mediated large-scale chromatin unfolding.

As shown in Fig. 3, cancer-predisposing mutations in the 3’ region of BRCA1 caused significant enhancement of the chromatin unfolding activity. Intriguingly, the same mutations (A1708E, M1775R, and Y1853X) also increased the affinity for COBRA1 in the coimmunoprecipitation assay (Fig. 5 B, compare lane 9 with lanes 10–12). A similar result was also observed in an in vitro glutathione S-transferase (GST) pull-down assay (Fig. 5 C). In this case, $^{35}$S-labeled, in vitro–translated COBRA1 was pulled down by both GST–AD and GST–AD2, but not by GST–AD1 (Fig. 5 C, bottom panel, lanes 3–5). Furthermore, COBRA1 displayed a higher affinity for the mutant (M1775R) than the wild-type GST–AD2 fusion (Fig. 5 C, bottom panel, lanes 5 and 6). As a control, we also used $^{35}$S-labeled CtIP, a transcriptional corepressor that binds to the COOH terminus of BRCA1 (Wong et al., 1998; Yu et al., 1998; Li et al., 1999). Consistent with previous findings, CtIP binds specifically to AD2 but, unlike COBRA1, its association with AD2 is abolished by the M1775R mutation (Fig. 5 C, top panel, lanes 5 and 6). Thus, the same cancer-predisposing mutations exert opposite effects on BRCA1 binding to two different partners.

Involvement of COBRA1 in BRCT1-mediated chromatin unfolding

To explore the role of COBRA1 in the BRCT1-mediated chromatin unfolding, we cotransfected FLAG-COBRA1 with various lac–BRCA1 fusion constructs into AO3_1 cells. As detected by confocal immunofluorescent microscopy, FLAG-COBRA1 and lac–BRCT1 co-localized in 96% of the cells that expressed both proteins (Fig. 6 A, a–c). In contrast, we did not detect any enrichment of the FLAG-COBRA1 signal at either the BRCT2- or AD1-unfolded chromatin regions (Fig. 6 A, BRCT2, d–f, and AD1, g–i). Thus, whereas all three subdomains are capable of inducing large-scale chromatin unfolding, they appear to recruit distinct cofactors to mediate this process.

Wild-type AD2, which failed to induce chromatin unfolding (Fig. 1), did not display any obvious colocalization with COBRA1 (Fig. 6 A, panels j–l). However, two 3’ cancer-predisposing mutations in the same context led to pronounced recruitment of COBRA1 to the unfolded chromatin regions (Fig. 6 A, BRCT2, d–f, and AD1, g–i). Thus, whereas all three subdomains are capable of inducing large-scale chromatin unfolding, they appear to recruit distinct cofactors to mediate this process.

To directly assess the impact of COBRA1 on large-scale chromatin structure, we used lac repressor to target COBRA1 to the lac binding sites in AO3_1 cells. As shown in Fig. 6 B, 61% of the cells that expressed lac–COBRA1 showed a comparable extent of chromatin unfolding as did lac–BRCT1.
The findings strongly implicate COBRA1 in BRCT1-mediated chromatin restructuring.

Discussion

Eukaryotic genomes are packaged through multiple steps into higher levels of chromatin structure. It is now well established that remodeling of local chromatin structure is a key step common to the initiation of multiple chromosomal functional events, including transcription, DNA replication, repair, and recombination (Elgin and Workman, 2000; Fyodorov and Kadonaga, 2001). Whereas intense research in the past decade has provided a wealth of information regarding the biochemical basis for chromatin remodeling at the nucleosome level, much less is known about reorganization of higher levels of chromatin structure. It remains unclear whether the known modifications of nucleosome organization are sufficient for changes in large-scale chromatin organization, or whether novel mechanisms acting at higher levels of chromatin structure are responsible for changes in large-scale chromatin organization.

A major difficulty in distinguishing these two possibilities is that most assays for identifying transcriptional activators or coactivators have used transcriptional activity as a final readout. Direct assays for changes in higher order chromatin structure have not been used previously. Because BRCA1 had been functionally implicated in a range of nuclear processes, it was reasonable to postulate that the regulation of these multiple nuclear events might occur through a general chromatin remodeling activity of BRCA1. The lac repressor–tethering system, while artificial in many aspects, provided an excellent assay to pursue this research direction.

Our findings in this current study strongly suggest that BRCA1 recruits COBRA1, a novel factor, to the lac operator–containing chromatin region. Within the constraints of the lac repressor–tethering assay, BRCA1-dependent unfolding of higher levels of chromatin structure appears to be at least partially mediated through recruitment of COBRA1. Notably, BRCA1-mediated chromatin decondensation is distinct from transcriptional activation per se and histone hyperacetylation. It is unclear how unique the histone acetylation–independent chromatin unfolding is. Although the chromatin unfolding produced by VP16, E2F1, and p53 is accompanied by histone hyperacetylation, no causal relationship between histone acetyltransferases recruitment and chromatin unfolding has been demonstrated. Moreover, preliminary data suggests that large-scale decondensation produced by estrogen receptor does not correlate with histone hyperacetylation (A. Nye and A. Belmont, personal communication).
Whereas the lac-based chromatin-unfolding assay provides a new tool for visualizing chromatin dynamics and in vivo protein–protein interactions in mammalian cells, it is important to point out that the molecular and biochemical basis for BRCA1-mediated chromatin decondensation is yet to be understood. Furthermore, utilization of a long tandem array of lac binding sites may raise the concern that the observed chromatin unfolding could simply be due to steric effects of the proteins/protein complexes that are brought to the lac binding sites. However, we believe this possibility is unlikely because our work does not indicate an obvious correlation between the potency of chromatin unfolding and the size or charge of the tethered protein fragments. For example, the minimal chromatin-unfolding domain defined in our study is only 50 aa long (BRCA1C). In contrast, several other BRCA1 fragments that range in size from 324 to 740 aa do not display any chromatin-unfolding activity (Fig. 1B). In addition, BRCT1 and BRCT2 have a net charge of +5 and −6, respectively, yet both demonstrate strong chromatin-unfolding activity. On the other hand, BRCTN carries more positive charges (+5) than BRCT1C (+1), but only the latter can induce chromatin decondensation. Finally, in previous work using either lac repressor tetramer, or lac repressor fused to several other protein domains up to ∼350 aa in size (i.e., GFP), no effect on large-scale chromatin structure has been observed (Robinnett et al., 1996, and A. Belmont, personal communication).

In our minds, a more serious caveats concerning the lac repressor–tethering system is the question of whether the observed effects produced by BRCA1 and other proteins on large-scale chromatin unfolding are physiologically relevant given the high numbers of lac operator repeats involved. In fact, the exact number of lac repressors binding per lac operator has not been determined and there is reason to believe that lac repressor binding may be significantly limited by steric constraints and phasing of lac operators relative to the nucleosome linker DNA. However, we note that a recent study on a transgene array containing a viral promoter with several glucocorticoid hormone response elements observed a very similar type of large-scale chromatin decondensation produced by glucocorticoid receptor (Muller et al., 2001). Ultimately, validation of the physiological significance of our observations of BRCA1-dependent large-scale chromatin unfolding will depend on the outcome of future experiments exploring the mechanisms of unfolding and identifying the biological functions of other transacting factors involved, such as COBRA1.

With these caveats in mind, we find it particularly intriguing that a subset of cancer-predisposing mutations of BRCA1 lead to increased chromatin unfolding and recruitment of COBRA1. Although the genotype–phenotype relationship in cancer-predisposing mutations of BRCA1 remains to be understood, it is generally assumed that most, if not all, BRCA1 mutations lead to loss of the biological functions of the protein. However, the behaviors of the BRCA1 mutants in the chromatin-unfolding assay clearly demonstrate an allele-specific effect. Consistent with this finding, it has been reported that mutations at different locations along the coding sequence of BRCA1 differentially affect the penetrance of BRCA1-dependent breast and ovarian cancer (Gayther et al., 1995; Risch et al., 2001). It remains to be determined whether the three groups of mutations that cause differential effects on chromatin unfolding (Fig. 3) may indeed lead to distinct clinical consequences in terms of risks, types, or prognosis of BRCA1-associated cancers. In particular, it will be interesting to see whether those 3′ mutations that enhance chromatin unfolding exhibit any dominant or semidominant phenotype in cancer genetics. It is conceivable that constitutive decondensation of large-scale chromatin structure may cause additional deleterious effects on genome stability and thus result in more severe clinical consequences in cancer development.

Our study also indicates that BRCT-mediated chromatin unfolding may be tightly regulated. As shown in Figs. 1 and 3, a single BRCT motif is more potent in chromatin unfolding than the larger fragments of the protein, that contain both BRCT repeats. Furthermore, the full-length wild-type BRCA1 only exhibits a moderate chromatin-unfolding activity, whereas the cancer-predisposing mutations in group c (Fig. 3B) that affect the integrity of the BRCT repeats significantly enhance the chromatin-unfolding activity and COBRA1 binding. These results lead us to the following two models that could explain negative regulation of BRCA1-mediated chromatin unfolding. In a “trans-inhibition” model, we speculate that binding of a putative inhibitor (i.e., CtIP) to AD2 region of BRCA1 may prevent BRCA1 from interacting with its cofactors for chromatin unfolding (i.e., COBRA1). In an alternative, “cis-inhibition” model, the two BRCT tandem repeats may form an intramolecular dimer. This in turn may reduce the affinity of both BRCT repeats for their corresponding cofactors. Conceivably, the “superactivating” mutations in group c may prevent binding of the putative inhibitor or the intramolecular interaction between the two BRCT motifs, thus rendering the protein constitutively active for binding to the cofactors that mediate chromatin unfolding.

It is plausible that BRCT-mediated chromatin unfolding may lead to a novel nuclear function of BRCA1 in global reorganization of the genome. However, in light of the known function of the COOH-terminal region of BRCA1 in transcription and DNA repair, the observed chromatin decondensation may represent the first step in BRCA1-mediated regulation of these two nuclear processes (Fig. 7). In such a
Materials and methods

Chromatin unfolding assay
To construct the EGFP-lac-E2F1 and EGFP-lac-p53 fusion expression vectors, the PCR fragments that encode the E2F1 (aa 368–437) and p53 (aa 1–73), respectively, were cloned into the Ascl site in the plasmid p3SS d th Cl EGFps (NYE4) (A.C. Nye and A.S. Belmont, personal communication). The correct orientation of the inserts was identified by colony hybridization and confirmed by DNA sequencing. To construct the lac-BRCA1 plasmids, the sequence for lac repressor was first amplified by PCR from the plasmid NYE4. The lac sequence was cloned into the HindIII–NotI sites of pRC-CMV (Invitrogen), generating pRC-lac. Various BRCA1 fragments and the COBRA1 sequence were amplified by PCR and inserted into the unique Ascl site of pRC-lac.

The chromatin unfolding experiments were performed as previously described (Tumbar et al., 1999). Briefly, AO3_1 cells were transiently transfected with the lac expression vectors using the FuGENE 6 transfection reagent (Roche). The medium was changed 24 h after transfection and cells were immunostained 48 h after transfection. Cells grown on glass coverslips were fixed with 1.6% paraformaldehyde for 30 min in PBS, permeabilized with 0.2% Triton X-100 in PBS for 5 min, and blocked in 1% normal goat serum in PBS for 1 h. The coverslips were then incubated with primary antibodies at room temperature for 1 h, followed by incubation with the appropriate secondary antibodies for 1 h. Unless otherwise specified, a rabbit polyclonal anti-lac repressor antibody (Stratagene) and mouse monoclonal anti-FLAG antibody (Sigma-Aldrich) were used in each immunoprecipitation. Immunoprecipitation was performed overnight at 4°C. The beads were centrifuged at 3,300 rpm for 2 min, and washed three times with washing buffer (50 mM Tris, pH 8.0, 150 mM NaCl, 1% NP-40, 0.1% SDS, and 0.5% sodium deoxycholate). Each wash was performed for at least 30 min. The precipitates were then eluted in 15 μl 2× SDS-PAGE sample buffer. Gel electrophoresis was followed by immunoblotting according to standard procedures.

GST pulldown assay
The PCR fragments encoding various BRCA1 fragments were cloned into pGEX-2T and the constructs were confirmed by sequencing. The GST-BRCA1 proteins were made and purified, with the induction of protein expression performed at 19°C overnight. pcDNA3 vector containing the COBRA1 gene was used for in vitro transcription and translation in the TNT Reticulocyte Lyase system (Promega). The 35S-labeled COBRA1 was translated in vitro according to the manufacturer’s instructions and mixed with 10 μg of the GST-bound bead in 0.5 ml binding buffer (50 mM Tris-HCl, pH 7.5, 150 mM NaCl, 1 mM EDTA, 0.3 mM DTT, 0.1% NP-40 and protease inhibitor tablet). The binding reaction was performed at 4°C overnight and the beads were subsequently washed four times with washing buffer (same as binding buffer except 0.5% NP-40 was used), 30 min each time. The beads were eluted in 10 μl 2× SDS-PAGE sample buffer and the proteins were resolved on 10% denaturing gel. The gel was then dried and exposed to x-ray films for overnight.

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