An insulator loop resides between the synthetically interacting elements of the human/rat conserved breast cancer susceptibility locus MCS5A/Mcs5a

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ABSTRACT

Many low-penetration breast cancer susceptibility loci are found to be located in non-protein-coding regions, suggesting their involvement in gene expression regulation. We identified the human/rat-conserved breast cancer susceptibility locus MCS5A/Mcs5a. This locus has been shown to act in a non-mammary cell-autonomous fashion through the immune system. The resistant Mcs5a allele from the Wistar–Kyoto (WKy) rat strain consists of two non-protein-coding genetic elements that must be located on the same chromosome to elicit the phenotype. In this study, we show the presence of a conserved higher order chromatin structure in MCS5A/Mcs5a located in between the synthetically interacting genetic elements. The looped elements are shown to be bound by CTCF and cohesin. We identify the downregulation of Fbxo10 expression in T cells as a strong candidate mechanism through which the interacting genetic elements of the resistant Mcs5a allele modulate mammary carcinoma susceptibility. Finally, we show that the human MCS5A polymorphisms associated with breast cancer risk are located at both sides of the looped structure and functionally interact to downregulate transcriptional activity, similar to rat Mcs5a. We propose a mechanistic model for MCS5A/Mcs5a in which a CTCF-mediated insulator loop encompassing the TOMM5/Tomm5 gene, resides in between and brings into closer physical proximity the synthetically and functionally interacting resistant genetic variants.

INTRODUCTION

The risk of developing breast cancer involves the interaction between a woman’s inherited genetics and her environment. The genetic component of risk for most common forms of breast cancer defines it as a complex trait consisting of numerous susceptibility alleles and their interactions. Approximately 25 of such alleles have been identified thus far using genome-wide association studies (GWAS) and comparative genetics and each is associated with a low relative risk (1–10). Currently, a major open question regarding these low-penetrance susceptibility alleles is defining their function in regard to breast cancer etiology. The great majority of these alleles are non-protein-coding, which implies that they may modulate gene expression regulation. For example, the alleles of a breast cancer risk variant in the second intron of FGFR2 have been shown to differentially regulate its expression (11). For the vast majority of low-penetrance breast cancer susceptibility alleles, it remains unclear if these are involved in gene expression regulation and what their target genes may be. Understanding mechanistically how these alleles modulate breast cancer susceptibility could go beyond population-based screening and lead to intervention strategies for cancer prevention and therapy.

The non-protein-coding locus Mcs5a/MCS5A modulates breast cancer susceptibility in both rats and women (5). We identified this locus via a comparative genetics approach consisting of linkage analysis in the backcross progeny of the mammary carcinoma susceptible Wistar–Furth (WF) rat strain and the resistant Wistar–Kyoto (WKy) rat strain (12), subsequent genetic fine-mapping using congenic recombinant rat lines (13,14), and a case-control association study using variants in the human orthologous locus (5). When the resistant allele of Mcs5a from the WKy rat strain is introgressed into the...
susceptible genetic background from the WF rat strain, a ∼50% reduction in mammary carcinoma multiplicity is observed (5). The resistant allele acts in a non-mammary cell-autonomous manner, during early mammary carcinoma progression through the immune system (15). We have also shown that the presence of the resistant Mcs5a allele is associated with T-cell homeostasis and functions (15). Specifically, T cells of the Mcs5a resistant congenic line show an increased mitogen-induced proliferation potential and Th1 cytokine production, suggesting that the Mcs5a allele exerts its effect on mammary carcinoma susceptibility through T cells (15). Mcs5a consists of two non-protein-coding, synthetically interacting elements (Mcs5a1, Mcs5a2), which have to be located on the same chromosome to elicit the resistance phenotype (5). The variants associated with breast cancer risk in women are located in the human orthologous loci, MCS5A1 and MCS5A2, respectively, at a distance of ∼60 kb of each other (5). Although identified by comparative genetics, the MCS5A variants resemble GWAS-identified risk alleles as the risk-associated polymorphisms are non-protein-coding, are common in the population and display low- genetic penetrance. The risk-associated variant in MCS5A1 marked by single nucleotide polymorphism (SNP) rs2182317 has four correlated polymorphisms. These are located in a ∼5.7-kb region around the CpG island that is associated with a predicted promoter of FBXO10, a gene encoding the uncharacterized E3 ubiquitin ligase F-box protein 10. Similarly, the MCS5A2 variant marked by SNP rs2182317 encompasses a total of 15 correlated polymorphisms spanning ∼26.1 kb. These are located around the CpG island that is associated with the predicted promoter of the FERM and PDZ domain containing 1 gene, FRMPD1. We have shown previously that between susceptible congenic control and Mcs5a resistant congenic rats, Fbxo10 and Frmpd1 are differentially expressed in the thymus and spleen, respectively, and not in the mammary gland (5). A third gene in the MCS5A locus, the translocase of the outer mitochondrial membrane5 gene, TOMM5, is not directly associated with the breast cancer risk variants of MCS5A1 and MCS5A2, but is located in the 60 kb of sequence that separates them. SNP rs2182317 in MCS5A2 has recently been shown to be among the most significant of a list of 710 candidate breast cancer risk alleles and to modify risk to both Estrogen Receptor-positive and -negative breast cancer (4).

The availability of both human cell based and congenic rat models allows us to address several mechanistic questions regarding the Mcs5a/MCS5A breast cancer susceptibility alleles. First, we mechanistically address the observation that the mammary carcinoma resistance phenotype requires two genetic Mcs5a elements, separated by Tomm5, to synthetically interact on the same chromosome. Next, we show that the synthetic interaction is also required for downregulation of the transcript levels of Fbxo10 in the thymus and in various sorted T-cell populations. Finally, we show that the resistant alleles of the human breast cancer risk-associated variants could functionally interact to downregulate transcriptional activity, resembling transcriptional regulation of Fbxo10 by the interacting genetic elements of the resistant rat Mcs5a allele.

MATERIAL AND METHODS

Animals

The congenic rat lines were established and maintained in an AAALAC-approved facility as previously published (14). All animal protocols were approved by the University of Wisconsin Medical School Animal Care and Use Committee. Congenics are defined as genetic lines developed on a WF-susceptible genetic background and carrying the selected WKy-resistant Mcs5a alleles. The resistant congenic line WF.WKy-Mcs5a ('Mcs5a', line WW) with a decreased 7,12-dimethylbenz(a)anthracene (DMBA)-induced mammary carcinoma susceptibility phenotype is WKy-homozygous at the entire Mcs5a locus (5). The susceptible congenic control line WF.WKy ('susc.') is WF-homozygous at Mcs5a and all other identified Mcs loci. The susceptible congenic lines WF.WKYa-Mcs5a1 ('Mcs5a1', line B3) and WF.WKYa-Mcs5a2 ('Mcs5a2', Line LL) are WKy-homozygous at Mcs5a1 and Mcs5a2, respectively (5). For all experiments, only female animals were used.

Flow cytometry, magnetic cell sorting and T-cell activation

For the gene expression studies, cells were prepared for immunostaining by submerging fresh thymus and/or spleen tissues in cold flow cytometry solution (FCS: PBS with 2% FBS and 25 mM HEPES). Tissues in 1 ml cold FCS were dispersed using a 50 μM Medicin (BD Biosciences) filter grinders and a BD Medimachine for ~1 min. The resulting cell suspension was filtered with 70 μm Ficlon (BD Biosciences) and centrifuged to pellet cells. In sterile 12 ml polystyrene round-bottom Falcon tubes (BD Biosciences), cells were resuspended in cold FCS (25 × 10⁶ cells/ml) and immunostained at 4°C with 0.5 μg antibody-conjugate per 10⁶ cells (25 μg each antibody) for 30 min. Monoclonal antibodies from BD Biosciences included anti-rat CD3 (clone 1F4), CD45RA (clone OX-33), CD4 (clone OX-35) and CD8a (clone OX-8). Cells were washed 1× with 1 ml cold FCS and filtered prior to flow cytometry sorting on a FACSVantageSE (BD Biosciences) at the University of Wisconsin Flow Cytometry Core Facility. For chromatin studies, the enrichment of CD3+ T cells was done by magnetic cell sorting. Single cell suspensions (in cold FCS) of anonymous cryopreserved human peripheral blood mononuclear cell samples and rat (red blood cell-reduced) splenocytes were incubated with CD3 and pan-T cell (OX-52) microbeads (Miltenyi), respectively. The labeled cell suspension was loaded onto a pre-equilibrated MACS cell separation column (Miltenyi), washed and eluted according to the manufacturer’s recommendations. CD3+ T-cell purity was estimated by flow cytometry to be >92%. For T cell culturing and activation, 2 million lymphocytes from spleen were stimulated with 1 μg/ml Concanavalin A.
(conA; Calbiochem/EMD Chemicals) and were cultured in 2 ml RPMI medium containing 10% FBS for 24 h at 37°C in a 95% air/5% CO₂ atmosphere. Unstimulated cells served as a control. Splenocytes were stained with anti-T-cell microbeads (OX52, Miltenyi) and T cells were sorted by positive selection using EasySep™ magnet (Stem Cell Technologies). Purity of T cells was >90% as determined by CD3 labeling and flow cytometry.

**Primary T-cell nucleofections with siRNAs**

Nucleofections were carried out by following the Mouse T-Cell Nucleofector Kit (Lonza) protocol. Briefly, 2 × 10⁶ T cells (from a pool of sorted splenic T cells with equal contribution from four female Mcs5a resistant congenic rats) were pelleted at low speed (10 min, 90g) and resuspended in 100 µl supplemented Nucleofector Solution. Two nanomoles of the desired siRNAs (ONTARGETplus Non-Targeting, and ONTARGETplus SMARTpool CTGF; Dharmacon) were mixed in, followed by nucleofection using program X-001 of the Nucleofector II Device. The cells were cultured in Nucleofection Media for 64–68 h at 37°C.

**Chromosome conformation capture assay**

Rat templates were prepared from T cells of six susceptible congenic control, and six Mcs5a resistant congenic animals. Human templates were prepared from three T-cell samples. Approximately 5 × 10⁶ T cells (or 2 × 10⁶ for the siRNA experiments) were diluted in 40 ml of PBS and 1.7 ml of 37% formaldehyde was added to fix chromatin. The solution was allowed to fix for 10 min at room temperature. To quench the reaction, 2.7 ml of 2 M glycine was added to each tube and incubated for 5 min at room temperature, after which the tubes were placed on ice for 15 min. To burst the cells and expose the fixed chromatin, the cells were centrifuged for 10 min at 800 g and resuspended in 0.5 ml ice-cold lysis buffer, containing 10 mM Tris–HCl pH 8.0, 10 mM NaCl, 0.2% Nonidet P-40 (NP-40), 1 x protease inhibitors (Roche Applied Science). The reaction was allowed to proceed on ice for 10 min. The cells were gently lysed via dounce homogenization using a tight pestle for 1 min on ice. The reaction sat on ice for 1 min and homogenization was repeated for 30 s. Next, the lysis buffer was removed, the nuclei were washed in 0.5 ml of 1 x restriction enzyme buffer (NEB3; New England Biolabs) and resuspended in 400 µl of 1 x restriction enzyme buffer containing 0.1% SDS. This mixture was incubated at 65°C for 10 min. To sequester SDS, Triton X-100 was thoroughly mixed in to a final concentration of 1%. To digest the DNA and obtain cross-linked DNA fragments, 400 U of BglII restriction enzyme (New England Biolabs) was added. The ligation was incubated overnight at 37°C under slow rotation. The following day, SDS was added to a concentration of 2%. The mixture was incubated at 65°C for 30 min. The ligation was done in a final volume of 7.5 ml, in the presence of 50 mM of Tris–HCl (pH 7.5), 10 mM of DTT, 10 mM of MgCl₂, 1% Triton X-100, 0.11 mg/ml of BSA, 1.07 mM of ATP and 4000 U of T4 DNA ligase (New England Biolabs). The reactions were incubated at 16°C for 4 h. To remove any residual proteins, 33 µl of Proteinase K (15 mg/ml; Roche Applied Science) was added to each tube and incubated at 65°C overnight. An additional 33 µl of Proteinase K was added and the mixture was incubated for 2 h at 42°C. Finally, the ligated DNA products were isolated using two phenol/chloroform (1:1) extractions, and one chloroform extraction in 7.5 ml volume, followed by ethanol precipitation. The pellets were resuspended in 1 ml of dH₂O containing RNase A (Roche Applied Science) in a final concentration of 10 µg/ml. Following a 30 min incubation at room temperature, final phenol/chloroform and chloroform extractions were performed, followed by ethanol precipitation. The pelleted DNA was washed five times with 70% ethanol and dissolved in 400 µl of dH₂O. The concentration of template from each individual tissue sample was determined by agarose gel electrophoresis, quantifying the template band by comparing it to the known amount of the highest band of a 1-kb DNA ladder (NEB). Equal amounts of each template were pooled. All pools were diluted to the same concentration. The amount to use in a PCR detection reaction (1 µl) was determined empirically. For preparing the control DNA templates, Bacterial Artificial Chromosome (BAC) DNA templates corresponding to the human MCS5A and rat Mcs5a regions (human: RP11-65G18; rat: CH230-298P15 and CH230-303N12) were obtained from Children’s Hospital Oakland Research Institute (CHORI). The BAC DNA was isolated using a Plasmid Maxi kit (Qiagen) and 500 ng was digested overnight at 37°C using 5 U of BglII restriction enzyme (NEB). The restriction fragments were recovered using phenol/chloroform and chloroform extractions, followed by ethanol precipitation. The digested BAC DNA was religated by adding 5 µl of 10 x ligation buffer and 2 U of T4 DNA Ligase (Roche Applied Science). The mixture was incubated overnight at 16°C. The religated fragments were recovered using phenol/chloroform and chloroform extraction, followed by ethanol precipitation. The DNA concentration was measured by nanodrop and diluted to 10 ng/µl. A working stock of 1:1000 dilution was prepared. The amount to use in a reaction (0.02 µl) was empirically determined. Empirically determined DNA template was added to each reaction well and stored on ice. The PCR mixture was prepared on ice, and mixed into the pre-cooled DNA templates. The reaction was done in the presence of 1 x Herculase reaction buffer, 0.2 mM of each NTP, 0.4 µM of each primer, 0.3 µl of Herculase Enhanced polymerase (5 U/µl; Stratagene) in a total volume of 25 µl. Primer sequences are listed in Supplementary Table S1. The PCR was performed using a Peltier Thermocycler (PTC-225; MJ Research), using the following cycling conditions: 95°C for 1 min, 35 cycles of 95°C for 30 s, 60°C for 30 s, 72°C for 20 s, followed by a final extension of 72°C for 8 min. When the program was complete, 5 µl of 6 x loading buffer (15 % w/v Ficoll-400, 0.25% w/v Xylene Cyanol FF) was added to the reaction and 12 µl of each reaction was analyzed by agarose gel electrophoresis. The gels were stained by soaking in a 1 µg/ml ethidium bromide solution for 10 min, and
destained in dH2O for 15 min. An image was taken under UV-exposure using the Gel Doc XR system (Biorad). PCR band intensity was quantified using ImageQuant software (GE Healthcare).

Chromatin immunoprecipitation

The JURKAT (clone E6-1; CD4+ leukemic T-lymphocytic cell line) cell line was obtained from ATCC and grown in GIBCO RPMI-1640 media (Invitrogen) supplemented with 10% decomplemented Hyclone Fetal Bovine Serum (FBS; Thermo Scientific) and 1% Penicillin/Streptomycin (Invitrogen). JURKAT cells were cultured to a density of ~3 x 10^6 cells/ml. Roughly 2 x 10^6 JURKAT cells or splenic T cells from female susceptible congenic control and Mct5a resistant congenic rats were fixed using formaldehyde to a final concentration of 1% at room temperature for 10 min. Fixation was followed by centrifugation (5 min, 800 g) to pellet the cells. The cells were washed twice with PBS. The chromatin immunoprecipitation (ChiP) assay was done using the ChIP Assay Kit (Upstate/Millipore) following manufacturer’s recommendations with one exception: the SDS lysis buffer was reconstituted to contain 0.1% of SDS instead of 1%. For one assay 4 x 10^6 cells were resuspended in 200 µl of SDS-lysis buffer. Chromatin was sheared to 200- to 1000-bp fragments by sonication using three pulses of 10 s on the third power level of a Branson sonifier (Cell disruptor 185; Branson). Roughly 1% of the sonicated chromatin was set aside as the input (positive control) sample. Immunoprecipitations were done on the remainder of the sonicated chromatin using the following antibodies: rabbit monoclonal to CTCF (07–729; Millipore), rabbit polyclonal to Rad21 (cohesin; ab992; Abcam), and rabbit control IgG (negative control; ab46540; Abcam). Immunoprecipitations and input DNA were recovered by phenol:chloroform and chloroform extractions, followed by ethanol precipitation. Input- and immunoprecipitation-recovered DNAs were tested in a dilution series as templates in detection PCRs. Empirically determined DNA template was added to each reaction well and stored on ice. The PCR mixture was prepared on ice, and mixed into the pre-cooled DNA templates. The reaction was done in the presence of 1 x Herculase reaction buffer, 0.2 mM of each NTP, 0.4 µM of each primer, 0.3 µl of Herculase Enhanced polymerase (5 U/µl) in a total volume of 20 µl. The PCR was performed using a Peltier Thermocycler, using the following cycling conditions for the human primers: 95°C for 1 min, 32 cycles of 95°C for 30 s, 58°C for 30 s, 72°C for 15 s, followed by a final extension of 72°C for 3 min, and the following cycling conditions for the rat primers: 95°C for 1 min, 32 cycles of 95°C for 30 s, 56°C for 30 s, 72°C for 15 s, followed by a final extension of 72°C for 3 min. Primer sequences are listed in Supplementary Table S1. When the program was complete, 5 µl of 6X loading buffer (15% w/v Ficoll-400, 0.25% w/v Xylene Cyanol FF) was added to the reaction and 10 µl of each reaction was analyzed by agarose gel electrophoresis. The gels were stained by soaking in a 1 µg/ml ethidium bromide solution for 10 min, and destained in dH2O for 10 min. Image was taken under UV exposure.

RNA extraction and quantitative real-time PCR

Congenic female rats (11 weeks of age) of line WF.WKy (susc.), line WW (Mct5a), line B3 (Mct5a1) and line LL (Mct5a2) were used as tissue donors. RNA was extracted from snap-frozen tissues and sorted cell samples using the MagMax-96 Total RNA isolation kit (Ambion) according to manufacturer’s directions. To synthesize cDNA from 500 ng total RNA treated with TURBO-free DNase (Ambion), the reverse transcriptase Superscript II kit (Invitrogen) was used according to manufacturer’s directions. Quantitative real-time PCR was used to quantify transcript levels. TaqMan quantitative PCR primers and probes (available in Supplementary Table S1) were either designed using Primer Express v 2.0 (ABI/Applied Biosystems) according to developer’s specifications, or ordered as made-to-order assays (Fbxo10 Rn01439970_m1; Tomm5 Rn01436622_g1; Mcart1 Rn01764629_g1; Dcaf10 Rn01764617_m1; ABI/Applied Biosystems). A microiler of cDNA (~12.5 ng RNA equivalent cDNA) was used in a 16 µl TaqMan QPCR reaction. Reaction components were 1 x TaqMan Buffer A (Applied Biosystems), 5.5 mM MgCl2, 400 µM each dATP, dCTP, dGTP, dTTP and 0.4 U Taq Gold DNA Pol (ABI). For the custom designed assays, 500 nM each experimental primer, 200 nM TaqMan experimental probe, 60 nM each Gapdh primer, 120 nM rodent Gapdh probe, were used per reaction. For the made-to-order assays, 0.8 µl of the experimental assay and 0.4 µl of the ActB standard assay were used per reaction. Cycling conditions were 50°C for 2 min, 95°C for 10 min, followed by 40 cycles of 95°C for 15 s and 60°C for 1 min. FAM (Fbxo10, Fmm1, Tomm5 probe) and VIC (Rodent Gapdh or ActB probe, ABI) fluorescence values were measured in real-time on an ABI 7900 real-time PCR machine using ABI SDS v2.2 software. Quantities of transcripts were measured by comparison of Ct values with a standard curve calculated from serial dilutions made from reverse transcriptase (RT) reactions that contained 2 µg of total RNA. Sample measurements are an average of four replicates. Sample measurements were normalized by dividing the gene-specific transcript quantity over the rodent Gapdh or ActB transcript quantity detected in the same reaction. Ratios were scaled to the ratio of the susceptible congenic control sample from the same experiment. Data were analyzed using Kruskal–Wallis or Mann–Whitney non-parametric tests. Transcript levels were only statistically compared if the samples were run in the same assay plate.

5’ RNA ligase mediated-rapid amplification of cDNA ends (5’ RLM-RACE)

To localize the exact transcriptional start site (TSS) of the Fbxo10 transcripts, the 5’ RLM-RACE assay was performed using the FirstChoice RLM-RACE kit (Ambion) following manufacturer’s recommendations. Thymic or splenic total RNA from five susceptible congenic
control and five Mcs5a resistant congenic rats was used. Equal amounts of RNA (as determined by Nanodrop quantification) were pooled and 10 μg of each pool was used as input RNA into the reaction. To identify the human FBXO10 TSS, 10 μg total RNA from breast, thymus or spleen (Ambion) were used as input. To amplify the 5'-ends of the Fbxo10/FBXO10 transcripts, a nested PCR was performed with the gene-specific primers (available in Supplementary Table S1) annealing to the translational start codon (ATG)-containing exon that is presumably shared by all Fbxo10/FBXO10 transcripts. A typical PCR product yielded multiple bands on an agarose gel, indicative of multiple TSS. The products were blunt-cloned using the ZEROBLENII vector system (Invitrogen) for subsequent resequencing. In total, 150 individual rat thymus or spleen TSS products and 54 individual human thymus, spleen or breast TSS products were analyzed by resequencing. The subcloned sequences were identified by performing BLAT searches using the UCSC genome browser.

Vector construction, transient transfection, luciferase assays and Bradford assays

Genomic fragments with the appropriate polymorphisms were amplified from MCS5A1 and MCS5A2 heterozygous genomic DNA using the Platinum HiFi Taq polymerase (Invitrogen) and primers containing the restriction sites (BamHI, Sall) used for cloning (available in Supplementary Table S1). First, the FBXO10-TSS constructs were made by inserting the BamHI-, Sall-digested genomic FBXO10-TSS fragment using the XhoI and BglII site located upstream of Luc in pGL3-Basic (Promega). The S and R versions of the FBXO10-TSS construct were digested with BamHI and Sall to insert the BamHI-, Sall-digested genomic fragments containing the MCS5A2 polymorphisms. To ensure integrity, all constructs underwent visual inspection by restriction enzyme digestion with NotI and SacI (Supplementary Figure S2), as well as resequencing of both inserts. The JURKAT (clone E6-1; CD4+ leukemic T-lymphocytic cell line) cell line was obtained from ATCC and grown in GIBCO RPMI-1640 media (Invitrogen) supplemented with 10% decomplemented Hyclone Fetal Bovine Serum (FBS; Thermo Scientific) and 1% Penicillin/Streptomycin (Invitrogen). JURKAT cells were cultured to a density of ∼3 × 10^6 cells/ml. The cells were diluted into fresh prewarmed RPMI/FBS media to a density of 0.5 × 10^6 cells/ml. A total of 500 μl cells (approximately 250,000 cells) were distributed into each well of a 24-wells culture plate. The cells were allowed to rest for at least 3 h. Transient transfection was performed using LTX Lipofectamine reagent (Invitrogen) according to manufacturer's recommendations. The DNA concentration of each construct was verified before each transfection. All DNA concentrations were between 90 and 110 ng/μl. For each transfection, 500 ng of construct DNA was diluted with a mixture of 100 μl GIBCO OPTI-MEM reduced serum media (Invitrogen), 5 μl of PLUS reagent (Invitrogen) and 5 ng of Renilla internal control vector pRL-TK (Promega). Following a 5-min incubation at room temperature, 5 μl of LTX Lipofectamine was added. The transfection solution was mixed by pipetting up-and-down, followed by a 30-min incubation at room temperature. A total of 100 μl of the transfection solution was added to each well containing the cells. The cells and the transfection solution were mixed by gently shaking the plate in multiple directions. The cells were cultured for 44–48 h. The Luciferase assay was performed using the Dual-Luciferase Reporter Assay kit (Promega) according to the manual. The cells were harvested, spun down and washed with PBS. The cell pellet was lysed in 100 μl 1× Passive Lysis Buffer and incubated at room temperature for 15 min. An amount of 20 μl of the lysate was measured in a 12 × 75 mm disposable culture tube (Fisher Scientific). Each transfection was measured three times, using a Luminometer Monolight 3010 (BD Biosciences). The Luminometer was set as follows: inject 100 μl of the Luciferase substrate, 2 s delay, 10 s of measuring time, inject 100 μl of the Stop-and-Glo solution, 2 s delay, 10 s measuring time. The remainder of each lysates was used to quantify total protein content using Bradford assays. To determine total protein content of the lysate, 5 μl of the lysate was mixed with 250 μl 1× QuickStart Bradford Dye reagent (Biorad) in a 96-well flat bottom plate. The plate was incubated for 15 min at room temperature. The optical density (OD) at 560 nm wavelength was determined using a Spectrophotometer for microtiter plates. The OD was correlated to a standard curve derived from a dilution series using Bovine Serum Albumin (BSA). Relative Luciferase/Renilla activity of each construct was determined by dividing the Luciferase/Bradford ratio over the Renilla value. Each construct was subsequently normalized against the relative Luciferase activity of the pGL3-control construct present in each 24-well plate. The SS and RR version of each construct were always present in the same 24-well plate. The normalized values for the SS and RR version of each construct from at least six transfections were tested for a significant difference using the Wilcoxon rank sum test.

RESULTS

The Chromosome conformation capture assay identifies a conserved higher order chromatin loop within Mcs5a/MCS5A

To investigate if chromatin looping could provide a mechanistic basis for the synthetic (genetic) interaction between the genetic elements in Mcs5a1 and Mcs5a2, we applied the chromosome conformation capture (3C) assay to the rat and human Mcs5a/MCS5A locus (Figure 1A). The 3C assay measures the relative (looping) interaction frequency of crosslinking captured, enzymatically digested (BgIII), ligated chromatin fragments (16). For the rat Mcs5a locus, the fixed fragment was initially chosen to encompass the genomic region orthologous to the 5.7-kb region in the human genome harboring the correlated polymorphisms in MCS5A1 that are associated with breast cancer risk (5). This fixed fragment also contained the
Figure 1. Analysis of the higher order chromatin structure of rat and human Mcs5a/MCS5A. (A) Maps of the rat and human Mcs5a/MCS5A locus. The relative position of Mcs5a1/MCS5A1, Mcs5a2/MCS5A2, all BglII restriction site, the transcripts Fbxo10/FBXO10, Tomm5/TOMM5 and Frmpd1/FRMPD1, and the human breast cancer risk-associated variants is indicated. The primers matching the fixed fragments used to generate panels (B–E) are indicated by black vertical triangles. (B–D) 3C profiles of the Mcs5a locus in T cells of susceptible congenic control (susc.) (continued)
CpG island associated with a predicted Fbxo10 promoter. Relative interaction frequencies with the majority of BglII fragments in Mcs5a were determined. Restriction fragments located close to the fixed fragment yielded a high relative interaction frequency that decreased with increasing genomic distance, indicative of random ligation, but no looping (Figure 1B). Similarly, taking a nearby BgIII fragment (two BgIII fragments equaling ~5 kb upstream of the one containing the predicted Fbxo10 promoter) as the fixed fragment yielded no looping (Supplementary Figure S1A and B). However, when the fixed fragment was shifted further upstream (five BgIII fragments equaling ~10 kb) of the one containing the predicted Fbxo10 promoter, two areas of locally increased relative interaction frequency were identified, indicative of looping (Figure 1C). Both looping areas are located in the Mcs5a2 region, close to the Mcs5a1–Mcs5a2 border. For confirmation, a BgIII fragment in the first looping area was chosen as the fixed fragment. The profile showed the same areas of looping, namely the previous fixed fragment in Mcs5a1 and the second looping area in Mcs5a2 (Figure 1D). Furthermore, none of the loops appeared to be influenced by the susceptible or resistant Mcs5a genotype as the 3C profiles show the same trends. All 3C experiments described above were performed on primary T cells, as we consider this a specific cell type of Mcs5a activity (15).

The spatial organization of the Mcs5a locus in the mammary gland was found to be similar to the profile in T cells, as the same regions were found to be involved in the chromatin looping interactions (Supplementary Figure S1C and D).

The 3C profile of the orthologous human MCS5A locus was found to be highly similar to rat Mcs5a (Figure 1E). No looping was found with the fixed fragment containing the MCS5A1 correlated polymorphisms that associate with breast cancer risk (Figure 1E). This fixed fragment also contained the CpG island associated with the predicted FBXO10 promoter. With the fixed fragment shifted upstream (two BgIII fragments equaling ~10 kb) of the CpG island associated with the predicted FBXO10 promoter, two areas of looping were found to be located in MCS5A2, close to the MCS5A1–MCS5A2 border (Figure 1E). The similarities in the 3C profiles of the rat and human Mcs5a/MCS5A suggest that the higher order chromatin structure of Mcs5a/MCS5A is evolutionary conserved.

CTCF and cohesin binding is confirmed in all interacting chromatin looping fragments

The position of the looping elements at either side of the Tomms/TOMMS gene, thus spatially isolating this gene, suggests a function as an insulator loop. The CCCTC-binding factor (CTCF) protein is widely known for its role as a vertebrate insulator protein (17). Additionally, CTCF has been shown to be essential for long-distance enhancer-promoter looping (18). Recently, the cohesin protein complex has also been shown to be essential for long-range chromatin looping in the developmentally controlled IFNG locus (19). We sought to investigate whether CTCF and/or cohesin binding could also underlie the observed higher order chromatin structure of MCS5A/Mcs5a. Using a ChIP assay with antibodies against CTCF and Rad21, a DNA-binding subunit of the cohesin complex, association of both CTCF and cohesin to all three interacting chromatin looping fragments was confirmed in both a human T-lymphocytic cell line (JURKAT, Figure 2A) and rat primary T cells (Figure 2B). No evidence of CTCF or cohesin binding was found in a location outside of the looping fragments (Figure 2A and B). CTCF binding was confirmed on a site in the H19 locus previously shown to bind CTCF in human (20) and rat (21). The location of the CTCF-binding site in the looping fragment MCS5A1 and the first looping fragment in MCS5A2 coincides with CTCF binding previously identified in a genome-wide ChIP-seq study (22). To test if CTCF is necessary for the higher order chromatin structure, a short-interfering RNA (siRNA) approach was undertaken to deplete primary rat T cells with CTCF. Nucleofection of siRNA against CTCF resulted in a significant reduction of CTCF transcript level (measured after 64–68 h) as compared with nucleofection with control siRNAs (Figure 2C). Interestingly, the transcript level of Tomms, the gene residing within the looped segments was also significantly reduced after nucleofection with siRNAs against CTCF (Figure 2D). The transcript levels of a housekeeping gene, Gapdh and two genes close to the Mcs5a locus, Meart1 and Dcaf10 were not affected by CTCF depletion (Figure 2D). Using the ChIP assay a reduction of enrichment of the Mcs5a1 3C looping fragment in the cells nucleofected with siRNAs against CTCF as compared with the cells nucleofected with control siRNAs was observed, consistent with CTCF depletion (Figure 2E).

Figure 1. Continued and Mcs5a resistant congenic (Mcs5a) rats. The position of the fixed fragments is indicated with a black bar and F, and the primer within the fixed fragment is indicated with a vertical triangle. In (B) the fixed fragment contains the CpG island associated with the predicted promoter of Fbxo10 in Mcs5a1. In (C) the fixed fragment is located five BgIII restriction fragments (~10 kb) upstream of the previous fixed fragment that contains the CpG island in Mcs5a1. In (D) the fixed fragment was located in the first looping fragment in Mcs5a2. (E) Comparison of the rat and human Mcs5a/MCS5A 3C profile. Dashed lines represent the 3C profile with the fixed fragment containing the predicted Fbxo10/FBXO10 promoter in Mcs5a1/MCS5A1 and does not display looping to Mcs5a2/MCS5A2. In the human, this fixed fragment also contains the four correlated MCS5A1 polymorphisms associated with breast cancer risk. Solid lines represents the 3C profile with the fixed fragment located five (rat) or two (human) BgIII restriction fragments (~10 kb) upstream of the BgIII fragment containing the predicted Fbxo10/FBXO10 promoter in Mcs5a1/MCS5A1. The chromatin fragments in Mcs5a2/MCS5A2 looping to the fixed fragment in Mcs5a1/MCS5A1 are indicated with PEAK 1 and PEAK 2. Graphed are the average ± SEM relative interaction frequencies of a fixed BgIII fragment with other BgIII fragments in Mcs5a/MCS5A. The genomic distance (in kilobases) represents the distance of the midpoint of a BgIII fragment to the Mcs5a1/MCS5A1-Mcs5a2/MCS5A2 border.
Finally, using the 3C assay a significant reduction in relative interaction frequency between the Mcs5a1 and first Mcs5a2 looping fragments and between the Mcs5a1 and second Mcs5a2 looping fragments was observed in the cells nucleofected with siRNAs against CTCF, as compared with cells nucleofected with control siRNAs (Figure 2F). We conclude that CTCF is necessary for the higher order chromatin structure of Mcs5a. Disruption of this structure resulted in lower transcript level of Tomm5.

Mcs5a controls the transcript level of Fbxo10 in T cells

We previously reported that the non-protein-coding resistant Mcs5a allele is associated with lower Fbxo10 and higher Frmpd1 transcript levels in thymus and spleen, respectively (5), and that the resistance phenotype is mediated by the immune system (15). Here, we explored by quantitative real-time PCR using TaqMan assays, if the difference in transcript levels of these two genes, like the mammary carcinoma resistance phenotype, is controlled by the two synthetically interacting genetic

Figure 2. CTCF and cohesin bind to the looping fragments in Mcs5a/MCS5A and CTCF is necessary for the higher order chromatin structure. (A and B) The Mcs5a1/MCS5A1 and Mcs5a2/MCS5A2 loci are depicted as black lines. The light gray bars within the black lines are the CpG islands associated with the promoters of the Fbxo10/FBXO10, Tomm5/TOMM5 and Frmpd1/FRMPD1 genes, respectively. The locations of the three interacting chromatin looping fragments identified in the 3C assay are indicated by light gray blocks. In the human, the location of known CTCF sites from a genome-wide CTCF ChIP-seq study is indicated (22). Amplicons within and outside the looping Mcs5a/MCS5A fragments, as well as an amplicon in the H19 locus (as a positive control) were analyzed by PCR on CTCF (C), cohesin (R; Rad21) and IgG (I; negative control) antibody immunoprecipitated chromatin samples, and an input (IN, positive control) sample, prepared from JURKAT cells (human) or primary rat splenic T cells (rat). Each gel image is accompanied by a 100-bp DNA ladder of which the lower three bands (100, 200, 300 bp) are shown. CTCF and cohesin binding was found in all looping fragments and the H19 positive control locus, but not in the Mcs5a2 fragment not located in a looping fragment. (C and D) Transcript level of CTCF, Tomm5, Gapdh, Mcart1 and Dcaf10 (normalized to ActB) measured 64–68 h after nucleofection of rat primary T cells with control siRNAs (siCONTROL; light gray bars) and siRNAs against CTCF (siCTCF; dark gray bars). The transcript level of CTCF and Tomm5 was significantly reduced (indicated with an asterisk) in the siCTCF-treated T cells. The transcript levels of the housekeeping gene Gapdh and genes Mcart1 and Dcaf10 located adjacent to the Mcs5a locus were not affected. (E) ChIP analysis of CTCF binding to the looping fragment in Mcs5a1 and the negative control fragment in Mcs5a2 in primary rat T-cells 64–68 h after nucleofection with control siRNAs and siRNAs against CTCF. CTCF binding to the Mcs5a1 looping fragment was reduced in the siCTCF-treated T cells. (F) 3C analysis of the Mcs5a locus in primary rat T cells 64–68 h after nucleofection with control siRNAs (light gray line) and siRNAs against CTCF (dark gray line). The looping fragment in Mcs5a1 was taken as the fixed fragment (indicated with a black bar and F). Looping of the fixed fragment to the looping fragments in Mcs5a2 was significantly reduced (indicated with an asterisk) in the siCTCF-treated T cells.

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elements of Mcs5a. Hence, thymic Fbxo10 and splenic Frmpd1 transcript levels were compared between the susceptible congenic control line (WF-homozygous at Mcs5a), the Mcs5a resistant congenic line (WKy-homozygous at Mcs5a) and the Mcs5a1 and Mcs5a2 susceptible congenic lines (WKy-homozygous at Mcs5a1 or Mcs5a2, respectively). The Fbxo10 transcript level was significantly reduced ($P = 0.01$) in the thymus of the Mcs5a congenic resistant animals and was not significantly different in thymus tissue from either the Mcs5a1 or Mcs5a2 susceptible congenic lines when compared to the susceptible congenic control line (Figure 3A). Thus, having lower thymic Fbxo10 transcript levels required the same synthetic interaction between Mcs5a and Mcs5a2 as was necessary for the mammary carcinoma resistance phenotype. Conversely, higher Frmpd1 transcript levels in the spleen did not require the synthetic interaction, as the Mcs5a resistant congenic line, and both Mcs5a1 and Mcs5a2 susceptible congenic lines had higher splenic Frmpd1 transcript levels as compared with the susceptible congenic control line (Figure 3A).

Subsequently, Fbxo10 transcript levels were monitored in various classes of sorted and unsorted thymocytes (Figure 3B). The reduction in Fbxo10 transcript levels associated with the resistant Mcs5a allele was observed in unsorted ($P = 0.03$), CD4+CD8+ ($P = 0.02$), CD4-CD8+ ($P = 0.03$) and CD4-CD8- ($P = 0.002$), but not in CD4-CD8- thymocytes ($P = 0.77$). The Fbxo10 transcript level difference was also found in CD3+ T cells sorted from the spleen ($P = 0.001$), whereas other classes of sorted and unsorted splenocytes did not display the differential transcript levels (Figure 3C). The Fbxo10 transcript level difference in splenic T cells persisted upon culturing ($P = 0.0003$) and concanavalin A (conA) stimulation ($P = 0.01$; Figure 3D).

C9ORF10/C9orf105 is a spliced transcript located in the 60-kb sequence between the human MCS5A1 and MCS5A2 risk-associated polymorphisms on the rat Mcs5a1–Mcs5a2 border and was recently annotated as the translocase of outer mitochondrial membrane 5 gene, TOMM5/Tomn5. We checked if the transcript level of C9orf105 is associated with the resistant rat Mcs5a allele in the immune system. Tomn5 transcript levels were not different between susceptible congenic control and Mcs5a resistant congenic rats in whole spleen ($P = 0.18$) and thymus ($P = 0.40$) tissues (Figure 3E). Similarly, in primary T cells sorted from the spleen, Tomn5 was not differentially expressed ($P = 0.94$; Figure 3E). In cultured T cells from susceptible congenic control and Mcs5a resistant congenic rats, the transcript levels of Tomn5 were found to be different ($P = 0.008$; Figure 3F). In conA-stimulated cultured T-cells Tomn5 transcript levels were not found to be different ($P = 0.95$) between the Mcs5a genotypes. Interestingly, in conA-stimulated T cells, Tomn5 transcript levels were found to be strongly increased (~3-fold) when compared to unstimulated cultured T-cell samples from the rat lines harboring both Mcs5a genotypes ($P < 0.0001$ for both the susceptible and the resistant congenic line; Figure 3F). For comparison, Fbxo10 transcript levels in conA-stimulated T cells were not different ($P = 0.45$) when compared to the unstimulated cultured T-cell samples from the susceptible congenic control line, but were increased (~1.5-fold, $P = 0.003$) in the resistant congenic line (Figure 3D). It should also be noted that Frmpd1 failed to amplify in TaqMan assays on primary, cultured unstimulated and cultured activated T-cell samples (data not shown).

Taken together, the gene expression data implicate that lower Fbxo10 transcript levels, and not Frmpd1 and Tomm5 transcript levels in the immune system are associated with the Mcs5a resistant allele. Furthermore, these data suggest that Tomm5 and Fbxo10 and Frmpd1 may play a role in T-cell activation.

The TSS cluster of FBXO10/Fbxo10 is located in MCS5A1

To understand the regulation of the Fbxo10/FBXO10 gene it is important to know the exact TSS. This facilitates localization of the putative promoter. Two areas of transcriptional initiation of the Fbxo10/FBXO10 gene in rats and humans are annotated in the UCSC genome browser, namely the CpG island in Mcs5a1/MCS5A1 and an area close to the Mcs5a1/MCS5A1–Mcs5a2/MCS5A2 border (Figure 4A). To identify the TSS of the Fbxo10 gene, we performed the 5′ RLM-RACE assay (RNA Ligase Mediated-Rapid Amplification of cDNA Ends). The assay makes use of an RNA adapter that is ligated to the de-CAP-ped 5′-end of transcripts, followed by a RT reaction to make cDNA. To amplify just the 5′-ends of the Fbxo10 gene, a nested PCR reaction was performed with primers annealing to the translatonal start codon (ATG)-containing exon and universal primers annealing to the 5′ RNA-adapter. The assay was done on pools of thymus or spleen RNA from four susceptible congenic control and four Mcs5a resistant congenic rats and on human RNA samples from thymus, spleen and breast tissue. A typical PCR product yielded multiple bands on an agarose gel, indicative of multiple TSSs. In total, 150 rat and 54 human TSS clones were sequenced to elucidate the exact start position of the Fbxo10/FBXO10 transcripts. In the rat, 14 TSS positions were found, all located in the CpG island of the Mcs5a1 locus (Figure 4B). In the human, three TSS positions were found, again, located in the CpG island of the MCS5A1 locus (Figure 4B). It should be noted that in the human breast RNA sample the FBXO10 TSS cluster was found to be located close to the MCS5A1/MCS5A2 border, indicating that the assay was able to identify such cases. Identification of the Fbxo10/FBXO10 TSS cluster in the CpG island of Mcs5a1/MCS5A1 suggests that potential regulatory elements (e.g. promoter or enhancers) may be located in the same region. The human TSS cluster is located within 150 bp of SNP rs6476643 (Figure 4B), which is one of the breast cancer risk-associated SNPs found in MCS5A1 (5).
Figure 3. Lower Fbxo10 transcript level in T cells is associated with the mammary carcinoma resistance phenotype mediated by Mcs5a. (A) The thymic Fbxo10, and not splenic Frmpd1 transcript level is associated with the resistant Mcs5a allele. Graphed are average ± SEM thymic Fbxo10 (dark gray) and splenic Frmpd1 (light gray) transcript levels (normalized to Gapdh) scaled to the thymic Fbxo10 or splenic Frmpd1 transcript level of the susceptible congenic control line. Significantly (P < 0.05) different Fbxo10 and Frmpd1 transcript levels as compared with the susceptible congenic (continued)
**Reporter constructs identify functional interactions between the MCS5A1 and MCS5A2 risk-associated polymorphisms**

In rat T cells, the Fbxo10 transcript level is under control of the synthetically interacting resistant Mes5a genetic elements. This observation led to the hypothesis that the polymorphisms of the resistant risk-associated alleles of MCS5A1 and MCS5A2 could interact similarly to downregulate Fbxo10 transcript levels in human T cells. To investigate this, Luciferase reporter assays were performed using a series of constructs containing selected MCS5A1 and MCS5A2 alleles. All constructs described below were visually inspected by restriction enzyme digest to ensure integrity (Supplementary Figure S2). A human T-lymphocytic cell line (JURKAT) was transiently transfected with each construct in the presence of a Renilla expressing vector as a control for transfection efficiency.

First, a 1464-bp fragment of MCS5A1 including the previously identified FBXO10 TSS cluster was inserted upstream of the Luciferase (Luc+) reporter gene in the pGL3-basic vector (Figure 5A). Two versions of the construct were made (Figure 5B), one containing the resistant (R) allele of rs6476643 (G) and one containing the susceptible (S) allele of rs6476643 (T). The transcriptional activity of the R and S version of the FBXO10 TSS-Luciferase construct is not significantly different (P = 0.45; Figure 5C).

Subsequently, the R and S Fbxo10 TSS-Luciferase constructs were modified to harbor MCS5A2 fragments containing the resistant or susceptible allele of one or two MCS5A2 SNPs (Figure 5A). The MCS5A2 fragments were inserted downstream of Luc+ (Figure 5B). The 15 correlated MCS5A2 polymorphisms were integrated into 10 MCS5A2 fragments of on average ~990 bp in length (Supplementary Table S2). This yielded a series of 10 constructs, each present in two versions, namely SNP rs6476643 susceptible plus MCS5A2 susceptible (SS) and SNP rs6476643 resistant plus MCS5A2 resistant (RR).

The transcriptional activity of the entire series of 10 RR constructs is significantly lower compared to the entire series of 10 SS combination constructs (Wilcoxon rank test P < 0.05; Figure 5D).

Next, the transcriptional activity of the SS and RR versions was compared for each construct (Figure 5D). The transcriptional activity was not significantly different between the SS and RR versions of constructs 1, 2, 5, 7, 8 (P > 0.05, Supplementary Table S2). However, the activity of constructs 3, 4, 6, 9 and 10 were significantly lower in the RR version compared to the SS version (P < 0.05, Supplementary Table S2). In addition, the MCS5A2 SNPs rs62534439 (from construct 4), and rs62534443 and rs62534444 (from constructs 5 and 6) were also represented in constructs 4a and 5a, respectively, again resulting in lower activities for the RR versions compared to the SS versions (Supplementary Figure S3), although only significant for construct 4a (P < 0.05, Supplementary Table S2).

The MCS5A2 fragments in construct 4, 6 and 10 (yielding lower activity in the RR combination constructs) were also tested on their own for transcriptional activity by transferring them to the cloning site upstream of Luc+ (instead of the FBXO10 TSS-containing fragment). All of these constructs had lower transcriptional activity as compared with the FBXO10 TSS-containing fragment (Supplementary Figure S4). The S and R version of these constructs were not significantly different (Supplementary Figure S4), suggesting that the R allele of the polymorphisms in these fragments act only when combined with the R allele of the MCS5A1 SNP rs6476643 in the FBXO10 TSS-containing fragment. The observation that the MCS5A1 and MCS5A2 R alleles functionally interact to downregulate transcriptional activity is consistent with our observation of lower Fbxo10 transcript levels in T cells of Mes5a resistant congenic animals compared to susceptible congenic control animals.

**Figure 3.** Continued

control line is indicated with one and two asterisks, respectively. Sample sizes were: ssuc., n = 9; Mes5a, n = 6; Mes5a1, n = 9; Mes5a2, n = 8. (B) Lower Fbxo10 transcript level associated with the resistant Mes5a allele is observed in unsorted and sorted thymocytes, except for double negative (CD4+CD8+) thymocytes. Graphed are average ± SEM Fbxo10 transcript levels (normalized to Gapdh) in unsorted and sorted thymocytes, scaled to the average Fbxo10 transcript level for the unsorted thymocytes of the susceptible congenic control line. Sample sizes for the susceptible congenic control line and Mes5a congenic resistant line were, respectively: unsorted, n = 16 and n = 9; CD4+CD8−, n = 4 and n = 4; CD4−CD8+, n = 17 and n = 19; CD4+CD8+, n = 15 and n = 18; CD8+CD4−, n = 18 and n = 17. (C) Lower Fbxo10 transcript level associated with the resistant Mes5a allele is observed in splenic T cells, but not in other splenocytes. Graphed are average ± SEM Fbxo10 transcript levels (normalized to Gapdh) in unsorted and sorted splenocytes, scaled to the Fbxo10 transcript level for the unsorted splenocytes of the susceptible congenic control line. Sample sizes for the susceptible congenic control line and Mes5a resistant congenic line were, respectively: unsorted, n = 16 and n = 12; B cells, n = 9 and n = 8; T cells, n = 16 and n = 13; non-B-/non-T cells, n = 3 and n = 5. (D) Lower Fbxo10 transcript level associated with the resistant Mes5a allele is observed in cultured unstimulated and conA-stimulated T cells. Graphed are average ± SEM Fbxo10 transcript levels (normalized to ActB) in cultured T cells, unstimulated (−) and stimulated (+) with conA, scaled to the average Fbxo10 transcript level of the unstimulated sample of the susceptible congenic control line. Sample sizes for the susceptible congenic control line and Mes5a resistant congenic line were, respectively: unstimulated, n = 11 and n = 10; conA-stimulated, n = 11 and n = 12. (E) Tomm5 transcript levels in the immune system are not associated with the resistant Mes5a allele. Graphed are average ± SEM Tomm5 transcript levels (normalized to ActB) in spleen, thymus and primary T cells, scaled to the average Tomm5 transcript level of the susceptible congenic control line. Sample sizes for the susceptible congenic control line and Mes5a resistant congenic line were, respectively: spleen, n = 7 and n = 7; thymus, n = 8 and n = 8; primary T cells, n = 12 and n = 9. (F) Tomm5 transcript levels in cultured T cells are strongly increased after conA-stimulation. Graphed are average ± SEM Tomm5 transcript levels (normalized to ActB) in cultured T cells, unstimulated and stimulated with conA, scaled to the average Tomm5 transcript level of the unstimulated sample of the susceptible congenic control line. Sample sizes for the susceptible congenic control line and Mes5a resistant congenic line were, respectively: unstimulated, n = 11 and n = 10; conA-stimulated, n = 11 and n = 12. In B-F, significantly different transcript level (P < 0.05) between the susceptible congenic control line (ssuc.; open bars) and the Mes5a resistant congenic line (Mes5a; filled bars) are indicated with an asterisk. In D and F, significantly different Fbxo10 or Tomm5 transcript level (P < 0.05) between samples without and with conA stimulation is indicated with two asterisks.
MCS5A/Mcs5a is a human/rat-conserved breast cancer susceptibility locus. In the rat, the resistant Mcs5a allele has previously been shown to consist of two interacting genetic elements that must be located on the same chromosome to elicit mammary carcinoma resistance (5). The carcinoma susceptibility phenotype of Mcs5a is not transferable from donor to recipient animals in a mammary gland transplantation assay, indicating that Mcs5a acts in a non-mammary cell-autonomous fashion (15). We have also demonstrated that carcinoma development is influenced by the immune system, as Mcs5a resistant congenic rats reconstituted with the susceptible congenic control line’s immune system develop more mammary carcinomas as compared with Mcs5a resistant congenic rats reconstituted with their own immune system. Finally, we have observed that T-cell homeostasis and functions are under control of the Mcs5a locus (15).

In this study, we begin to translate the association of genetic variants with the complex trait breast cancer susceptibility into molecular mechanisms. As the genetic variants underlying the MCS5A/Mcs5a locus are found in non-protein-coding sequences, we sought to investigate the gene regulatory function of MCS5A/Mcs5a. We showed previously that the expression level of genes located within 1 Mb surrounding the locus, including Fbxo10, Frmpd1 and Tomm5 that are partially located within Mcs5a, were not differentially expressed in the

Figure 4. Fbxo10/FBXO10 TSS analysis by the 5’ RLM-RACE assay. (A) Schematic representation of three putative Fbxo10/FBXO10 transcripts annotated in the UCSC genome browser, indicated relative to the position of the Mcs5a1/MCS5A1 locus (in black). The location of the Fbxo10/FBXO10 specific primers in the first coding exon is indicated (P). The 5’ RLM-RACE revealed a cluster of 14 rat Fbxo10 TSSs within the Mcs5a1 CpG island (indicated in light gray). A cluster of three human FBXO10 TSSs in the orthologous MCS5A1 CpG island (indicated in light gray) was identified. The position of the first non-coding exon of Fbxo10/FBXO10 within the CpG island is indicated in dark gray. The human TSS cluster was found to be located at 150-bp distance from breast cancer risk-associated SNP rs6476643. The transcripts not identified in rat or human immune tissue are indicated with a X. (B) Sequence information of the rat and human Mcs5a1/MCS5A1 CpG island, first non-coding Fbxo10/FBXO10 exon (highlighted in gray) and TSS identified (in bold). In the human sequence, the breast cancer risk-associated SNP rs6476643 (T/G) is indicated.
Figure 5. Transcriptional activity analysis of the human breast cancer risk-associated susceptible and resistant alleles in Luciferase reporter assays. (A) Schematic representation of the position of the genomic fragments derived from the MCS5A1 and MCS5A2 loci combined into the Luciferase constructs. The MCS5A1 and MCS5A2 loci are shown as black lines. The light gray bars within the black lines represent the CpG islands located in the locus. The three genes FBXO10, FRMPD1 and TOMM5 are shown in dark gray. The breast cancer risk-associated polymorphisms are represented as vertical gray lines. The fragments subcloned into the reporter constructs are indicated as horizontal light gray bars. The susceptible alleles of the MCS5A2 polymorphisms were combined with the susceptible allele of the MCS5A1 SNP rs6476643 (SS constructs 1–10). Similarly, the resistant alleles were combined (RR constructs 1–10). (B) Map of the FBXO10 TSS-Luciferase reporter construct. A MCS5A1 fragment containing the FBXO10 TSSs and the risk-associated SNP rs6476643 was inserted upstream of the Luciferase reporter gene (Luc+) in reverse genomic orientation. Two versions of the construct were created, namely having the susceptible (S; T allele) or resistant (R; G allele) of SNP rs6476643. Other features of the construct include the ampicillin resistance gene (AmpR), origin of replication derived from filamentous phage (f1 ori), origin of replication in E. coli (ori), a synthetic poly(A) signal/transcriptional pause site for background reduction [synthetic poly(A)] and a cloning site downstream of Luc+. (C) Box plot of the relative Luciferase activity of the R and S constructs. n = 30 transient transfection assays. (D) Average ± SEM relative Luciferase activity of constructs SS 1–10 and RR 1–10. On the x-axis, the genomic distance of the MCS5A2 polymorphisms to the MCS5A1 SNP rs6476643 is plotted. The data points at genomic distance 0 correspond to the FBXO10 TSS-Luciferase constructs S and R. The measurements indicated with an asterisk are significantly different between SS and RR (P < 0.05). The entire series of RR1–10 is significantly lower as compared with the entire series of SS 1–10 (P < 10^-8). n = 6 or more transient transfection assays per data point.
mammary gland of susceptible congenic control and Mcs5a resistant congenic rats. Two of these genes, however, were found to be differentially expressed in the immune system, namely Fbxo10 in the thymus, and Frmpd1 in the spleen (5). Here, we have shown that the genetic interaction that is necessary for the mammary carcinoma resistance phenotype, is also essential to downregulate Fbxo10 transcript levels in the thymus. The transcript levels of Frmpd1 and Tomm5 were not differentially associated with the Mcs5a alleles. Lower Fbxo10 transcript levels in the resistant congenic rat line as compared with the susceptible congenic control line were detectable in CD4+CD8– thymocytes, CD8+CD4– thymocytes, CD4+CD8+ thymocytes and CD3+ primary, cultured unstimulated and cultured activated T cells from the spleen. Analysis of some aspects of the gene regulatory mechanism mediated by the Mcs5a/MCS5A locus revealed striking similarities between rats and humans. First, the TSSs of the Fbxo10/FBXO10 transcripts were found to be located in the CpG island in Mcs5a/MCS5A in both rat and human thymus and spleen RNA samples. The human Fbxo10 TSS cluster was found to be located 150-bp away from a breast cancer risk-associated SNP (rs6476643). Second, both the rat and human Mcs5a/MCS5A locus display a similar profile of chromatin looping and CTCF/cohesin binding. Third, functional assessment of the transcriptional regulatory properties of the resistant (R) and susceptible (S) allele of rs6476643 in combination with the R and S allele of each of the 15 MCS5A2 risk-associated SNPs, revealed an overall significantly lower transcriptional activity for the RR combinations compared to the SS combinations, thus mimicking the resistant Mcs5a1–Mcs5a2 interaction controlling Fbxo10 transcript levels and mammary carcinoma resistance in the rat. Interestingly, the R allele of multiple MCS5A2 SNPs combined with the R allele of rs6476643 can independently lower transcriptional activity compared to the combined SS alleles, perhaps underlying a haplotype effect between the MCS5A1 and MCS5A2 breast cancer risk-associated polymorphisms, which are separated by >60 kb in the genome. Physical interactions between the fragments containing the MCS5A1 and MCS5A2 risk-associated polymorphisms, however, were not directly observed, likely due to limited sensitivity of the 3C assay and the transient, stochastic and/or structurally diffuse nature of the putative interactions.

Mcs5a/MCS5A harbors a higher order chromatin structure, which spatially isolates the Tomm5/TOMM5 gene and its flanking regulatory region by locating them within the loop (Figure 6). The chromatin loop also functionally isolates Tomm5 as its transcript level was found to be strongly upregulated in T-cell activation, whereas Fbxo10 or Frmpd1 transcript levels were not. The higher order chromatin structure is thought to be an insulator loop. There appear to be three looping fragments involved that are all bound by CTCF and cohesin. CTCF has been recognized as a DNA-binding protein with diverse functions (23), for example insulator loop formation (24), and enhancer–promoter looping, e.g. in the β-globin locus (25). CTCF has also been associated with boundary/barrier elements, which demarcate separately regulated chromatin domains (26). The transcription of a growing number of genes has been found to be dependent on CTCF- and/or cohesin-binding and -looping. For example, gene expression in the developmentally regulated HoxA cluster is under control of CTCF-mediated heterochromatin partitioning, with involvement of Oct4 and cohesin loading at the CTCF site in undifferentiated embryonic stem cells and differentiated neuronal progenitor cells, respectively (27). Another example of CTCF-mediated looping and cooperation of a transcription factor (T-bet) in cell type specific gene transcription is the Th1 cell-specific expression of Ifng (28). Ablation of

Figure 6. A model of the MCS5A locus in a folded configuration. The first (non-coding, 5'-UTR) and second (coding, ATG-containing) exons of the Fbxo10 transcript are displayed in orange. The Tomm5 transcript is indicated in dark blue. The first (non-coding, 5'-UTR) exon of the Frmpd1 transcript is shown in light blue. The CpG islands associated with their promoters are indicated in dark green. The correlated polymorphisms that associate with breast cancer risk are depicted as purple bars. The looping fragments are shown in light green and are shown to be bound by CTCF. The looped structure involves three elements that may loop simultaneously (as depicted) or loop in specific combinations in a single nucleus. The polymorphisms associated with breast cancer risk in MCS5A1 and MCS5A2 are located at both sides of the looped structure and are in closer proximity as may be derived from a linear genome view. We hypothesize that the MCS5A1 and MCS5A2 breast cancer risk variants interact to regulate the transcript levels of Fbxo10.
the Mcs5a loop by depletion of CTCF yielded a lower Tomm5 transcript level (whereas the transcript levels of two other genes adjacent to Mcs5a were unaffected), possibly through misregulation by the adjacent CTCF.
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