



FANCM c.5101C>T mutation associates with breast cancer survival and treatment outcome

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Breast cancer (BC) is a heterogeneous disease, and different tumor characteristics and genetic variation may affect the clinical outcome. The *FANCM* c.5101C>T nonsense mutation in the Finnish population associates with increased risk of breast cancer, especially for triple-negative breast cancer patients. To investigate the association of the mutation with disease prognosis, we studied tumor phenotype, treatment outcome, and patient survival in 3,933 invasive breast cancer patients, including 101 *FANCM* c.5101C>T mutation carriers and 3,832 non-carriers. We also examined association of the mutation with nuclear immunohistochemical staining of DNA repair markers in 1,240 breast tumors. The *FANCM* c.5101C>T mutation associated with poor 10-year breast cancer-specific survival (hazard ratio (HR)=1.66, 95% confidence interval (CI) 1.09–2.52, $p=0.018$), with a more pronounced survival effect among familial cases (HR = 2.93, 95% CI 1.5–5.76, $p=1.80 \times 10^{-3}$). Poor disease outcome of the carriers was also found among the estrogen receptor (ER) positive subgroup of patients (HR = 1.8, 95% CI 1.09–2.98, $p=0.021$). Reduced survival was seen especially among patients who had not received radiotherapy (HR = 3.43, 95% CI 1.6–7.34, $p=1.50 \times 10^{-3}$) but not among radiotherapy treated patients (HR = 1.35, 95% CI 0.82–2.23, $p=0.237$). Significant interaction was found between the mutation and radiotherapy ($p=0.040$). Immunohistochemical analyses show that c.5101C>T carriers have reduced PAR-activity. Our results suggest that *FANCM* c.5101C>T nonsense mutation carriers have a reduced breast cancer survival but postoperative radiotherapy may diminish this survival disadvantage.

Key words: FANCM, breast cancer, survival, DNA repair, radiotherapy

Abbreviations: BC: breast cancer; BER: base excision repair; CI: confidence interval; CISH: chromogenic in situ hybridization; DSB: double strand break; ER: estrogen receptor; FA: Fanconi anemia; HR: hazard ratio; HRM: high resolution melt; IHC: immunohistochemistry; KBCP: Kuopio breast cancer project; NHEJ: non-homologous end joining; PARP: poly(ADP-ribose) polymerase; PR: progesterone receptor; SSB: single strand break; TNBC: triple-negative breast cancer

Additional Supporting Information may be found in the online version of this article.

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What's new?

Variations in DNA repair genes can predispose individuals to breast cancer, with one example being *FANCM* c.5101C>T, a nonsense mutation in the Fanconi Anemia DNA repair pathway. In previous work, *FANCM* c.5101C>T was associated with increased breast cancer risk in the Finnish population. Here, the mutation is further shown to be associated with adverse breast cancer outcome. Mutation-positive Finnish patients exhibited reduced long-term survival and increased risk of disease recurrence. Survival was worse particularly for patients who were not treated with radiotherapy, indicating that *FANCM* c.5101C>T may interact with radiotherapy to improve disease outcome in mutation carriers.

Breast cancer is the most common cancer among women worldwide, and also the leading cause of female cancer death.¹ Most breast cancer cases are sporadic, but around 15% have familial background. Hereditary predisposition to breast cancer is caused by variation in multiple genes commonly involved in DNA repair, especially with homologous recombination repair pathway.² Recently, we identified a new breast cancer allele in the Finnish population in the *FANCM* gene, that functions in the Fanconi Anemia (FA) DNA repair pathway. The *FANCM* c.5101C>T (p.Q1701X, rs147021911) nonsense mutation increased the risk of breast cancer over twofold, and 3.5-fold increased frequency was seen among the triple-negative breast cancer (TNBC) cases.³

Predisposing mutations may associate with specific breast cancer phenotype or subgroup, as well as with patient prognosis and treatment outcome. *CHEK2* and *PALB2* truncating mutations, as well as *FANCM* c.5101C>T mutation, confer moderate risk for breast cancer, with a higher risk among patients with family history of breast cancer.^{3–7}

CHEK2 c.1100delC and *PALB2* c.1592delT mutations are associated also with an increased risk of breast cancer death or second breast cancer. Among patients with ER positive breast cancer, *CHEK2* c.1100delC heterozygosity is associated with 1.6-fold risk of breast cancer specific death and 3.5-fold risk of a second breast cancer.^{8,9} A significant proportion of *PALB2* tumors are triple-negative and the *PALB2* mutation carriers have about 2-fold increased risk of breast cancer death, independently of the triple-negative status.^{7,10}

Here, we studied tumor characteristics, patient survival, and treatment outcome associated with the *FANCM* c.5101C>T mutation among 3,933 breast cancer patients in four breast cancer patient series from Finland. In addition, we examined the nuclear immunohistochemical staining of DNA repair markers in the mutation carrier and non-carrier tumors from 1,240 invasive breast cancer cases.

Material and Methods**Subjects**

Helsinki breast cancer series. The unselected breast cancer patient samples from Helsinki were collected at Helsinki University Central Hospital. From this cohort, 884 samples, including 79% of all consecutive, newly diagnosed breast cancer cases during the collection periods were collected at Department of Oncology in 1997–1998 and 2000.^{11,12} In

addition, 986 samples, including 87% of all consecutive, newly diagnosed breast cancer cases were collected at Department of Surgery in 2001–2004.¹³ Of these series, 397 cases had family history of breast cancer.

Additional familial breast cancer series was collected at Helsinki University Central Hospital Departments of Oncology and Clinical Genetics.^{13,14} When combining the unselected and the additional familial samples, 524 patients had strong family history with at least three breast or ovarian cancers among first or second degree relatives (including the proband) and 568 patients had at least one first degree relative affected with breast or ovarian cancer. All the patients with strong family history were tested negative for *BRCA1/2* mutations and the patients with one affected relative were tested negative for Finnish *BRCA1/2* founder mutations as previously described.^{5,15,16} Only invasive cases were included in the analyses ($N = 2,337$).

All samples are genomic DNA isolated from peripheral blood. The patient genealogies were confirmed with population registries or hospital records and cancer diagnoses through the hospital records and the Finnish Cancer Registry. ER and progesterone hormone receptor (PR) status (positive when >10% of cells were stained) and tumor histology information were collected from pathology reports, HER2-status is based on immunohistochemistry and gene amplification as described earlier.^{17–19} Information on breast cancer death was obtained from the Finnish Cancer Registry.

Tampere breast cancer series. The unselected breast cancer patient samples from Tampere area were collected in 1997–1999 and additional 336 incident cases in 1996–2004 at Tampere University Hospital as previously described.^{11,13} Only invasive cases were included in the analysis ($N = 650$). All samples are genomic DNA isolated from peripheral blood. ER and PR hormone receptor status (positive when >10% of cells were stained), HER2-status, and other clinicopathological information was obtained from patient and pathology reports and information on breast cancer death from the Finnish Cancer Registry.

Oulu breast cancer series. The unselected breast cancer patient samples from Northern Finland were collected at the Oulu University Hospital between the years 2000 and 2007. Only invasive cases were included in the analysis ($N = 516$). All samples are genomic DNA isolated from peripheral

blood. HER2-status was studied by means of immunohistochemistry (positivity defined as weak, moderate or strong levels of staining and negativity completely negative staining) and chromogenic *in situ* hybridization (CISH). ER and PR hormone receptor status (positive when >10% of cells were stained) and tumor histology information was collected from the pathology reports as described earlier.^{20,21} Information on breast cancer death was obtained from the Oulu University Hospital.

Kuopio breast cancer series. For this study a sample set was used from The Kuopio Breast Cancer Project (KBCP), a prospective population-based case-control study conducted in 1990–1995. Women entering Kuopio University Hospital due to breast symptoms were invited to take part in the study at their first visit to the hospital. Altogether 516 women out of 1,919 were eventually diagnosed to have breast cancer. Hospital registries were used to collect information concerning clinicopathological features of the breast cancer, surgical and oncological treatments, and follow-up.^{22,23} ER and PR hormone receptors were classified as positive if the percentage of positive cells with nuclear staining was $\geq 10\%$. HER2 status assessment was conducted by immunohistochemistry (IHC). Samples with IHC score 2+ or 3+ were classified as HER2 positive (HER2+). Altogether, 430 female patients with invasive breast cancer were included in the survival analysis. All samples are genomic DNA isolated from peripheral blood.

This study was performed with informed consent from the patients and permission from the ethics committees of Helsinki University Hospital, Oulu University Hospital, Tampere University Hospital, University of Eastern Finland, and Kuopio University Hospital Board on Research Ethics.

Genotyping

FANCM c.5101C>T genotyping for the Helsinki and Tampere sample sets was performed with Sequenom MassARRAY system as previously described³ and for Oulu and Kuopio sample sets by using PCR-based high resolution melt (HRM)—analysis and Sanger sequencing. The HRM PCR reactions were performed in 96 well plates using Type-it HRM PCR Kit (Qiagen, Hilden, Germany) and CFX96 Real-Time PCR Detection System (CFX96, Bio-Rad, Hercules, CA). Primers used for the genotyping and sequencing *FANCM* c. 5101C>T mutation for Oulu and Kuopio cohorts were: F: 5'TCAAGTGAGGAGGAGAACAATG3', R: 5'TCAGCGATGTCTGTTTGCTC3'.

Statistical Analyses

All four datasets including altogether 3,933 invasive breast cancer patients from Helsinki, Tampere, Oulu, and Kuopio areas of Finland, were pooled for statistical analyses. All statistical analyses were performed using the R version 3.0.2 statistical software (<http://www.r-project.org/>). Kaplan–Meier survival curves and uni and multivariate Cox's proportional hazard models were used to estimate the hazard ratios and

confidence intervals for survival and forest plots were drawn for visualization. All analyses were stratified by the study. The primary end point of the survival analyses was breast cancer death with 10-year follow-up time. In addition, 5-year survival analysis with local recurrence as an endpoint was used for survival analyses in the radiotherapy-based subgroups in the Helsinki data set ($N = 2,337$), where the information about local recurrence of the disease was available. Time-to-event was calculated from the date of the patient diagnosis and to account for the latency between diagnosis and recruitment into the study, all follow-up times were left-truncated. Cases with missing data were excluded from the analyses.

The multivariate analyses included the common clinically relevant factors (ER, grade, tumor size, nodal status) and/or cancer treatments (radiotherapy, endocrine therapy, and chemotherapy) as categorical co-variables and were stratified by the study; inclusion of the study as a categorical co-variant did not affect the result. In addition, the *FANCM* c.5101C>T genotype from the pooled data set was fitted into two Cox's proportional hazard models in order to test the interaction between the mutation and radiotherapy treatment. One model included the treatment and *FANCM* c.5101C>T genotype as individual covariates and the other included an interaction term between these two. Two-way anova was used as a likelihood-ratio test to compare the two models.^{24,25}

The *p*-values for comparisons of histopathological features of mutation carriers and non-carriers were calculated with Pearson's chi squared test or Fisher exact test (for $n \leq 5$). Logistic regression was used for histopathological features with more than two categories. *p*-values <0.05 were considered statistically significant.

To test whether *FANCM* mutation status correlates with immunohistochemical expression of markers involved in DNA damage response and repair, we analyzed a number of markers that have been stained and scored as described in our previous studies: BRCA1, FANCD2, RAD51, XPF, PAR²⁶; ATM,¹⁸ gamma-HA2X,²⁷ and TP53.²⁸ For the continuously scored markers (BRCA1, FANCD2, RAD51, XPF, and PAR; % positive nuclei and staining intensity score as determined by automated analysis), association with *FANCM* mutation status was tested using a Kruskal–Wallis test. All other markers used categorical scoring and a χ^2 test was employed as the test for association. Further information is available in Supporting Information Appendix.

Results

All survival analysis results are based on the 3,933 invasive breast cancer cases in the pooled data set with 581 breast cancer deaths, except the survival analysis among radiotherapy-based subgroups with local recurrence as an endpoint is based on the Helsinki data set with 2,337 invasive samples, including 344 breast cancer deaths. The pooled data set includes 101 *FANCM* c.5101C>T mutation carriers and

3,832 non-carriers, Helsinki data set includes 61 mutation carriers and 2,276 non-carriers. The tumor characteristics of the patients and detailed description of all the datasets used are presented in Table 1.

Histopathological Features of the *FANCM* C.5101C>T Positive Tumors

The association of the *FANCM* c.5101C>T mutation with histopathological features of the tumors was studied in the pooled data among all cases and separately among ER positive cases (Table 2). The mutation did not associate with any common clinical feature, however the breast tumors from the c.5101C>T mutation carriers were more often of triple negative phenotype ($p = 0.060$, compared with tumors from non-carriers).

FANCM C.5101C>T Mutation Associates with Breast Cancer Survival

To evaluate the association of the *FANCM* c.5101C>T mutation with the disease outcome, we examined 10-year breast cancer specific survival by Cox's univariate proportional hazard analysis in 3,933 invasive breast cancer patients from Helsinki, Tampere, Oulu, and Kuopio data sets. The mutation was associated with poor breast cancer-specific survival in the pooled data set stratified for study (HR = 1.66, 95% CI 1.09–2.52, $p = 0.018$). Absolute uncorrected survival rates are illustrated in Figure 1a. However, in the multivariate survival analysis including the common clinical features (ER, grade, tumor size, nodal status) and the conventional cancer treatments (radiotherapy, chemotherapy, endocrine treatment) the mutation was not significantly and independently prognostic in the pooled data set (HR = 1.44, 95% CI 0.91–2.26, $p = 0.133$) (Supporting Information Table 1).

As the mutation associates with triple-negative phenotype with poor survival as such, we analyzed the survival specifically also among ER positive cases. The mutation associated with reduced survival also in the ER-positive group of patients in the pooled data set stratified for study (HR = 1.8, 95% CI 1.09–2.98, $p = 0.021$). Absolute uncorrected survival rates are illustrated in Figure 1b. Furthermore, as the *FANCM* c.5101C>T mutation associates with familial breast cancer risk, we performed the survival analysis for the invasive familial cases ($N = 1,006$) among the Helsinki dataset in which familial status was available for the samples. The breast cancer specific survival was worse for mutation carriers among patients with family history of the disease (HR = 2.93, 95% CI 1.5–5.76, $p = 1.80 \times 10^{-3}$; Fig. 1c).

Survival in Subgroups Defined by Tumor Phenotype and Treatment

To examine the survival effect of the *FANCM* c.5101C>T mutation in more detail, we performed univariate Cox's proportional hazard analysis (endpoint: breast cancer death in 10 years) in subgroups based on the tumor phenotype (ER, PR, TN, nodal status, tumor size, grade) among the pooled

data set ($N = 3,933$). In addition, we performed univariate Cox's proportional hazard analysis by the conventional cancer treatment options (endocrine treatment, radiotherapy, and/or chemotherapy) to examine the treatment outcome of the *FANCM* c.5101C>T mutation carriers. Forest plot was drawn for visualizing hazard ratios and confidence intervals (Fig. 2). As the worse survival was also seen among the ER-positive patients, we performed similar subgroup analyses (PR, TN, nodal status, tumor size, grade, and the anticancer treatments) among ER-positive patients ($N = 3,013$) (Supporting Information Fig. 1). Heterogeneity in the survival effect was seen for the c.5101C>T mutation carriers related to radiotherapy treatment, with significantly reduced survival especially among patients who had not received radiotherapy (HR = 3.43, 95% CI 1.6–7.34, $p = 1.50 \times 10^{-3}$) but not among radiotherapy treated patients (HR = 1.35, 95% CI 0.82–2.23, $p = 0.237$).

To further examine the radiotherapy outcome among the c.5101C>T carriers, we performed survival analysis with local recurrence (within 5 years) as an endpoint in the Helsinki data set where the recurrence information was available ($N = 2,337$). Increased risk for local recurrence was observed for mutation carriers who had not received radiotherapy (HR = 6.19, 95% CI 1.46–26.2, $p = 0.013$, Supporting Information Table 2) but not among radiotherapy treated patients (HR = 0.98, 95% CI = 0.24–4.00, $p = 0.979$). In the multivariate model, the *FANCM* c.5101C>T mutation is only borderline significant ($p = 0.086$), however the hazard ratios remain consistent.

Next, we tested interaction between *FANCM* c.5101C>T genotype and radiotherapy treatment with Cox's proportional hazard model stratified with study among pooled data set, including 2,996 patients who had received radiotherapy and 864 who had not (Table 3A). A significant interaction was seen between the mutation and radiotherapy treatment ($p = 0.032$), with a protective hazard ratio (HR = 0.37, 95% CI 0.15–0.92). A likelihood-ratio test comparing models with interaction term and model with independent covariates displayed an interactive effect between the covariates ($p_{(\text{interaction})} = 0.040$). These results suggest that *FANCM*-mutation positive breast cancer patients may benefit from radiotherapy more than non-carriers, an issue that should be further investigated to clarify the absolute benefits from radiotherapy to such patients.

We further studied the survival interaction of *FANCM* mutation with radiotherapy using similar interaction model with local recurrence (within 5 years) as an endpoint in the Helsinki data set ($N = 2,069$) (Table 3B). Due to the smaller sample size and thus loss of statistical power, the significance of the interactive effect is not apparent (likelihood-ratio test p values 0.090). However, even more pronounced protective hazard ratio was seen for *FANCM* c.5101C>T mutation and radiotherapy interaction (HR = 0.16), compared to significantly increased hazard ratio for mutation alone (HR = 5.96).

Table 1. Description of the patient data sets used in this study

	Helsinki	Tampere	Oulu	Kuopio
No. of cases	2,337	650	516	430
No. of mutation carriers	61 (2.6%)	26 (4%)	5 (1%)	9 (2%)
Vital status				
Alive	1,482 (64%)	448 (69%)	362 (70%)	176 (41%)
Deceased: all-cause	511 (21%)	118 (18%)	94 (18%)	161 (37%)
Deceased: breast cancer	344 (15%)	84 (13%)	60 (12%)	93 (22%)
Follow-up mean \pmSD (years)	8.16 \pm 2.4	7.44 \pm 2.13	5.17 \pm 2.92	7.78 \pm 3.08
Age at diagnosis, mean [range]	56.3 [21–95]	58.9 [30–88]	57.4 [28–92]	58.1 [23–91]
Estrogen receptor				
Negative	430 (18%)	128 (20%)	96 (19%)	101 (23%)
Positive	1,803 (77%)	508 (78%)	385 (75%)	300 (70%)
Missing data	104 (5%)	14 (2%)	35 (7%)	29 (7%)
Grade				
1	580 (25%)	197 (30%)	76 (15%)	115 (27%)
2	980 (42%)	226 (35%)	212 (41%)	196 (46%)
3	651 (28%)	133 (20%)	177 (34%)	115 (27%)
Missing data	126 (5%)	94 (14%)	51 (10%)	4 (1%)
T/tumor size category				
1	1,409 (60%)	401 (62%)	238 (46%)	229 (53%)
2	743 (32%)	213 (33%)	226 (44%)	161 (37%)
3	69 (3%)	24 (4%)	15 (3%)	23 (5%)
4	82 (4%)	–	–	17 (4%)
Missing data	34 (1%)	12 (2%)	37 (7%)	–
N (nodal metastasis)				
Negative	1,263 (54%)	390 (69%)	265 (51%)	251 (58%)
Positive	1,036 (44%)	260 (40%)	216 (42%)	171 (40%)
Missing data	38 (2%)	–	35 (7%)	8 (2%)
M (distant metastasis)				
Negative	2,253 (96.5%)	630 (97%)	492 (95%)	419 (97%)
Positive	73 (3%)	12 (2%)	24 (5%)	11 (3%)
Missing data	11 (0.5%)	8 (1%)	–	–
Histological type				
Ductal	1,597 (68%)	537 (83%)	371 (71%)	281 (65%)
Lobular	470 (20%)	86 (13%)	78 (15%)	73 (17%)
Medullar	29 (1%)	–	2 (1%)	8 (2%)
Other	240 (10%)	18 (3%)	30 (6%)	68 (16%)
NA	1	9 (1%)	35 (7%)	–
Radiotherapy				
Yes	1,829 (78%)	493 (76%)	423 (82%)	251 (58%)
No	443 (19%)	155 (24%)	87 (17%)	179 (42%)
Missing data	65 (3%)	2	6 (1%)	–
Chemotherapy				
Yes	870 (37%)	131 (20%)	215 (42%)	83 (19%)
No	1,405 (60%)	511 (79%)	297 (58%)	347 (81%)
Missing data	62 (3%)	8 (1%)	4 (1%)	–
Endocrine therapy				
Yes	1,055 (45%)	204 (32%)	243 (47%)	105 (24%)
No	1,207 (52%)	444 (68%)	268 (52%)	325 (76%)
Missing data	65 (3%)	2	5 (1%)	–

Table 2. Histopathological features of *FANCM* c.5101C>T-mutation carriers and wild type tumors

Category	FANCM c.5101C>T	%	FANCM wt	%	<i>p</i>	Model
All breast cancer cases						
Grade					0.263	Logistic regression
1	25	26.00%	943	26.00%		
2	36	37.00%	1,578	44.00%		
3	36	37.00%	1,040	30.00%		
T					0.255	Logistic regression
1	53	53.00%	2,224	59.00%		
2	41	41.00%	1,302	34.50%		
3	2	2.00%	129	3.00%		
4	4	4.00%	95	2.50%		
N					0.380	Pearson chisq.
neg	52	52.00%	2,117	56.20%		
pos	48	48.00%	1,653	43.80%		
M					0.770	Fisher
neg	99	99.00%	3,695	96.90%		
pos	2	1.00%	118	3.10%		
ER					0.432	Pearson chisq.
neg	23	23.00%	726	19.80%		
pos	77	77.00%	2,936	80.20%		
PR					0.380	Pearson chisq.
neg	39	39.00%	1,271	34.80%		
pos	61	61.00%	2,386	65.20%		
Her2					0.167	Pearson chisq.
neg	67	90.50%	2,336	91.50%		
pos	7	9.50%	422	8.50%		
TN					0.060	Pearson chisq.
TN	13	14.00%	297	8.50%		
NOT TN	80	86.00%	3,215	91.50%		
Morphology					0.366	Logistic regression
Ductal	78	77.00%	2,708	71.50%		
Lobular	14	14.00%	693	18.30%		
Medullar	1	1.00%	38	1.00%		
Other	8	8.00%	348	9.20%		
ER-positive breast cancer cases						
Grade					0.813	Logistic regression
1	25	33.50%	874	31.50%		
2	33	44.00%	1,379	49.50%		
3	17	22.50%	524	19.00%		
T					0.279	Logistic regression
1	45	57.00%	1,827	63.00%		
2	27	35.00%	934	32.00%		
3	1	3.00%	87	3.00%		
4	4	5.00%	67	2.00%		
N					0.500	Pearson chisq.
neg	40	52.50%	1,265	43.50%		
pos	36	47.50%	1,644	46.50%		

Table 2. Histopathological features of *FANCM* c.5101C>T-mutation carriers and wild type tumors (Continued)

Category	FANCM c.5101C>T	%	FANCM wt	%	<i>p</i>	Model
M					0.775	Fisher
neg	76	98.70%	2,841	98.00%		
pos	1	1.30%	65	2.00%		
PR					0.754	Pearson chisq.
neg	17	22.00%	604	20.00%		
pos	60	88.00%	2,326	80.00%		
Her2					1	Fisher
neg	52	91.00%	1,943	90.00%		
pos	5	9.00%	219	10.00%		
Morphology					0.587	Logistic regression
Ductal	56	73.00%	2,007	68.00%		
Lobular	14	18.00%	635	22.00%		
Medullar	0	0.00%	4	0.00%		
Other	7	9.00%	286	10.00%		

Abbreviations: T: tumor size class; M: distant metastasis; ER: estrogen receptor; PR: progesterone receptor

Immunohistochemical Analyses

In the association analysis between *FANCM* c.5101C>T mutation status and DNA repair related immunohistochemical markers, a statistically significant association was detected between nuclear poly-ADP-ribose (PAR; a measurement of PARP activity) staining and mutated *FANCM*. PAR staining was reduced in *FANCM* c.5101C>T mutation carrier tumors, both in terms of the proportion of positively stained tumor nuclei ($p = 0.016$, Kruskal-Wallis test) and staining intensity ($p = 0.011$, Kruskal-Wallis test) (Supporting Information Fig. 2). No other immunohistochemical markers were associated with mutated *FANCM* (Supporting Information Table 3).

Discussion

This study evaluated the survival association, tumor characteristics, and treatment outcome for Finnish breast cancer patients carrying the *FANCM* c.5101C>T mutation. We detected an association between the *FANCM* c.5101C>T mutation and adverse breast cancer outcome (HR = 1.66, 95% CI 1.09 – 2.52, $p = 0.018$, $N = 3,832$ [non-carriers], $N = 101$ [mutation carriers]). The breast cancer specific survival was worse among familial cases (HR = 2.93, 95% CI 1.5–5.76, $p = 1.80 \times 10^{-3}$, $N = 981$ [non-carriers], $N = 25$ [mutation carriers]).

When examining the tumors of the *FANCM* c.5101C>T mutation carriers, a borderline significant association of the mutation was seen with triple-negative tumors ($p = 0.060$, compared with tumors from non-carriers). This is in line with the previous risk analysis, in which the *FANCM* c.5101C>T mutation was found to be associated with 3.6-

fold increased risk for triple-negative subtype of breast cancer.³ This type of breast cancer is generally aggressive with poor prognosis and no effective therapies available.²⁹ However, our survival analysis indicates that the poor prognosis associated with *FANCM* c.5101C>T mutation is not only a result of the higher incidence of the triple-negative tumors, as the mutation also associates with worse survival among the ER-positive subgroup of patients. Yet in the multivariate survival analysis including conventional prognostic markers and treatments, the *FANCM* c.5101C>T mutation was not independently prognostic (HR = 1.44, 95% CI 0.91–2.26, $p = 0.133$).

The comprehensive survival analyses revealed an association with *FANCM* c.5101C>T mutation and radiotherapy outcome. Interaction analyses with a hazard ratio of 0.37 (95% CI 0.15–0.95, $p = 0.032$) for the mutation:radiotherapy interaction compared to the HR of 3.72 for the mutation alone (95% CI 1.74–7.95, $p = 7.00 \times 10^{-4}$) in the interaction model indicate that the mutation carriers may benefit from radiotherapy. To this end, we performed the interaction analyses also with local recurrence in five years as an endpoint, as radiotherapy is commonly used to prevent such events. While this interaction model is not statistically significant in the smaller sample set, the more pronounced protective hazard ratio of 0.16 for the radiotherapy and *FANCM* c.5101C>T interaction further supports our observations that carrying the *FANCM* c. 5101C>T mutation increases the risk for local recurrence and subsequently also death from breast cancer, however the mutation carriers seem to benefit from postoperative radiotherapy. From the pathobiological point of view, we propose that the increased risk of local recurrence and death may reflect enhanced genomic

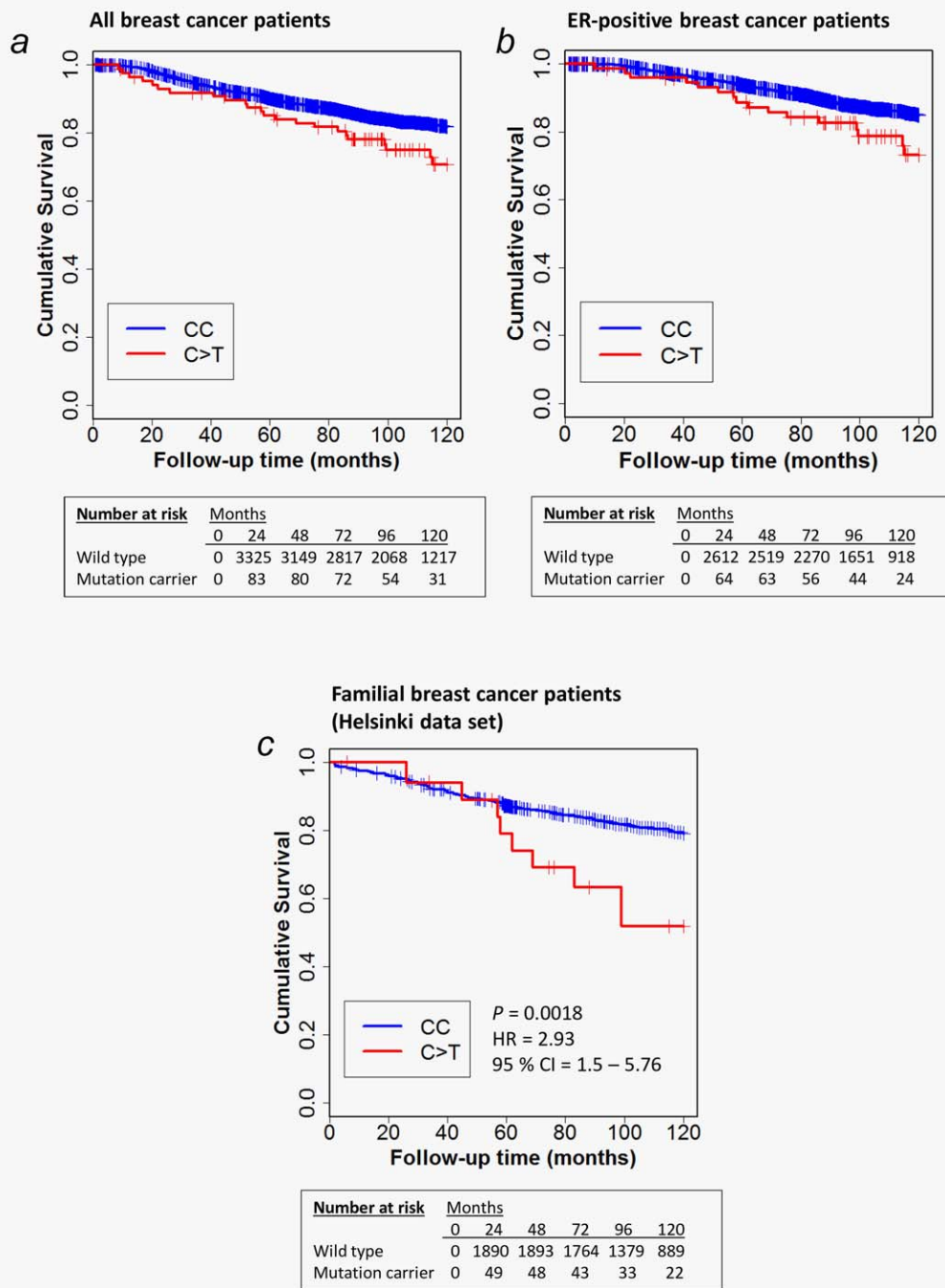


Figure 1. Kaplan–Meier plots of cumulative survival for breast cancer death in 10 years. Absolute uncorrected survival rates are presented among the pooled data set ($HR = 1.62$, $95\% CI 1.07–2.46$, $p_{Cox's\ regression} = 0.023$; (a) and among ER-positive patients ($HR = 1.8$, $95\% CI 1.09–2.98$, $p_{Cox's\ regression} = 0.021$; (b) Results for survival analysis among familial cases (C) from Helsinki.

instability and hence aggressiveness due to impaired DNA repair in the tumors with the *FANCM* c. 5101C>T mutation. On the positive side, such enhanced genetic instability and suboptimal repair capacity seem to represent a specific vulnerability of such tumors, manifest particularly after an

extra burden of difficult-to repair DNA damage caused by ionizing radiation treatment. Overall, these results are especially interesting, as markers associated with radiotherapy treatment outcome for cancer patients have not been previously described. However, further studies in larger datasets

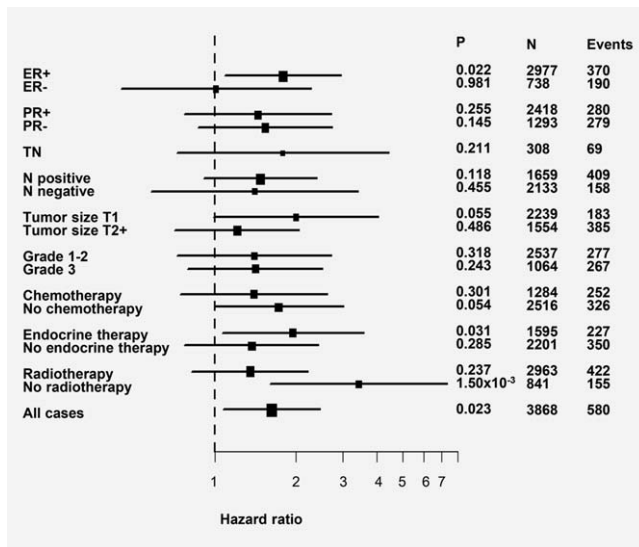


Figure 2. Forest plot of hazard ratios and their confidence intervals for the *FANCM* c.5101 C > T mutation in the pooled data set and in different subgroups including the clinical factors and conventional cancer treatments. The Cox proportional hazard model was used for 10-year breast-cancer specific survival. Horizontal lines represent 95% confidence intervals. ER = estrogen receptor, PR = progesterone receptor, TN = triple negative, N = nodal metastasis status. All cases = all cases after samples with missing data are excluded.

are needed to validate the radiotherapy outcome for *FANCM* mutation carriers.

FANCM is a multifunctional protein, acting as an anchor protein for both Fanconi Anemia and Bloom syndrome complexes, two molecular pathways that functionally overlap in these genetic disorders.³⁰⁻³² As a part of the FA pathway, *FANCM* operates in the interstrand crosslink repair to facilitate various DNA repair processes, such as homologous recombination and non-homologous end-joining (NHEJ) pathway.^{30,33} Inactivation of the FA pathway leads to hypersensitivity to DNA crosslinking agents, and in the absence of *FANCM*, the formation of the FA and Bloom's complexes is unsuccessful and this may explain the tumorigenic characteristics of defective *FANCM* protein.³² Interestingly, in addition to *BRCA*-genes, recent studies link several Fanconi anemia pathway genes also with sensitivity to PARP inhibition, including *PALB2*, *RAD51C*, and *SLX4*,³⁴⁻³⁶ as well as *FANCM*.³⁵ Mutations in *FANCM* were found to cause hypersensitivity to PARP inhibitors, indicating that *FANCM* actually has a role in the cellular defense against PARP inhibition.³⁷ This may reflect the several roles *FANCM* has in cells also outside the Fanconi Anemia pathway, including replisome stability and cell cycle checkpoint activation when DNA repair is needed.³⁸⁻⁴⁰

Taking the DNA repair functions of *FANCM* in consideration, we examined nuclear immunohistochemical staining

Table 3. A) Cox's proportional hazard model to test the interaction between radiotherapy treatment and *FANCM* c.5101C > T mutation with breast cancer death as an endpoint; B) Local recurrence as an endpoint

Covariate	HR	p	95% CI	Endpoint
A				
Model 1: no interaction				Breast cancer death (10 yrs)
RS147021911	1.71	0.011	1.13-2.60	
Radiotherapy	0.70	1.0 × 10 ⁻⁴	0.58-0.84	
Model 2: interaction				
RS147021911	3.72	7.00 × 10 ⁻⁴	1.74-7.95	
Radiotherapy	0.72	8.40 × 10 ⁻⁴	0.59-0.87	
RS147021911:Radiotherapy	0.37	0.032	0.15-0.92	
Likelihood ratio test p values		0.040		
B				
Model 1: no interaction				Local recurrence (5 yrs)
RS147021911	1.71	0.298	0.62-4.64	
Radiotherapy	0.48	1.05 × 10 ⁻³	0.31-0.75	
Model 2: interaction				
RS147021911	5.96	1.50 × 10 ⁻³	1.42-25.11	
Radiotherapy	0.52	4.05 × 10 ⁻³	0.33-0.81	
RS147021911:Radiotherapy	0.16	0.080	0.02-1.23	
Likelihood ratio test p values		0.090		

profiles of DNA repair markers of the *FANCM* c.5101C>T mutation carriers. Among eight examined markers, the mutation was associated with low expression of poly (ADP-ribose) marker (PAR), which measures the activity of the PARP enzymes participating in DNA repair processes in cells,⁴¹ indicating that the mutation carriers have decreased PARP-activity. It must be noted that our immunohistochemical method measures the overall poly(ADP-ribosyl)ation levels in tumor nuclei, and is therefore not specific to any particular PARP enzyme or biological process. The best known example of PARylation occurs in response to DNA damage, where the binding and activity of PARP promotes DNA repair through the single-strand break (SSB), double-strand break (DSB), or base excision repair (BER) pathways.⁴² In the case of excessive DNA damage, hyper-PARylation may also be a signal for cell death.⁴³ PARylation has additionally been reported to play a role in mitosis, chromatin remodeling, regulation of transcription, and the organization of genomic regulatory regions via insulator elements.⁴⁴ We can therefore only speculate on the specific functional significance of the *FANCM*-associated reduction in PARylation observed in our breast tumor samples. We did not detect a change in gamma-H2AX staining, suggesting that a major quantitative change in overall DNA damage is not the case here. Since both *FANCM* and PARP are involved in resolving replication stress,^{45–47} it is possible that the *FANCM* c.5101C>T mutation-associated reduction in PAR staining indicates a replication stress sensitive phenotype that would respond strongly to the extreme replication stress caused by radiation therapy. While the causal relationship of *FANCM* with reduced PARylation levels remains unclear, our data may have therapeutic implications.^{27,48} Given the role of *FANCM* in resolving replication stress, the *FANCM*-mutant tumors may be especially sensitive to drugs that further exacerbate the extent of replication stress, such as PARP inhibitors. Based on our present results and the emerging knowledge in the field, we suggest that the subset of *FANCM*-mutant tumors may be particularly vulnerable to PARP inhibitors, used either as a monotherapy or, as our data indicate, combined with radiotherapy. Future preclinical and clinical studies should test the feasibility of these conceptually plausible options.

Conclusions

Our findings indicate that the *FANCM* c.5101C>T mutation in Fanconi Anemia pathway associates with the disease outcome of breast cancer. Based on the large series of Finnish breast cancer patients, we have shown here that the mutation

carriers have worse long-term survival and increased risk for local recurrence, however the survival may be improved with radiotherapy. Further analyses in larger datasets are warranted to clarify the survival effects and functional mechanisms associated with the mutation, especially on the efficacy of radiotherapy. Such studies may eventually help to understand the biological mechanisms affecting tumor progression and further support efforts for creating more targeted treatment combinations and risk estimation.

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Author Contributions

J.I.K., R.F., C.B. and H.N. designed the study and drafted the manuscript.

J.I.K. analyzed and pooled the data.

J.I.K. and A.T. carried out the molecular genetic studies.

J.I.K. performed the statistical analyses with R.F., S.K., M.J. and L.M.P.

J.B. and J.I.B. performed and evaluated some of the immunohistochemical analyses and J.B. contributed to conceptual discussions and manuscript writing.

T.M., K.P., A.M., M.T., V.-M.K., R.W., A.K. and K.A. contributed samples and patient information. All authors read and approved the final manuscript.

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