



Distinct gene mutation profiles among luminal-type and basal-type breast cancer cell lines

Antoinette Hollestelle, Jord H. A. Nagel, Marcel Smid, Suzanne Lam, Fons Elstrodt, Marijke Wasielewski, Ser Sue Ng, Pim J. French, Justine K. Peeters, Marieke J. Rozendaal, et al.

► To cite this version:

Antoinette Hollestelle, Jord H. A. Nagel, Marcel Smid, Suzanne Lam, Fons Elstrodt, et al.. Distinct gene mutation profiles among luminal-type and basal-type breast cancer cell lines. *Breast Cancer Research and Treatment*, 2009, 121 (1), pp.53-64. <10.1007/s10549-009-0460-8>. <hal-00485063>

HAL Id: hal-00485063

<https://hal.science/hal-00485063v1>

Submitted on 20 May 2010

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



HAL Authorization

Distinct Gene Mutation Profiles Among Luminal-Type and Basal-Type Breast Cancer Cell Lines

Antoinette Hollestelle, Jord H.A. Nagel, Marcel Smid, Suzanne Lam, Fons Elstrodt, Marijke Wasielewski, Ser Sue Ng, Pim J. French, Justine K. Peeters, Marieke J. Rozendaal, Muhammad Riaz, Daphne G. Koopman, Timo L.M. ten Hagen, Bertie H.C.G.M. de Leeuw, Ellen C. Zwarthoff, Amina Teunisse, Peter J. van der Spek, Jan G.M. Klijn, Winand N.M. Dinjens, Stephen P. Ethier, Hans Clevers, Aart G. Jochemsen, Michael A. den Bakker, John A. Foekens, John W.M. Martens, and Mieke Schutte

A. Hollestelle (✉), J. H. A. Nagel, M. Smid, F. Elstrodt, M. Wasielewski, M. J. Rozendaal, M. Riaz, D. G.

Koopman, J. G. M. Klijn, J. A. Foekens, J. W. M. Martens, M. Schutte (✉)

Department of Medical Oncology, Josephine Nefkens Institute, Erasmus University Medical Center, P.O. Box 2040, 3000 CA Rotterdam, the Netherlands

e-mail: a.schutte@erasmusmc.nl or a.hollestelle@erasmusmc.nl

M. Smid, M. Wasielewski, J. A. Foekens

Cancer Genomics Centre, Rotterdam, the Netherlands

P. J. French, B. H. C. G. M. de Leeuw

Department of Neurology, Josephine Nefkens Institute, Erasmus University Medical Center, Rotterdam, the Netherlands

E. C. Zwarthoff, W. N. M. Dinjens, M. A. den Bakker

Department of Pathology, Josephine Nefkens Institute, Erasmus University Medical Center, Rotterdam, the Netherlands

J. K. Peeters, P. J. van der Spek

Department of Bioinformatics, Erasmus University Medical Center, Rotterdam, the Netherlands

T. L. M. ten Hagen

Laboratory Experimental Surgical Oncology, Section Surgical Oncology, Department of Surgery, Daniel den Hoed Cancer Center, Erasmus University Medical Center, Rotterdam, the Netherlands

S. Lam, A. Teunisse, A. G. Jochemsen

Department of Molecular Cell Biology, Leiden University Medical Center, Leiden, the Netherlands

S. Ng, H. Clevers

Hubrecht Institute for Developmental Biology and Stem Cell Research, Utrecht, the Netherlands

S. P. Ethier

Barbara Ann Karmanos Cancer Institute, Detroit, Michigan

Note: A. Hollestelle and J. H. A. Nagel contributed equally to this work.

Key Words: Breast cancer subtypes, Histological classification, Intrinsic subtypes, Molecular classification, Mutation analysis.

Abstract

Breast cancer has for long been recognized as a highly diverse tumor group, but the underlying genetic basis has been elusive. Here, we report an extensive molecular characterization of a collection of 41 human breast cancer cell lines. Protein and gene expression analyses indicated that the collection of breast cancer cell lines has retained most, if not all, molecular characteristics that are typical for clinical breast cancers. Gene mutation analyses identified 146 oncogenic mutations among 27 well-known cancer genes, amounting to an average of 3.6 mutations per cell line. Mutations in genes from the p53, RB and PI3K tumor suppressor pathways were widespread among all breast cancer cell lines. Most important, we have identified two gene mutation profiles that specifically associated with luminal-type and basal-type breast cancer cell lines. The luminal mutation profile involved *E-cadherin* and *MAP2K4* gene mutations and amplifications of *Cyclin D1*, *ERBB2* and *HDM2*, whereas the basal mutation profile involved *BRCA1*, *RB1*, *RAS* and *BRAF* gene mutations and deletions of *p16* and *p14ARF*. These subtype-specific gene mutation profiles constitute a genetic basis for the heterogeneity observed among human breast cancers, providing clues for their underlying biology and providing guidance for targeted pharmacogenetic intervention in breast cancer patients.

Introduction

Breast cancer is a heterogeneous disease with respect to prognosis and treatment response of patients. Clinical outcome of cancer patients is to a large extent driven by the biology of their tumors, rendering accurate classification of breast cancers of major importance. Expression of the estrogen receptor (ER), progesterone receptor (PR) and/or epidermal growth factor receptor 2 (ERBB2) has for long guided breast cancer classification. This classification has been refined by cytokeratin (CK) protein expression patterns. About 70% of primary breast cancers express at least one of the luminal CK proteins, whereas almost 30% of primary breast cancers express at least one of each of the luminal and basal CKs (CK7/8/18/19 and CK5/6/14, respectively; [1]). Two minor subtypes express only basal CKs or are negative for both luminal and basal markers, each representing less than 1% of tumors [1]. Luminal breast cancers are more often ER and/or PR positive or have overexpression of

ERBB2. The so-called triple-negative phenotype of tumors that express neither ER, PR nor ERBB2 is observed primarily among basal breast cancers. Basal breast cancers also more often have EGFR expression [2]. Histological classification of breast cancers has clinical relevance in that patients with basal breast cancers more often have a worse prognosis than patients with luminal breast cancers [1, 2]. Perou and colleagues used a 4-protein signature that defined four groups of breast cancers: ERBB2-overexpressing (ERBB2+), luminal (ERBB2- and ER+), basal-like (ERBB2/ER- and CK5+ and/or EGFR+), and the negative group that lacks expression of all four proteins [3]. A major discrepancy among these and other definitions lies in breast cancers that express basal CKs as well as ER, which may be as much as one-third of all basal CK expressing breast cancers [4].

Classification of breast cancers by gene expression profiling has defined five subtypes of breast cancer based on the intrinsic gene set [5]. ER expression was a major classifier, including the luminal A and luminal B subtypes of ER-positive breast cancers. ER-negative breast cancers included the ERBB2+, basal-like and normal-like subtypes. The intrinsic subtypes of breast cancer have clinical relevance [5-7], as have gene expression signatures that had been defined based on clinical outcome of breast cancer patients [8-11]. Importantly, the success of gene expression profiling in breast cancer classification implies a relatively restricted set of recurrent expression patterns among breast cancer subtypes.

Cancer is a genetic disease. Mutations in a cancer gene often induce constitutive changes in the expression of its downstream pathway members or targets. The subtype-specific gene expression patterns among breast cancers were therefore likely to reflect, at least in part, the accumulation of mutations in subtype-specific cancer genes. Here, we have evaluated this concept by extensive molecular characterization of 41 human breast cancer cell lines at protein, transcript and gene level.

Materials and Methods

Cell Lines

The 41 human breast cancer cell lines used in this study are listed in Fig. 1 and described in detail elsewhere [12]. Melanoma cell line M14 was obtained from its provider at Micromet Inc. (Munich, Germany; [13]).

Expression Analyses

Protein expression was determined by western blotting or immunohistochemistry on a tissue microarray of paraffin-embedded cell line samples (antibodies are specified in Supplementary Table S1). Transcript expression was determined by qPCR or by Affymetrix U133A microarrays. qPCR was performed as described [14] on an ABI7700 Taqman Analyzer (Applied Biosystems, Foster City, CA). Ct-values were normalized according *HPRT* and *PBDG* housekeeper Ct-values. DNaseI-treated RNA was antisense biotinylated using the MEGAScript T7 labeling kit (Ambion, Foster City, CA) and Affymetrix U133A microarrays were hybridized according Affymetrix GeneChip Manual, both performed on commercial basis by ServiceXS (Leiden, the Netherlands). Intensity values for all genes were scaled using the global normalization factor as specified by GCOS 1.1, and further normalized with Omniviz software 3.6 (Biowisdom, Maynard, MA). Intensity values <30 were set to 30. Differential gene expression was based on log2 transformed distances to the geometric mean for each probe set. Patterns of correlation were revealed by applying the Pearson matrix-ordering method that sorts samples into correlated blocks. The Stanford intrinsic gene set [15] was translated into an Affymetrix intrinsic gene set, including 451 probe sets from the Stanford list of 496 genes [7]. Intrinsic subtypes were classified by average distance linkage hierarchical clustering with non-centered correlation as distance metric [16]. Microarray data are available at GEO (<http://www.ncbi.nlm.nih.gov/geo/>), accession number GSE16795.

Gene Mutation Analyses

Gene sequence alterations typically were identified by direct sequencing of PCR-amplified genomic templates that included exon sequences and intron/exon boundaries. For *RB1*, exons 3 through 27 were analyzed by sequencing of RT-PCR amplified templates of emetine-treated cell lines. The complete coding sequences have been analyzed for *BRCA1* [17], *CHEK2* [18], *E-cadherin/CDH1*

([19] and Hollestelle *et al.*, submitted for publication), *MAP2K4* [20], *MSH6*, *p14ARF*, *p16*, *p53* [21], *PIK3CA* [12], *PTEN* [12] and *RB1*, whereas only known mutational hotspots have been analyzed for *AKT1* (codon 17), *APC* (mutation E1577X), *BRAF* (exons 7, 11 and 15; [12]), *CDK4* (p16-binding domain in exon 2), *FGFR3* (codons 248, 249, 372, 375, 393 and 652), and codons 12, 13 and 61 for *HRAS*, *KRAS* and *NRAS* [12]. All oncogenic mutations have been confirmed by sequencing of an independently amplified DNA and/or RNA template.

Amplifications of *AKT1*, *AKT2*, *AKT3*, *CDK4*, *c-MYC*, *Cyclin D1*, *EGFR*, *ERBB2* and *HDM2* were defined by genomic gain of the gene locus with concurrent transcript and/or protein overexpression. Genomic gains were determined by screening of publicly available CGH and SNP data ([22] and www.sanger.ac.uk), requiring gain of at least four allele copies. Genomic gain of the *ERBB2* and *EGFR* loci was determined by chromogenic in situ hybridization (SPoT-Light® CISH™ kit, Zymed).

Promoter hypermethylation was determined for *BRCA1*, *E-cadherin*, *p16*, *p14ARF* and *RB1*, by azacytidine assays and/or methylation specific PCR ([23] and Hollestelle *et al.*, submitted for publication). Wnt pathway activation was determined by TOP/FOP reporter construct assays ([19]; Ng *et al.*, submitted for publication).

Results

Expression Analysis of Breast Cancer Cell Lines

We have characterized 41 human breast cancer cell lines by extensive expression analyses, using immunohistochemistry and microarray expression profiling (Fig. 1 and Supplementary Table S2). CK protein expression patterns classified the breast cancer cell lines in four histological phenotypes: luminal (n=22), combined luminal/basal (n=4), null (n=7) and basal (n=1) [1]. Gene expression profiling identified two main clusters of cell lines (Fig. 2A). The major cluster contained all cell lines that (over)expressed ER, PR and/or ERBB2 proteins and that were luminal by CK expression (Fig. 1 and Supplementary Table S2). The minor cluster contained triple-negative cell lines only, with a subdivision in combined luminal/basal and null phenotype cell lines by CK expression. Protein expression of EGFR and particularly of the EMT protein markers Vimentin and N-cadherin was mostly

restricted to combined luminal/basal and null phenotype cell lines. A similar clustering of breast cancer cell lines was obtained by hierarchical clustering based on the intrinsic gene set (Fig. 2B; [7, 15]). Luminal phenotype cell lines by CK expression classified either as luminal or ERBB2+ by intrinsic gene expression. Similarly, combined luminal/basal and null phenotype cell lines were basal-like and normal-like, respectively, by intrinsic gene expression. The 4-protein signature of ERBB2, ER, CK5 and EGFR [3] was not able to distinguish combined luminal/basal or basal-like cell lines from null or normal-like cell lines. However, this signature identified all thirteen cell lines with ERBB2 overexpression. The results of these different classification methods were highly concordant and revealed a dichotomy among the breast cancer cell lines that we have designated "luminal-type" (n=25) and "basal-type" (n=15), irrespective of their further subclassification.

Mutation Analysis of Breast Cancer Cell Lines

In an ongoing effort to characterize the breast cancer cell lines, we have performed mutation analyses of 27 well-known cancer genes (results of 16 genes are first reported here; all mutations are detailed in Supplementary Table S3). Oncogenic mutations were defined as those mutations that predicted a premature termination or a significant deletion/insertion of the encoded proteins, splice site mutations, and missense mutations that previously had been identified in clinical cancers or had been associated with hereditary cancer syndromes. For oncogenes that reportedly are amplified in clinical cancers, we considered chromosomal gains with concurrent transcript and/or protein overexpression also as oncogenic mutations or amplifications. Together, we have identified 146 oncogenic mutations among 24 cancer genes (Fig. 1). Oncogenic mutations in tumor suppressor genes included 30 sizeable deletions, 26 nonsense mutations and small deletions/insertions causing premature terminations, 7 splice site mutations and 29 missense mutations, totaling to 92 mutations. In concordance with the recessive nature of tumor suppressor gene mutations, 87 (95%) of these mutations were bi-allelic. Oncogenic mutations in oncogenes included 29 amplifications and 25 missense mutations, totaling to 54 mutations. Three tumor suppressor genes had transcriptional silencing by promoter hypermethylation, together in 20 cell lines. The 146 oncogenic mutations were identified among 40 of

the 41 breast cancer cell lines, resulting in an average of 3.6 mutations per cell line and ranging from no to seven mutations each identified in a single cell line (Fig. 1).

For most cancer genes, we also determined transcript and protein expression patterns. As anticipated, transcript expression of mutant genes with premature terminations typically was low or absent whereas missense mutations resulted in relatively normal transcript expression levels. Protein expression patterns mostly followed transcript expression patterns and/or were in concordance with those reported for clinical cancers. For example, *p53* mutants with premature terminations typically had low transcript and protein expression whereas *p53* missense mutants had relatively normal transcript expression but high protein expression (Table 1). For some mutant genes, we observed expression modulation of other proteins in the same tumor suppressor pathway. In the RB pathway, for example, most *RB1* mutants had high p16 protein expression and very low Cyclin D1 protein expression whereas *Cyclin D1* mutants had low p16 protein expression (Table 2 and Fig. 3). Similar observations had been made by Gauthier *et al.*, who have associated deregulation of RB signaling with basal-like breast cancers [24]. Together, the transcript and protein expression patterns of mutant cancer genes thus further confirmed the functional significance of the identified oncogenic gene mutations.

Gene Mutation Profiles among Breast Cancer Cell Lines Associate with Breast Cancer Subtypes

Compilation of the mutation data revealed two prevalent gene mutation patterns among the breast cancer cell lines (Fig. 1). The first pattern involved frequent mutations among the cell lines in genes from the same tumor suppressor pathway. These included the p53 pathway in 90% of the cell lines (*p53*, *CHEK2*, *HDM2*, *c-MYC* and *p14ARF*; Table 1), the RB pathway in 64% (*RB1*, *p16*, *Cyclin D1* and *CDK4*; Table 2) and the PI3K pathway in 56% (*PTEN*, *PIK3CA*, *AKT1*, *AKT2* and *AKT3*; [12]). Oncogenic mutations within the RB pathway were essentially mutually exclusive among cell lines. Similar to other reports [25-27], *PIK3CA* mutations were biallelic in two cell lines and monoallelic concurrent with *AKT1* amplification or monoallelic *PTEN* mutation in two other cell lines [12]. Somewhat surprising was the frequent concurrent mutation of *p53* or *CHEK2* mutations with

mutations in *p14ARF*, *C-MYC* or *HDM2*. It may be of interest that all *p53/p14ARF* and *p53/c-MYC* double mutants had *p53* missense mutations, suggesting some residual activity of these *p53* mutants (Table 1 and Supplementary Table S3). Whatever the functional significance of these double mutations [28], the frequent mutational abrogation of the *p53*, *RB* and *PI3K* pathways among the cell lines suggests that their functionality is of major importance in breast carcinogenesis.

Most exciting, the second gene mutation pattern involved oncogenic mutations that cluster in a particular subset of breast cancer cell lines (Fig. 1). These included a "luminal mutation profile" involving *E-cadherin* and *MAP2K4* gene mutations and amplifications of *Cyclin D1*, *ERBB2* and *HDM2*, and a "basal mutation profile" involving *BRCA1*, *RB1*, *RAS* and *BRAF* gene mutations and deletions of *p16* and *p14ARF*. Specifically, the cohort of 25 luminal-type breast cancer cell lines included 35 of 36 mutations in luminal mutation profile genes (Fisher's exact $P < 0.0001$). The cohort of 15 basal-type breast cancer cell lines included 33 of 39 mutations in basal mutation profile genes ($P < 0.0001$). The existence of specific gene mutation profiles for each of the two major subtypes of breast cancer cell lines suggests a dichotomy in the genetic basis of human breast cancer.

Discussion

Expression Analyses Suggest Two Major Types of Luminal and Basal Breast Cancer Cell Lines

Classification of the breast cancer cell lines was remarkably consistent, whether they were classified by classical breast cancer parameters (ER, PR and ERBB2), histological criteria (CK protein expression or the 4-protein signature), or gene expression profiling (intrinsic subtypes; Fig. 1). Classification of the ERBB2+ subtype is ambiguous in breast cancer cell lines as well as clinical breast cancers due to differences between ERBB2 protein overexpression, gene amplification, and expression of genes from the intrinsic gene set. Our mutation analyses strongly suggested that the basal-like or combined luminal/basal and normal-like or null subtypes of cell lines represent two ends of a spectrum of basal-type breast cancer (Fig. 1). This was supported by the protein expression profiling data, specifically expression of E-cadherin, P-cadherin, N-cadherin, Vimentin and EGFR. Most notable, a similar interrelationship has been proposed for clinical breast cancers [29] and

accordingly, Neve *et al.* had designated these two subtypes of breast cancer cell lines as basal A and basal B [22]. Our molecular characterization of breast cancer cell lines is concordant with the concept that two major types of luminal and basal breast cancers exist that may be subdivided further by protein expression and/or intrinsic gene expression profiling.

Accumulation of Many Mutations in Well-known Cancer Genes in Breast Cancer Cell Lines

We have identified 146 oncogenic mutations among 41 breast cancer cell lines. This mutation frequency was much higher than anticipated. For example, the mutation frequencies of *p53* and *E-cadherin* were about twice as high as reported for clinical breast cancers [30, 31]. This high mutation frequency likely reflects a higher rate of mutation detection in cell lines. Importantly, the presence of non-neoplastic cells in clinical breast cancer samples essentially precludes identification of homozygous deletions, whereas these represent 20% of the mutations that we have identified. The Sanger Institute also has screened known cancer genes in breast cancer cell lines, with 15 genes and 23 cell lines overlapping with our study (www.sanger.ac.uk as per February 2009). Sanger failed to identify 23 (35%) of 65 mutations that we had detected and we failed to identify one mutation (that we subsequently confirmed). The reason for this discrepancy is unclear, but we expect that our mutation screens may involve more detailed primer design, PCR optimization and data analysis. Sample swapping appears unlikely since seven cell lines with discrepant results also carried signature mutations identified by both Sanger and us (Supplementary Table S4). Our results for the other two cell lines have been confirmed by literature reports [25, 32-35]. Overall, 14 of 23 discrepant mutations had been confirmed in literature reports, also rendering incidental sample swapping unlikely [25, 32-38]. We consider it unlikely that a significant proportion of mutations has been acquired during *in vitro* propagation of the cell lines, although there is no hard evidence to support this notion nor, for that matter, to reject it. Most important, the majority of the identified gene alterations appeared to be of functional relevance as they had previously been reported in clinical cancer samples, in the germline of patients with a hereditary cancer syndrome, or were sizeable deletions or intragenic mutations that predicted a premature protein termination (and thus are less likely to be recurrent). Silent mutations or

missense mutations of unclear significance represented less than 10% of all identified deviations from the reference coding gene sequences (5 and 7 mutations, respectively). A more likely explanation for the high mutation frequency among cell lines is a bias for breast cancers that are more easily propagated *in vitro*, such as *p53* mutant breast cancers and likely also those of the null or normal-like subtype both being cancers that are known to have a more aggressive clinical course. This might, however, not necessarily be disadvantageous as it allows one to study the more clinically-needy breast cancers.

Gene Mutation Profiles Provide a Genetic Basis for Luminal-type and Basal-type Breast Cancers

We have identified subtype-specific gene mutation profiles among the breast cancer cell lines. Of the mutations in luminal mutation profile genes, 97% had been identified among the luminal cell lines. The DU4475 cell line was the single non-luminal cell line that had a mutation in a luminal mutation profile gene: *MAP2K4*. Basal-type breast cancer cell lines carried 85% of the mutations in basal mutation profile genes. DU4475 again was atypical with mutations in *RB1* and *BRAF*. The MDA-MB-435s cell line was the only basal-type cell line that did not express EGFR proteins. MDA-MB-435s was recently shown to be genetically identical to the M14 melanoma cell line from the NCI-60 cell line panel (several MDA-MB-435s vials versus a single M14 vial; [39]). We have obtained M14 cells from several laboratories, including a vial of passage-10 cells from its originator, and MDA-MB-435s was obtained from ATCC at passage-239. Indeed, microsatellite analysis confirmed that the two cell lines were identical. It thus remains unclear which is the correct origin of the cell lines, although recent evidence suggests that they may be of breast origin [40]. Be as it may, it is not likely that the other basal-type breast cancer cell lines are melanocytic. Four of the basal-type cell lines carry *BRCA1* mutations and one carries the *CHEK2* 1100delC mutation, both well-known breast cancer susceptibility genes [41]. In fact, the consistent histological and intrinsic breast cancer classification of the basal-type cell lines strongly suggests that a melanoma-like gene mutation profile (that includes mutations in *p16*, *RB1* and *RAS* pathway genes [42]) is characteristic for basal-type breast cancers. Comparable gene mutation profiles among basal-type breast cancers and melanomas, and likely also pancreatic

cancers [43], might be conceivable given the typically aggressive clinical course of these tumor types and their frequent inherent therapy resistance. In this respect, distinct gene mutation profiles among breast cancers could very well explain the significant heterogeneity in clinical outcome of breast cancer patients.

It is of interest that recent large-scale sequencing efforts have identified a similar number of somatic oncogenic mutations among breast cancers and colorectal cancers, but that the spectrum of mutated pathways was far more diverse among breast cancers (an average of 14 and 15 mutations per tumor among 108 and 38 pathways for breast cancer and colorectal cancer, respectively; [44, 45]). These differences in mutation spectrum suggested genetic heterogeneity among breast cancers, and are thus in line with our finding of two distinct gene mutations profiles. After all, the existence of two gene mutation profiles among breast cancers implies more mutant genes when considering breast cancer at large. Similarly, mutations in subtype-specific cancer genes will then be less prevalent. Our study has focused on mutations in well-known cancer genes and the identified mutations therefore are more likely to represent drivers of breast carcinogenesis. Although this approach does not draw the full genomic landscape of breast cancer, it did allow us to decipher specific patterns of gene mutations among the two major breast cancer types. Of course, breast cancer cell lines are not clinical breast cancers. As a proof of principle, we have shown that *E-cadherin* mutation and *E-cadherin* promoter methylation indeed also associate with distinct pathological types of clinical breast cancers (Hollestelle *et al.*, submitted for publication). It also is significant that mutations in three other genes follow patterns reported for clinical breast cancers, including the exclusive presence of *Cyclin D1* amplifications in ER-positive breast cancers, *RB1* mutations in ER-negative breast cancers and the concurrent mutation of *BRCA1* and *p53* in ER-negative breast cancers [24, 46-49]. If the evidence is to be extended to other genes from the mutation profiles, we may at last have begun to elucidate the genetic basis for the inherent biological and clinical heterogeneity among human breast cancers.

Acknowledgements

Grant support: Susan G. Komen Breast Cancer Foundation, Erasmus MC Mreace, Dutch Cancer Society, Netherlands Genomics Initiative (NGI)/Netherlands Organization for Scientific Research (NWO), Association for International Cancer Research, and Survivorsland Foundation, Vlissingen, The Netherlands.

We appreciate the technical assistance of Michel Molier, Sofia Zuñiga, Mieke Timmermans, Jurjen J. Boonstra, Hein Sleddens and members of the Pathology Immunohistochemistry laboratories at Erasmus MC. We thank Dr. Jörg Volkland, Dr. Nelleke Gruis, Dr. Goos van Muijen and Dr. Jan Verheijen for kindly providing us with M14 cells.

References

- 1 Abd El-Rehim DM, Pinder SE, Paish CE, Bell J, Blamey RW, Robertson JF, Nicholson RI, Ellis IO (2004) Expression of luminal and basal cytokeratins in human breast carcinoma. *J Pathol* 203:661-671.
- 2 Korsching E, Jeffrey SS, Meinerz W, Decker T, Boecker W, Buerger H (2008) Basal carcinoma of the breast revisited: an old entity with new interpretations. *J Clin Pathol* 61:553-560.
- 3 Nielsen TO, Hsu FD, Jensen K, Cheang M, Karaca G, Hu Z, Hernandez-Boussard T, Livasy C, Cowan D, Dressler L, Akslen LA, Ragaz J, Gown AM, Gilks CB, van de Rijn M, Perou CM (2004) Immunohistochemical and clinical characterization of the basal-like subtype of invasive breast carcinoma. *Clin Cancer Res* 10:5367-5374.
- 4 Rakha EA, El-Sayed ME, Green AR, Paish EC, Lee AH, Ellis IO (2007) Breast carcinoma with basal differentiation: a proposal for pathology definition based on basal cytokeratin expression. *Histopathology* 50:434-438.
- 5 Sorlie T, Tibshirani R, Parker J, Hastie T, Marron JS, Nobel A, Deng S, Johnsen H, Pesich R, Geisler S, Demeter J, Perou CM, Lonning PE, Brown PO, Borresen-Dale AL, Botstein D (2003) Repeated observation of breast tumor subtypes in independent gene expression data sets. *Proc Natl Acad Sci U S A* 100:8418-8423.

- 6 Rouzier R, Perou CM, Symmans WF, Ibrahim N, Cristofanilli M, Anderson K, Hess KR, Stec J, Ayers M, Wagner P, Morandi P, Fan C, Rabiul I, Ross JS, Hortobagyi GN, Pusztai L (2005) Breast cancer molecular subtypes respond differently to preoperative chemotherapy. *Clin Cancer Res* 11:5678-5685.
- 7 Smid M, Wang Y, Zhang Y, Sieuwerts AM, Yu J, Klijn JGM, Foekens JA, Martens JWM (2008) Subtypes of breast cancer show preferential site of relapse. *Cancer Research* 68:3108-3114.
- 8 van 't Veer LJ, Dai H, van de Vijver MJ, He YD, Hart AA, Mao M, Peterse HL, van der Kooy K, Marton MJ, Witteveen AT, Schreiber GJ, Kerkhoven RM, Roberts C, Linsley PS, Bernards R, Friend SH (2002) Gene expression profiling predicts clinical outcome of breast cancer. *Nature* 415:530-536.
- 9 Wang Y, Klijn JG, Zhang Y, Sieuwerts AM, Look MP, Yang F, Talantov D, Timmermans M, Meijer-van Gelder ME, Yu J, Jatkoe T, Berns EM, Atkins D, Foekens JA (2005) Gene-expression profiles to predict distant metastasis of lymph-node-negative primary breast cancer. *Lancet* 365:671-679.
- 10 Paik S, Shak S, Tang G, Kim C, Baker J, Cronin M, Baehner FL, Walker MG, Watson D, Park T, Hiller W, Fisher ER, Wickerham DL, Bryant J, Wolmark N (2004) A multigene assay to predict recurrence of tamoxifen-treated, node-negative breast cancer. *N Engl J Med* 351:2817-2826.
- 11 Foekens JA, Wang Y, Martens JWM, Berns EMJJ, Klijn JGM (2008) The use of genomic tools for the molecular understanding of breast cancer and to guide personalized medicine. *Drug Discovery Today* 13:481-487.
- 12 Hollestelle A, Elstrodt F, Nagel JH, Kallemeijn WW, Schutte M (2007) Phosphatidylinositol-3-OH kinase or RAS pathway mutations in human breast cancer cell lines. *Mol Cancer Res* 5:195-201.
- 13 Chee DO, Boddie AW, Roth JA, Holmes EC, Morton DL (1976) Production of melanoma-associated antigen(s) by a defined malignant melanoma cell strain grown in chemically defined medium. *Cancer Res* 36:1503-1509.

- 14 Sieuwerts AM, Meijer-van Gelder ME, Timmermans M, Trapman AM, Garcia RR, Arnold M, Goedheer AJ, Portengen H, Klijn JG, Foekens JA (2005) How ADAM-9 and ADAM-11 differentially from estrogen receptor predict response to tamoxifen treatment in patients with recurrent breast cancer: a retrospective study. *Clin Cancer Res* 11:7311-7321.
- 15 Perou CM, Sorlie T, Eisen MB, van de Rijn M, Jeffrey SS, Rees CA, Pollack JR, Ross DT, Johnsen H, Akslen LA, Fluge O, Pergamenschikov A, Williams C, Zhu SX, Lonning PE, Borresen-Dale AL, Brown PO, Botstein D (2000) Molecular portraits of human breast tumours. *Nature* 406:747-752.
- 16 Eisen MB, Spellman PT, Brown PO, Botstein D (1998) Cluster analysis and display of genome-wide expression patterns. *Proc Natl Acad Sci U S A* 95:14863-14868.
- 17 Elstrodt F, Hollestelle A, Nagel JH, Gorin M, Wasielewski M, van den Ouweland A, Merajver SD, Ethier SP, Schutte M (2006) BRCA1 mutation analysis of 41 human breast cancer cell lines reveals three new deleterious mutants. *Cancer Res* 66:41-45.
- 18 Wasielewski M, Hanifi-Moghaddam P, Hollestelle A, Merajver SD, van den Ouweland A, Klijn JGM, Ethier SP, Schutte M (2008) Deleterious CHEK2 1100delC and L303X mutants identified among 38 human breast cancer cell lines. *Breast Cancer Res Treat* DOI 10.1007/310549-008-9942-3.
- 19 van de Wetering M, Barker N, Harkes IC, van der Heyden M, Dijk NJ, Hollestelle A, Klijn JG, Clevers H, Schutte M (2001) Mutant E-cadherin breast cancer cells do not display constitutive Wnt signaling. *Cancer Res* 61:278-284.
- 20 Su GH, Song JJ, Repasky EA, Schutte M, Kern SE (2002) Mutation rate of MAP2K4/MKK4 in breast carcinoma. *Hum Mutat* 19:81.
- 21 Wasielewski M, Elstrodt F, Klijn JG, Berns EM, Schutte M (2006) Thirteen new p53 gene mutants identified among 41 human breast cancer cell lines. *Breast Cancer Res Treat* 99:97-101.
- 22 Neve RM, Chin K, Fridlyand J, Yeh J, Baehner FL, Fevr T, Clark L, Bayani N, Coppe JP, Tong F, Speed T, Spellman PT, DeVries S, Lapuk A, Wang NJ, Kuo WL, Stilwell JL, Pinkel D, Albertson DG, Waldman FM, McCormick F, Dickson RB, Johnson MD, Lippman M, Ethier S,

- Gazdar A, Gray JW (2006) A collection of breast cancer cell lines for the study of functionally distinct cancer subtypes. *Cancer Cell* 10:515-527.
- 23 Esteller M, Silva JM, Dominguez G, Bonilla F, Matias-Guiu X, Lerma E, Bussaglia E, Prat J, Harkes IC, Repasky EA, Gabrielson E, Schutte M, Baylin SB, Herman JG (2000) Promoter hypermethylation and BRCA1 inactivation in sporadic breast and ovarian tumors. *J Natl Cancer Inst* 92:564-569.
- 24 Gauthier ML, Berman HK, Miller C, Kozakeiwicz K, Chew K, Moore D, Rabban J, Chen YY, Kerlikowske K, Tlsty TD (2007) Abrogated response to cellular stress identifies DCIS associated with subsequent tumor events and defines basal-like breast tumors. *Cancer Cell* 12:479-491.
- 25 Saal LH, Holm K, Maurer M, Memeo L, Su T, Wang X, Yu JS, Malmstrom PO, Mansukhani M, Enoksson J, Hibshoosh H, Borg A, Parsons R (2005) PIK3CA mutations correlate with hormone receptors, node metastasis, and ERBB2, and are mutually exclusive with PTEN loss in human breast carcinoma. *Cancer Res* 65:2554-2559.
- 26 Stemke-Hale K, Gonzalez-Angulo AM, Lluch A, Neve RM, Kuo WL, Davies M, Carey M, Hu Z, Guan Y, Sahin A, Symmans WF, Pusztai L, Nolden LK, Horlings H, Berns K, Hung MC, van de Vijver MJ, Valero V, Gray JW, Bernardis R, Mills GB, Hennessy BT (2008) An integrative genomic and proteomic analysis of PIK3CA, PTEN, and AKT mutations in breast cancer. *Cancer Res* 68:6084-6091.
- 27 Oda K, Okada J, Timmerman L, Rodriguez-Viciano P, Stokoe D, Shoji K, Taketani Y, Kuramoto H, Knight ZA, Shokat KM, McCormick F (2008) PIK3CA cooperates with other phosphatidylinositol 3'-kinase pathway mutations to effect oncogenic transformation. *Cancer Res* 68:8127-8136.
- 28 Sherr CJ (2006) Divorcing ARF and p53: an unsettled case. *Nat Rev Cancer* 6:663-673.
- 29 Fulford LG, Easton DF, Reis-Filho JS, Sofronis A, Gillett CE, Lakhani SR, Hanby A (2006) Specific morphological features predictive for the basal phenotype in grade 3 invasive ductal carcinoma of breast. *Histopathology* 49:22-34.
- 30 Borresen-Dale AL (2003) TP53 and breast cancer. *Hum Mutat* 21:292-300.

- 31 Berx G, Becker KF, Hofler H, van Roy F (1998) Mutations of the human E-cadherin (CDH1) gene. *Hum Mutat* 12:226-237.
- 32 Concin N, Zeillinger C, Tong D, Stimpfl M, Konig M, Printz D, Stonek F, Schneeberger C, Hefler L, Kainz C, Leodolter S, Haas OA, Zeillinger R (2003) Comparison of p53 mutational status with mRNA and protein expression in a panel of 24 human breast carcinoma cell lines. *Breast Cancer Res Treat* 79:37-46.
- 33 Hiraguri S, Godfrey T, Nakamura H, Graff J, Collins C, Shayesteh L, Doggett N, Johnson K, Wheelock M, Herman J, Baylin S, Pinkel D, Gray J (1998) Mechanisms of inactivation of E-cadherin in breast cancer cell lines. *Cancer Res* 58:1972-1977.
- 34 Prosperi MT, Dupre G, Lidereau R, Goubin G (1990) Point mutation at codon 12 of the Ki-ras gene in a primary breast carcinoma and the MDA-MB-134 human mammary carcinoma cell line. *Cancer Lett* 51:169-174.
- 35 Teng DH, Perry WL, 3rd, Hogan JK, Baumgard M, Bell R, Berry S, Davis T, Frank D, Frye C, Hattier T, Hu R, Jammulapati S, Janecki T, Leavitt A, Mitchell JT, Pero R, Sexton D, Schroeder M, Su PH, Swedlund B, Kyriakis JM, Avruch J, Bartel P, Wong AK, Tavtigian SV, et al. (1997) Human mitogen-activated protein kinase kinase 4 as a candidate tumor suppressor. *Cancer Res* 57:4177-4182.
- 36 Saal LH, Gruvberger-Saal SK, Persson C, Lovgren K, Junttinen M, Staaf J, Jonsson G, Pires MM, Maurer M, Holm K, Koujak S, Subramaniam S, Vallon-Christersson J, Olsson H, Su T, Memeo L, Ludwig T, Ethier SP, Krogh M, Szabolcs M, Murty VV, Isola J, Hibshoosh H, Parsons R, Borg A (2008) Recurrent gross mutations of the PTEN tumor suppressor gene in breast cancers with deficient DSB repair. *Nat Genet* 40:102-107.
- 37 Tomlinson GE, Chen TT, Stastny VA, Virmani AK, Spillman MA, Tonk V, Blum JL, Schneider NR, Wistuba, II, Shay JW, Minna JD, Gazdar AF (1998) Characterization of a breast cancer cell line derived from a germ-line BRCA1 mutation carrier. *Cancer Res* 58:3237-3242.
- 38 Runnebaum IB, Nagarajan M, Bowman M, Soto D, Sukumar S (1991) Mutations in p53 as potential molecular markers for human breast cancer. *Proc Natl Acad Sci U S A* 88:10657-10661.

- 39 Rae JM, Creighton CJ, Meck JM, Haddad BR, Johnson MD (2007) MDA-MB-435 cells are derived from M14 melanoma cells--a loss for breast cancer, but a boon for melanoma research. *Breast Cancer Res Treat* 104:13-19.
- 40 Chambers AF (2009) MDA-MB-435 and M14 Cell Lines: Identical but not M14 Melanoma? *Cancer Res* 69:5292-5293.
- 41 Walsh T, King MC (2007) Ten genes for inherited breast cancer. *Cancer Cell* 11:103-105.
- 42 Gray-Schopfer V, Wellbrock C, Marais R (2007) Melanoma biology and new targeted therapy. *Nature* 445:851-857.
- 43 Jaffee EM, Hruban RH, Canto M, Kern SE (2002) Focus on pancreas cancer. *Cancer Cell* 2:25-28.
- 44 Sjoblom T, Jones S, Wood LD, Parsons DW, Lin J, Barber TD, Mandelker D, Leary RJ, Ptak J, Silliman N, Szabo S, Buckhaults P, Farrell C, Meeh P, Markowitz SD, Willis J, Dawson D, Willson JK, Gazdar AF, Hartigan J, Wu L, Liu C, Parmigiani G, Park BH, Bachman KE, Papadopoulos N, Vogelstein B, Kinzler KW, Velculescu VE (2006) The consensus coding sequences of human breast and colorectal cancers. *Science* 314:268-274.
- 45 Wood LD, Parsons DW, Jones S, Lin J, Sjoblom T, Leary RJ, Shen D, Boca SM, Barber T, Ptak J, Silliman N, Szabo S, Dezso Z, Ustyansky V, Nikolskaya T, Nikolsky Y, Karchin R, Wilson PA, Kaminker JS, Zhang Z, Croshaw R, Willis J, Dawson D, Shipitsin M, Willson JK, Sukumar S, Polyak K, Park BH, Pethiyagoda CL, Pant PV, Ballinger DG, Sparks AB, Hartigan J, Smith DR, Suh E, Papadopoulos N, Buckhaults P, Markowitz SD, Parmigiani G, Kinzler KW, Velculescu VE, Vogelstein B (2007) The genomic landscapes of human breast and colorectal cancers. *Science* 318:1108-1113.
- 46 Bieche I, Lidereau R (2000) Loss of heterozygosity at 13q14 correlates with RB1 gene underexpression in human breast cancer. *Mol Carcinog* 29:151-158.
- 47 Barnes DM, Gillett CE (1998) Cyclin D1 in breast cancer. *Breast Cancer Res Treat* 52:1-15.
- 48 Gasco M, Yulug IG, Crook T (2003) TP53 mutations in familial breast cancer: functional aspects. *Hum Mutat* 21:301-306.
- 49 Turner NC, Reis-Filho JS (2006) Basal-like breast cancer and the BRCA1 phenotype. *Oncogene* 25:5846-5853.

Legends to the Tables and Figures

Figure 1. Molecular characterization of 41 human breast cancer cell lines. **Left panel,** classification of the cell lines by expression analyses. Classification by cytokeratin expression: L, luminal: CK8/18+ and/or CK19+, CK5-; C, combined luminal/basal: CK8/18+ and/or CK19+, CK5+; N, null: CK8/18 low, CK19-, CK5-; B, basal: CK8/18 low, CK19-, CK5+. Classification by the 4-protein signature: E, ERBB2 overexpression; L, luminal: ERBB2-, ER+; B, basal: ERBB2/ER-, CK5+ and/or EGFR+; N, negative: ERBB2/ER/CK5/EGFR-. Classification by intrinsic gene expression: E, ERBB2+; L, luminal; B, basal-like; N, normal-like; O, other subtype. For individual proteins: P, expression and PP, overexpression in blue and no expression in white. The two major types of breast cancer cell lines are indicated at the far left. Classification of the ERBB2 subtype varies with definitions, but is clinically defined by protein overexpression (score 3) or by indecisive overexpression (score 2) with concurrent gene amplification. **Middle panel,** gene mutation analysis of the cell lines. M, oncogenic mutations, D, sizeable deletions and A, amplifications in red; M*, heterozygous oncogenic mutations in tumor suppressor genes in pink; wild-type genes in white; H, promoter hypermethylation in yellow and Y, constitutive Wnt pathway activation in green. nd, not determined. The black borders indicate the two gene mutation profiles. **Right panel,** number of oncogenic mutations in each cell line and microsatellite instability (MSI) with BAT 25, 26 and 40: N, no; Y, yes; 40, MSI with BAT 40 only. Ki-67 immunohistochemistry: 1, less than 33% of cells positive; 2, 33-66% of cells positive; 3, more than 66% of cells positive.

Figure 2. Global gene expression and intrinsic subtypes of breast cancer cell lines. **A.** Pearson correlation plot based on the subset of 5,490 log₂GM <-2 and >2 differentially expressed probe sets. The Pearson correlation coefficient algorithm positions samples according to their overall similarity in gene expression, where red indicates high overall similarity and blue indicates low similarity. **B.** Dendrogram of hierarchical clustering based on the intrinsic gene set. See Fig. 1 for color coding of intrinsic subtypes.

Figure 3. Protein expression of RB pathway genes by western blotting.

Table 1. Mutation Analysis of p53 Pathway Genes.

Breast cancer cell lines	<i>p14ARF</i> gene sequence	<i>p14ARF</i> transcript expression	<i>p53</i> gene sequence	<i>p53</i> transcript expression	<i>p53</i> protein expression	<i>CHEK2</i> gene sequence	<i>CHEK2</i> protein expression	<i>c-MYC</i> gene locus	<i>c-MYC</i> transcript expression	<i>HDM2</i> gene locus
ZR75-30	methyalted	+/-	wild-type	+	+	nd	+	gain	+	balanced
DU4475	methyalted	+	wild-type	+	+	wild-type	+	balanced	+	balanced
MCF-7	deleted	-	wild-type	+	+	wild-type	+	gain	+	gain ^{amp}
SK-BR-7	deleted	-	wild-type	+	+	wild-type	+	nd	+	nd
SUM102PT	deleted	-	wild-type	+	nd	mutant*	-	balanced	+	nd
BT20	deleted	-	mutant AA	+	++	wild-type	+	balanced	+	balanced
SUM149PT	deleted	-	mutant AA	+	++	wild-type	+	balanced	+	balanced
SUM229PE	deleted	-	mutant AA	+	++	wild-type	+	balanced	+	nd
SUM1315M02	deleted	-	mutant AA	+	++	wild-type	+	gain ^{amp}	++	balanced
Hs578T	deleted	-	mutant AA	++	++	wild-type	+	gain	+	balanced
MDA-MB-231	deleted	-	mutant AA	++	++	wild-type	+	balanced	+	balanced
OCUB-F	mutant	+	mutant AA*	+	nd	nd	+	gain	+	balanced
MDA-MB-361	mutant?	+/-	mutant TR	+	+/-	wild-type	+	balanced	+	balanced
SUM159PT	wild-type	-	mutant AA	+	++	wild-type	+	gain	+	balanced
BT483	wild-type	+	mutant AA	+	nd	wild-type	+	gain	+	balanced
MDA-MB-330	wild-type	+	mutant AA	+	++	wild-type	+	nd	+/-	nd
MDA-MB-435S	wild-type	+	mutant AA	+	++	wild-type	+	balanced	+	balanced
MDA-MB-468	wild-type	+	mutant AA	+	++	wild-type	+	balanced	+	balanced
T47D	wild-type	+	mutant AA	+	++	wild-type	+	balanced	+	balanced
SK-BR-5	wild-type	+	mutant AA	+	++	wild-type	+	nd	+/-	nd
BT474	wild-type	+	mutant AA	++	++	nd	+	balanced	+	balanced
EVSA-T	wild-type	+	mutant AA	++	++	wild-type	+	gain	+	balanced
SK-BR-3	wild-type	+	mutant AA	++	++	wild-type	+	gain ^{amp}	++	balanced
MDA-MB-415	wild-type	+	mutant AA	++	+	wild-type	+	balanced	+	balanced
BT549	wild-type	++	mutant AA	++	++	wild-type	+	balanced	+	balanced
CAMA-1	wild-type	++	mutant AA	+	++	wild-type	+	gain	+	balanced
MDA-MB-134VI	wild-type	++	mutant AA	+	+/-	wild-type	+	balanced	+	balanced
SUM225CWN	nd	nd	mutant AA	nd	nd	wild-type	+	nd	nd	balanced
SUM190PT	nd	nd	mutant TR	nd	+	wild-type	+	gain	nd	balanced
HCC1937	wild-type	++	mutant TR	+/-	+	wild-type	+	balanced	+	balanced
MDA-MB-157	wild-type	++	mutant TR	+/-	+/-	wild-type	+	gain	+	balanced
MDA-MB-436	wild-type	+	mutant TR	+/-	+/-	wild-type	+	balanced	+	balanced
SUM44PE	wild-type	+	mutant TR	+/-	+/-	nd	+	nd	+	balanced
SUM185PE	wild-type	+	mutant TR	+/-	+/-	wild-type	+	balanced	+/-	balanced
UACC893	wild-type	+	mutant TR	+/-	+	wild-type	+	gain	+	balanced
SUM52PE	wild-type	+	mutant TR	+/-	+/-	wild-type	+	gain	+	gain ^{amp}
MDA-MB-453	wild-type	+	deleted	nd	+	wild-type	+	balanced	+	balanced
UACC812	wild-type	+/-	wild-type	+	+	mutant	-	balanced	+	gain ^{amp}
ZR75-1	wild-type	+/-	wild-type	+	+	wild-type	+	balanced	+	gain ^{amp}
MDA-MB-175VII	wild-type	+	wild-type	+	+	wild-type	+	gain	+/-	balanced
MPE600	wild-type	+	wild-type	+	+	wild-type	+	balanced	+	balanced

Breast cancer cell lines are organized by their gene mutation status. Protein and transcript expression: -, no detectable expression; +/-, barely detectable; +, expression at apparently normal level; ++, overexpression. Mutant AA, missense mutant; mutant TR, truncating mutant; *, heterozygous oncogenic mutation in a tumor suppressor gene; ?, missense mutation of unclear significance; amp, amplified; nd, not determined.

Table 2. Mutation Analysis of RB Pathway Genes.

Breast cancer cell lines	Allelic loss at 13q	<i>RB1</i> gene sequence	<i>RB1</i> transcript expression	<i>RB1</i> protein expression	Allelic loss at 9p	<i>p16</i> gene sequence	<i>p16</i> transcript expression	<i>p16</i> protein expression	<i>Cyclin D1</i> gene locus	<i>Cyclin D1</i> transcript expression
DU4475	yes	deleted	-	-	no	wild-type	+	-	balanced	+/-
MDA-MB-468	yes	deleted	-	-	no	wild-type	++	++	balanced	+/-
BT549	yes	deleted	-	-	no	wild-type	++	++	balanced	+/-
MDA-MB-436	yes	mutant	+/-	-	yes	wild-type	++	++	balanced	+/-
HCC1937	yes	deleted	+	-	nd	wild-type	++	++	balanced	+
BT20	yes	mutant?	+	+	yes	deleted	-	-	balanced	+
MCF-7	yes	wild-type	+	+	yes	deleted	-	-	balanced	+
SUM229PE	yes	wild-type	+	+	yes	deleted	-	-	balanced	+
MDA-MB-231	yes	wild-type	+	++	yes	deleted	-	-	balanced	+
Hs578T	yes	wild-type	+/-	+	yes	deleted	-	-	balanced	+
SUM149PT	yes	wild-type	+/-	+	yes	deleted	-	-	balanced	+
SUM1315MO2	no	wild-type	+	+	yes	deleted	-	-	balanced	+
SK-BR-7	no	wild-type	+	+	yes	deleted	-	-	balanced	+
SUM102PT	no	wild-type	+	nd	yes	deleted	-	nd	balanced	++
MDA-MB-435s	yes	wild-type	+	+	yes	mutant	++	+	balanced	+/-
SUM52PE	yes	wild-type	+	+	yes	mutant	+	+	balanced	+
MDA-MB-361	yes	wild-type	+	+	yes	mutant	+	+/-	balanced	+
OCUB-F	no	wild-type	+	nd	yes	mutant	+	nd	balanced	+
UACC812	yes	wild-type	+	+	no	methylated	+/-	+	balanced	+
UACC893	yes	wild-type	+/-	+	no	methylated	+/-	+	balanced	+
T47D	yes	wild-type	+	+	yes	methylated	-	-	balanced	+
ZR75-1	yes	wild-type	+	+	no	methylated	+/-	+	gain ^{amp}	++
MPE600	no	wild-type	+	+	yes	methylated	+/-	+	gain ^{amp}	++
SUM44PE	nd	nd	+	+	no	methylated	+/-	+/-	gain ^{amp}	++
MDA-MB-415	no	nd	++	+	no	mutant*	++	+/-	gain ^{amp}	++
CAMA-1	no	wild-type	+	+	no	wild-type	+	+	gain ^{amp}	++
MDA-MB-134VI	no	wild-type	+	+	no	wild-type	+	-	gain ^{amp}	++
MDA-MB-330	nd	nd	+	+	no	wild-type	+	-	gain ^{amp}	++
BT474	yes	wild-type	+	+	yes	wild-type	+	+	gain	+
MDA-MB-175VII	no	wild-type	+	+	no	wild-type	+	+/-	gain	++
SK-BR-3	yes	wild-type	+	+	no	wild-type	+	+	balanced	+
SK-BR-5	yes	wild-type	+	+	yes	wild-type	+	+	nd	+
EVSA-T	yes	wild-type	+	+	no	wild-type	+	+	balanced	+/-
MDA-MB-157	yes	wild-type	+/-	+	yes	wild-type	++	++	balanced	+
ZR75-30	nd	nd	+	+	no	wild-type	+	+	balanced	+
BT483	no	wild-type	+	nd	no	wild-type	+	nd	balanced	+
MDA-MB-453	no	wild-type	+	+	yes	wild-type	+	+/-	balanced	+
SUM185PE	no	wild-type	+	++	nd	wild-type	+/-	+/-	balanced	+
SUM159PT	no	wild-type	+	++	no	wild-type	-	+/-	balanced	+

See legend of Table 1 for explanation.



