## Package 'breastCancerMAINZ'

January 4, 2024

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Type Package
<b>Title</b> Gene expression dataset published by Schmidt et al. [2008] (MAINZ).
<b>Version</b> 1.41.0
Date 2011-02-10
<b>Description</b> Gene expression data from the breast cancer study pub- lished by Schmidt et al. in 2008, provided as an eSet.
<b>biocViews</b> ExperimentData, CancerData, BreastCancerData, MicroarrayData, GEO
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<b>Depends</b> R (>= 2.5.0)
Suggests survcomp, genefu, Biobase
LazyLoad yes
License Artistic-2.0
<pre>URL http://compbio.dfci.harvard.edu/</pre>
git_url https://git.bioconductor.org/packages/breastCancerMAINZ
git_branch devel
git_last_commit 11facf7
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#### Description

This dataset contains the gene expression, annotations and clinical data as published in Schmidt et al. 2008.

#### Usage

data(mainz)

#### Format

ExpressionSet with 22283 features and 200 samples, containing:

- exprs(mainz): Matrix containing gene expressions as measured by Affymetrix hgu133a technology (single-channel, oligonucleotides).
- fData(mainz): AnnotatedDataFrame containing annotations of Affy microarray platform hgu133a.
- pData(mainz): AnnotatedDataFrame containing Clinical information of the breast cancer patients whose tumors were hybridized.
- experimentalData(mainz): MIAME object containing information about the dataset.
- annotation(mainz): Name of the affy chip.

#### Details

This dataset represents the study published by Schmidt et al. 2008.

• Abstract: Estrogen receptor (ER) expression and proliferative activity are established prognostic factors in breast cancer. In a search for additional prognostic motifs, we analyzed the gene expression patterns of 200 tumors of patients who were not treated by systemic therapy after surgery using a discovery approach. After performing hierarchical cluster analysis, we identified coregulated genes related to the biological process of proliferation, steroid hormone receptor expression, as well as B-cell and T-cell infiltration. We calculated metagenes as a surrogate for all genes contained within a particular cluster and visualized the relative expression in relation to time to metastasis with principal component analysis. Distinct patterns led to the hypothesis of a prognostic role of the immune system in tumors with high expression of proliferation-associated genes. In multivariate Cox regression analysis, the proliferation metagene showed a significant association with metastasis-free survival of the whole discovery cohort [hazard ratio (HR), 2.20; 95% confidence interval (95% CI), 1.40-3.46]. The B-cell metagene showed additional independent prognostic information in carcinomas with high proliferative activity (HR, 0.66; 95% CI, 0.46-0.97). A prognostic influence of the B-cell metagene was independently confirmed by multivariate analysis in a first validation cohort enriched for high-grade tumors (n = 286; HR, 0.78; 95% CI, 0.62-0.98) and a second validation cohort enriched for younger patients (n = 302; HR, 0.83; 95% CI, 0.7-0.97). Thus, we

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could show in three cohorts of untreated, node-negative breast cancer patients that the humoral immune system plays a pivotal role in metastasis-free survival of carcinomas of the breast.

#### Source

http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE11121

#### References

Marcus Schmidt and Daniel Boehm and Christian von Toerne and Eric Steiner and Alexander Puhl and Heryk Pilch and Hans-Anton Lehr and Jan G. Hengstler and Hainz Koelbl and Mathias Gehrmann (2008)"The Humoral Immune System Has a Key Prognostic Impact in Node-Negative Breast Cancer", *Cancer Research*, **68**(13):5405-5413

#### Examples

## load Biobase package library(Biobase) ## load the dataset data(mainz) ## show the first 5 rows and columns of the expression data exprs(mainz)[1:5,1:5] ## show the first 6 rows of the phenotype data head(pData(mainz)) ## show first 20 feature names featureNames(mainz)[1:20] ## show the experiment data summary experimentData(mainz) ## show the used platform annotation(mainz) ## show the abstract for this dataset abstract(mainz)

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\* datasets mainz, 2

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