

Supplementary Table S1: Specific analysis roles and blinding status of contributing teams

Order of analysis	Role	Team members	Location/Affiliation	Blinding status
1	Pathological analyses, tissue banking, sample provision	CD, KE, BS, CSo, IS, TF	Local trial units, central pathology	blinded to molecular and clinical study data
2	RNA preparation and sequencing	BMY, BLJ	Avera Cancer Institute	fully blinded
3	Primary RNA-Seq raw data analysis and QC	TM, JB	Avera Cancer Institute	fully blinded
4	Sample coding, dataset assembly and distribution	KW, VN	GBG-Statistics Dept.	unblinded
5	Blinded gene expression analysis, development of statistical analysis plan and SPSS script	TK, UH	Goethe University	fully blinded
6	Correlation with patient data according to predeveloped SPSS script	KW, VN	GBG-Statistics Dept.	unblinded
7	Blinded interpretation of summary results	TK, UH	Goethe University	no patient level data
8	Monitoring and review of results	SL, MU, PAF, FM, VM, BG, CSch, CH, ES, JH, MvM	GBG-Boards	no patient level data

Supplementary Table S2: Pre-defined analytical aims of the study:

Pre-defined analytical aims of the study:	
1.	Concordance of RNA-Seq-derived genomic ER-/PR-status, proliferation, immune signature expression, and molecular subtype with pathology-derived IHC-based ER-/PR-status, histological grading, and tumor-infiltrating-lymphocyte (TIL)-scoring, respectively.
2.	Robustness of the above concordances with regard to sample quality (QC class).
3.	Univariate predictive value for pCR of RNA-Seq-derived molecular subtypes, and signatures for proliferation, stroma, T-cell signature, and hypoxia signature.
4.	Multivariate logistic regression of pCR including the following predictor variables: a) Hormone receptor status, treatment arm (+/- Bev), hypoxia signature, and the interaction between hypoxia signature and treatment arm. b) All predictor variables from (a), with additional clinical variables (age, cT, cN, histological grade) as predictors.

Supplementary Table S3: Comparison of clinical parameters of the complete trial cohort and the RNA-Seq cohorts

Parameter	Category	Total cohort 1540 [#] (100.0%)	RNA-Seq data 289 (100.0%)	P-Value [§]	High Quality RNA-Seq data 221 (100.0%)	P-Value [§]
Age	median	48	46	<0.001	46	0.009
clin. tumor status	T1	261 (17.0%)	45 (15.6%)	0.028	37 (16.8%)	0.035
	T2	883 (57.5%)	188 (65.3%)		145 (65.9%)	
	T3	224 (14.6%)	33 (11.5%)		23 (10.5%)	
	T4a-c	79 (5.1%)	8 (2.8%)		8 (3.6%)	
	T4d	89 (5.8%)	14 (4.9%)		7 (3.2%)	
	missing	4	1		1	
clin. lymph node status	negative	772 (51.0%)	148 (51.6%)	n.s.	115 (52.5%)	n.s.
	LN1-3	668 (44.1%)	124 (43.2%)		93 (42.5%)	
	LN4-9	57 (3.8%)	11 (3.8%)		7 (3.2%)	
	LN>=10	17 (1.1%)	4 (1.4%)		4 (1.8%)	
	missing	26	2		2	
HER2 status	negative	1540 (100.0%)	289 (100.0%)		221 (100.0%)	
Hormone receptor status	negative	558 (36.2%)	133 (46.0%)	<0.001	102 (46.2%)	0.001
	positive	982 (63.8%)	156 (54.0%)		119 (53.8%)	
Histological grade	G1	53 (3.5%)	9 (3.1%)	<0.001	7 (3.2%)	<0.001
	G2	781 (51.0%)	119 (41.5%)		86 (39.3%)	
	G3	697 (45.5%)	159 (55.4%)		126 (57.5%)	
	missing	9	2		2	
Histological subtype	Ductal invasive	1241 (80.8%)	238 (82.4%)	n.s.	179 (81.0%)	n.s.
	Lobular invasive	162 (10.5%)	26 (9.0%)		23 (10.4%)	
	other	133 (8.7%)	25 (8.7%)		19 (8.6%)	
	missing	4	0		0	
Treatment arm	EC-T	743 (48.2%)	150 (51.9%)	n.s.	117 (52.9%)	n.s.
	ECB-TB	797 (51.8%)	139 (48.1%)		104 (47.1%)	
pCR	no	1156 (75.1%)	149 (51.6%)	<0.001	112 (50.7%)	<0.001
	yes	384 (24.9%)	140 (48.4%)		109 (49.3%)	

[§] P-values are the result of Fisher's exact tests for binary variables, of chi-square tests for variables with three or more levels, and of Wilcoxon test for continuous variables, respectively.

[#] 1540 total patients with HER2 negative disease with response data from treatment arms out of the ITT population of 2572 patients from the GeparQuinto trial.

Supplementary Table S4: Gene lists of signatures

Signature	Gene	EntrezID	Signature	Gene	EntrezID	Signature	Gene	EntrezID	Signature	Gene	EntrezID	Signature	Gene	EntrezID	Signature	Gene	EntrezID	Signature	Gene	EntrezID	Signature	Gene	EntrezID	Signature	Gene	EntrezID	
Adipocyte	AASS	10157	Basal	ABC13		ER-related	ABCC8	6833	Hypoxia	CSTB	1476	IFN	CMPK2		Ribosomal	CSDE1	7812	Stroma	ADAM12	8038	T-Cell	B2M		567	T-Cell	B2M	
	ABCA10		Basal	COL27A1		ER-related	AFF3	3899	Hypoxia	NDRG1	10397	IFN	DDX58	23586	Ribosomal	EEF1A1	1915	Stroma	ADAMTS12	81792	T-Cell	CD74		972	T-Cell	CD74	
	ABCA6	23460	Basal	CRYAB	1410	ER-related	AGR2	10551	Hypoxia	VEGFA	7422	IFN	DDX60	55601	Ribosomal	EEF1G	1937	Stroma	BNC2	54796	T-Cell	CIT2A		4261	T-Cell	CIT2A	
	ABCA8	10351	Basal	ELF5	2001	ER-related	AR	367				IFN	DDX60L		Ribosomal	HSP01	3329	Stroma	CACNA1C	775	T-Cell	CYBB		1536	T-Cell	CYBB	
	ABCA9		Basal	FOXK1	2296	ER-related	CA12	771				IFN	EIF2AK2	5610	Ribosomal	LRPPRC	10128	Stroma	COL10A1	1300	T-Cell	DOCK10		55619	T-Cell	DOCK10	
Adipocyte	ADH1B	125	Basal	GABRP	2568	ER-related	CAPN8					IFN	EPSTI1		Ribosomal	NACA	4666	Stroma	COL11A1	1301	T-Cell	DOCK2		1794	T-Cell	DOCK2	
Adipocyte	ANK2	287	Basal	KRT23	25984	ER-related	CCDC170	80129				IFN	HERC5	51191	Ribosomal	PABPC1	26986	Stroma	COL5A3	50509	T-Cell	DOCK8			T-Cell	DOCK8	
Adipocyte	APOD	347	Basal	PROM1	8842	ER-related	CYP2B7P1					IFN	HERC6	55008	Stroma	PRKDC	5591	Stroma	COL8A1		T-Cell	HLA-DPA1		3113	T-Cell	HLA-DPA1	
Adipocyte	COL14A1	7373	Basal	RGMA		ER-related	ERBB4	2066				IFN	IFI44	10561	Ribosomal	RPL10	6134	Stroma	COMP	1311	T-Cell	HLA-DRA		3122	T-Cell	HLA-DRA	
Adipocyte	FAT4	79633	Basal	SFRP1	6422	ER-related	ESR1	2099				IFN	IFI44L	10964	Ribosomal	RPL11	6135	Stroma	CERS1D2	83716	T-Cell	HLA-DRB1		3123	T-Cell	HLA-DRB1	
Adipocyte	FREM1		Basal	SLC34A2	10568	ER-related	FOXA1	3169				IFN	IFIH1	64135	Ribosomal	RPL12	6136	Stroma	CTGF	1490	T-Cell	IL10RA		3587	T-Cell	IL10RA	
Adipocyte	IGF1	3479	Basal	SLC6A14	11254	ER-related	FSIP1					IFN	IFIT1	3434	Ribosomal	RPL19	6143	Stroma	DCHS1	8642	T-Cell	NCKAP1L		3071	T-Cell	NCKAP1L	
Adipocyte	LAMA2	3908	Basal	SOX10	6663	ER-related	GATA3	2625				IFN	IFIT2	3433	Ribosomal	RPL23A		Stroma	DPVSL3	1809	T-Cell	PARP14			T-Cell	PARP14	
Adipocyte	PLEKHH2		Basal	TCF7L1	83439	ER-related	GFRA1	2674				IFN	MX1	4599	Ribosomal	RPL27A	6157	Stroma	FAP	2191	T-Cell	PARP9			T-Cell	PARP9	
Adipocyte	SPARCL1	8404	Basal	ZNF462		ER-related	GREB1	9687				IFN	MX2	4600	Ribosomal	RPL3	6122	Stroma	FBLN2	2199	T-Cell	PTPRC		5788	T-Cell	PTPRC	
Adipocyte	SVEP1	79987	Basal	DSC3	1825	ER-related	GRPR	2925				IFN	OAS1	4938	Ribosomal	RPL32	6161	Stroma	FLNC	2318	T-Cell	SAMD9		54809	T-Cell	SAMD9	
Adipocyte	TNXB	7148	Basal	DSG3	1830	ER-related	KDM4B	23030				IFN	OAS2	4939	Ribosomal	RPL4	6124	Stroma	FNDC1		T-Cell	SAMD9L			T-Cell	SAMD9L	
			Basal	FAT2	2196	ER-related	MAPT	4137				IFN	OAS3	4940	Ribosomal	RPL5	6125	Stroma	GAS6	2621	T-Cell	SAMHD1		25939	T-Cell	SAMHD1	
			Basal	KRT14	3861	ER-related	MLPH	79083				IFN	PARP12	64761	Ribosomal	RPL9	6133	Stroma	GAS7	8522	T-Cell	STAT1		6772	T-Cell	STAT1	
			Basal	KRT16		ER-related	NAT1	9				IFN	PLSCR1	5359	Ribosomal	RPLP0	6175	Stroma	GLIS2		T-Cell	STAT2		6773	T-Cell	STAT2	
			Basal	KRT17	3872	ER-related	PGR	5241				IFN	RSAD2	91543	Ribosomal	RPS11	6205	Stroma	IGFBP4	3487	T-Cell	TRIM22		10346	T-Cell	TRIM22	
			Basal	KRT5	3852	ER-related	SCUBE2	57758				IFN	SP100	6672	Ribosomal	RPS18	6222	Stroma	ITGA11		T-Cell	XRN1			T-Cell	XRN1	
			Basal	KRT6A		ER-related	SLC44A4	80736				IFN	SP110	3431	Ribosomal	RPS20	6224	Stroma	ITGA5	3678							
			Basal	KRT6B	3854	ER-related	TBC1D9	23158				IFN	UBA7	7318	Ribosomal	RPS24	6229	Stroma	ITGB5	3693							
			Basal	TRIM29	23650	ER-related	THSD4	79875				IFN	XAF1	54739	Ribosomal	RPS27A	6233	Stroma	ITGBL1	9358							
												IFN	ZNFX1		Ribosomal	RPS4X	6191	Stroma	KANK2	25959							
															Ribosomal	RPS6	6194	Stroma	KIAA1462	57608							
															Ribosomal	RPS7	6201	Stroma	KIF26B	55083							
															Ribosomal	RPSA		Stroma	LMOD1	25802							
																		Stroma	LRRC15	131578							
																		Stroma	MAP1A	4130							
																		Stroma	MICAL2	9645							
																		Stroma	MMP11	4320							
																		Stroma	MMP14	4323							
																		Stroma	MRC2	9902							
																		Stroma	MXRA8	54587							
																		Stroma	MYH1	4619							
																		Stroma	MYL9	10398							
																		Stroma	NID1	4811							
																		Stroma	NID2	22795							
																		Stroma	PCOLCE	5118							
																		Stroma	PDLIM7	9260							
																		Stroma	PHLD81	23187							
																		Stroma	PMEPA1	56937							
																		Stroma	PMP22	5376							
																		Stroma	PODN								
																		Stroma	RIN2	54453							
																		Stroma	SFRP2								
																		Stroma	SFRP4	6424							
																		Stroma	SPON1	10418							
																		Stroma	SSCSO								
																		Stroma	TAGLN	6876							
																		Stroma	TENM4	26011							
																		Stroma	THY1	7070							
																		Stroma	TIMP2	7077							
																		Stroma	TIMP3	7078							
																		Stroma	UNC5B	219699							
																		Stroma	ZFHX4	79776							

Supplementary Table S4: Gene lists of signatures

PROVIDED AS EXCEL FILE: *Suppl_Table_S4_SignatureList.xlsx*

Supplementary Table S5: Core genes of the hypoxia signature cluster from different datasets with correlated expression

Gene symbol	Gene name	Category	Details	FFPE-RNA-Seq data
VEGFA	Vascular endothelial growth factor	Angiogenesis	Growth factor active in angiogenesis, vasculogenesis and endothelial cell growth. Induces endothelial cell proliferation, promotes cell migration, inhibits apoptosis and induces permeabilization of blood vessels. Binds to the FLT1/VEGFR1 and KDR/VEGFR2 receptors, heparan sulfate and heparin.	yes
NDRG1	N-myc downstream regulated gene 1	Stress response	Involved in stress responses, hormone responses, cell growth, and differentiation. Necessary for p53-mediated caspase activation and apoptosis.	yes
ANGPTL4	Angiopoietin-like 4	Angiogenesis, hypoxia	Hypoxia-induced expression in endothelial cells. May act as a regulator of angiogenesis and modulate tumorigenesis. In response to hypoxia, the unprocessed form of the protein accumulates in the subendothelial extracellular matrix.	no
ADM	Adrenomedullin	Angiogenesis	Adrenomedullin functions include vasodilation, regulation of hormone secretion, and promotion of angiogenesis.	no
DDIT4	DNA damage induced transcript 4	Stress response	Regulates cell growth, proliferation and survival via inhibition of mTORC1. Important role in responses to cellular energy levels and cellular stress, including responses to hypoxia and DNA damage.	no
CSTB	Cystatin-B	Proteinase inhibitor	Intracellular thiol proteinase inhibitor thought to play a role in protecting against proteases leaking from lysosomes.	yes

Supplementary Table S6: Accuracy of IHC detection of NDRG1 as marker of the hypoxia signature

Cohort	Finding (N=23)	Finding (N=23)	Finding (N=23)	Full (N=193)
Source	Whole slide	Whole slide	TMA	TMA
Method	pathological scoring	digital image analysis	digital image analysis	digital image analysis
Cutoff	pos/neg	>10% positive cells	>10% positive cells	>10% positive cells
Positive by RNA-Seq*	19	19	19	20
Negative by RNA-Seq*	4	4	4	173
Accuracy	91.3 %	73.9 %	47.8 %	81.9 %
Sensitivity	89.5 %	68.4 %	36.8 %	40.0 %
Specificity	100.0 %	100.0 %	100.0 %	86.7 %
PPV	100.0 %	100.0 %	100.0 %	25.8 %
NPV	66.7 %	40.0 %	25.0 %	92.6 %

* based on cutoff z-score 1.5 from RNA-Seq

Supplementary Table S7: Comparison of hormone receptor status from RNA-Seq and IHC

High Quality samples	Sensitivity	Specificity	PPV	NPV	Accuracy
ER _{RNA-Seq} vs. ER _{IHC} (N=221)	75.7 %	93.4 %	92.6 %	78.0 %	84.2 %
PR _{RNA-Seq} vs. PR _{IHC} (N=221)	76.6 %	83.5 %	77.4 %	82.8 %	80.5 %
Low Quality samples	Sensitivity	Specificity	PPV	NPV	Accuracy
ER _{RNA-Seq} vs. ER _{IHC} (N=68)	72.7 %	88.6 %	85.7 %	77.5 %	80.9 %
PR _{RNA-Seq} vs. PR _{IHC} (N=67)	61.3 %	86.1 %	79.2 %	72.1 %	74.6 %

Supplementary Table S8: Comparison of Molecular Subtyping between High Quality and Low Quality Samples

Group	Basal-like	Her2-enrich.	LumA	LumB	Normal-like
Total (N=289)	119 (41.2%)	60 (20.8%)	33 (11.4%)	46 (15.9%)	31 (10.7%)
HQ no dupl. (N=221, %)	103 (46.6%)	33 (14.9%)	28 (12.7%)	42 (19.0%)	15 (6.8%)
LQ no dupl. (N=68, %)	16 (23.5%)	27 (39.7%)	5 (7.4%)	4 (5.9%)	16 (23.5%)

The distributions of molecular subtypes according to AIMS differ significantly ($P = 8.6 \times 10^{-9}$, Fisher's Exact Test) between samples with high and low quality.

Supplementary Table S9: Univariate logistic regression of pCR by molecular markers (N=221 High Quality samples)

Molecular marker	OR	95% CI	P-value
Basal-like*	8.88	2.34-33.6	0.001
HER2-enriched*	3.33	0.79-14.1	0.10
Lum-A*	0.87	0.18-4.28	0.86
Lum-B*	2.22	0.54-9.14	0.27
T-cell signature [#]	1.60	1.21-2.12	0.001
Proliferation signature [#]	2.88	2.00-4.16	<0.001
Hypoxia signature [#]	1.92	1.41-2.60	<0.001

* vs. Normal-like subtype, [#] z-score

Supplementary Table S10: Multivariate logistic regression of pCR with NDRG1 from TMA analysis

	OR	95% CI	P-value
Hormone receptor (neg. vs. pos.)	4.35	2.34-8.07	<0.001
NDRG1-TMA-IHC (z-score >1.5)	3.79	0.85-16.9	0.080
Bevacizumab therapy	0.92	0.53-1.60	0.766
Interaction NDRG1-TMA-IHC * bevacizumab	1.31	0.78-2.21	0.309
cN (≥10 vs 4-9 vs 1-3 vs 0 positive nodes)	0.80	0.52-1.24	0.323
cT (T4d vs T4a-c vs T3 vs T2 vs T1)	0.80	0.59-1.0	0.173
Grading (G3 vs G2 vs G1)	1.43	0.83-2.48	0.197