

VARIANT[™]Plex Comprehensive Tissue & Blood

Description

The VARIANT[™]Plex Comprehensive Tissue & Blood panel is a balanced pool of gene-specific primer (GSP) oligonucleotides that is optimized for use with VARIANT[™]Plex reagents and molecular barcode (MBC) adapters to produce targeted NGS libraries. This product insert should be used in conjunction with VARIANT[™]Plex HS/HGC protocol for Illumina® (RA-DOC-056) or VARIANT[™]Plex-LAC protocol for Illumina® (RA-DOC-470).

VARIANT[™]Plex Comprehensive Tissue & Blood contains **26,535** GSPs targeting **460** genes commonly mutated in solid tumor and blood cancer types as well as microsatellite instability (**MSI**), tumor mutational burden (**TMB**), and homologous recombination deficiency (**HRD**) status.

Description	Part number	Storage
VARIANT [™] Plex Comprehensive Tissue & Blood GSP1 - 8 reactions	SA24315081	-20°C ± 10°C
VARIANT [™] Plex Comprehensive Tissue & Blood GSP2 - 8 reactions	SA24315082	

Required reagent volumes

Protocol reference	Protocol step	Reagent	HS/HGC RA-DOC-056 Volume per reaction (µL)	LAC RA-DOC-470 Volume per reaction (µL)
A	Ligation Step 2 Elution	5mM NaOH	32	32
B	First PCR	VARIANT [™] Plex Comprehensive Tissue & Blood GSP1	8	8
C	First PCR	10mM Tris-HCl pH 8.0	34	30
D	First PCR	Purified PCR1 eluate	32	28
E	Second PCR	VARIANT [™] Plex Comprehensive Tissue & Blood GSP2	8	8

Recommended PCR cycling

	Step	Temperature (°C)	Time	Cycles
First PCR reaction	1	95	3 min	1
	2	95	30 sec	
	3	58	10 sec	15
	4	60	10 min (100% ramp rate)	
	5	72	3 min	1
	6	4	Hold	1
Second PCR reaction	1	95	3 min	1
	2	95	30 sec	
	3	58	10 sec	20 [†]
	4	65	10 min (100% ramp rate)	
	5	72	3 min	1
	6	4	Hold	1

[†]The number of PCR2 cycles may be decreased if you regularly experience library yields greater than 200 nM.

Recommended reads and multiplexing

VARIANT[™]Plex Comprehensive Tissue & Blood libraries should be sequenced to a minimum of **61M** reads. Starting read depth recommendations for standard profiling may be adjusted based on user needs.

Archer™ Analysis settings

Sequencing data should be processed using Archer Analysis (v7.4, or greater). The VARIANT[™]Plex Comprehensive Tissue & Blood panel is compatible with the **SNV/Indel, Copy Number Variation, CNV 2.0, ASCN, Structural Variation, MSI, TMB, and HRD** pipelines, found under the **DNA** Input Type. The recommended TMB settings in Archer Analysis are *High Threshold for TMB = 17.5* and *Low Threshold for TMB = 6* for VARIANT[™]Plex Comprehensive Tissue & Blood. As with most Analysis settings, these may be adjusted

based on user needs. Selection of the DNA Target Coverage pipeline is optional (see the Archer Analysis User Guide for more details on setting up your analysis).

Archer Analysis processing of VARIANTPlex Comprehensive Tissue & Blood libraries requires a one-time upload of the Panel GTF. When performing DNA Target Coverage analysis, users must also select a Region of Interest BED file. Users may optionally add a Targeted Mutations VCF file for targeted SNV/Indel detection. Files can be obtained by contacting archer-tech@idtdna.com

Assay targets

Integrative genomic biomarkers

MSI, TMB, HRD

Genes

The following genes in the panel have full coding sequence coverage for SNV/Indel analysis. Coding sequence was determined using NCBI RefSeq transcripts[†]. These genes are also enabled for Copy Number Variation (CNV) analysis.

<i>ABL1</i>	<i>CCND3</i>	<i>EPCAM</i>	<i>FOXQ1</i>	<i>KIT*</i>	<i>NBN</i>	<i>PPP2R2A</i>	<i>SF3B1</i>	<i>TXNIP</i>
<i>ABL2</i>	<i>CCNE1</i>	<i>EPHA2</i>	<i>FUBP1</i>	<i>KLF4</i>	<i>NCOR1</i>	<i>PPP6C</i>	<i>SH2D1A</i>	<i>U2AF1</i>
<i>ACVR1</i>	<i>CD70</i>	<i>EPHA3</i>	<i>GATA1</i>	<i>KLF5</i>	<i>NF1</i>	<i>PRDM1</i>	<i>SLX4</i>	<i>UNCX</i>
<i>ACVR2A</i>	<i>CD79A</i>	<i>EPHA7</i>	<i>GATA2</i>	<i>KMT2A (MLL)</i>	<i>NF2</i>	<i>PRKAR1A</i>	<i>SMAD2</i>	<i>USP9X</i>
<i>AKT1</i>	<i>CD79B</i>	<i>EPHB1</i>	<i>GATA3</i>	<i>KMT2B</i>	<i>NFE2L2</i>	<i>PRKD1</i>	<i>SMAD3</i>	<i>VHL</i>
<i>AKT2</i>	<i>CD274 (PD-L1)</i>	<i>ERBB2</i>	<i>GEN1</i>	<i>KMT2C</i>	<i>NFKBIA</i>	<i>PRKDC</i>	<i>SMAD4</i>	<i>WT1</i>
<i>AKT3</i>	<i>CDC73</i>	<i>ERBB3</i>	<i>GLI1</i>	<i>KMT2D (MLL2)</i>	<i>NIPBL</i>	<i>PSIP1</i>	<i>SMARCA4</i>	<i>XPO1</i>
<i>ALK</i>	<i>CDH1</i>	<i>ERBB4</i>	<i>GNA11</i>	<i>KRAS</i>	<i>NKX2-1</i>	<i>PTCH1</i>	<i>SMARCB1</i>	<i>XRCC2</i>
<i>APC</i>	<i>CDK4</i>	<i>ERCC1</i>	<i>GNA13</i>	<i>KRT222</i>	<i>NOTCH1*</i>	<i>PTEN</i>	<i>SMC1A</i>	<i>XRCC3</i>
<i>AR</i>	<i>CDK6</i>	<i>ERCC2</i>	<i>GNAQ</i>	<i>LAMP1</i>	<i>NOTCH2</i>	<i>PTMA</i>	<i>SMC3</i>	<i>ZBTB20</i>
<i>ARAF</i>	<i>CDK8</i>	<i>ERCC4</i>	<i>GNAS</i>	<i>LATS1</i>	<i>NOTCH3</i>	<i>PTPDC1</i>	<i>SMO</i>	<i>ZFHX3</i>
<i>ARHGAP35</i>	<i>CDK12</i>	<i>ERG</i>	<i>GPS2</i>	<i>LATS2</i>	<i>NOTCH4</i>	<i>PTPN11</i>	<i>SOCS1</i>	<i>ZMYM2</i>
<i>ARID1A</i>	<i>CDKN1A</i>	<i>ERRFI1</i>	<i>GRIN2D</i>	<i>LEMD2</i>	<i>NPM1</i>	<i>PTPRC</i>	<i>SOX2</i>	<i>ZMYM3</i>
<i>ARID1B</i>	<i>CDKN1B</i>	<i>ESR1</i>	<i>GRM3</i>	<i>LRP1B</i>	<i>NRAS</i>	<i>PTPRD</i>	<i>SOX9</i>	<i>ZNF750</i>
<i>ARID2</i>	<i>CDKN2A</i>	<i>EWSR1</i>	<i>H3F3A</i>	<i>LZTR1</i>	<i>NSD1</i>	<i>PTPRT</i>	<i>SOX17</i>	<i>ANKRD26</i>

ARID5B	CDKN2B	EZH2	H3F3B	MACF1	NTRK1	RAC1	SPEN	CALR
ASXL1	CDKN2C	FAM46C	H3F3C	MAP2K1 (MEK1)	NTRK2	RAD21	SPOP	CBLB
ASXL2	CEBPA	FAM46D	HGF	MAP2K2 (MEK2)	NTRK3	RAD50	SPTA1	CBLC
ATM*	CHD1	FAM175A	HIST1H1C	MAP2K4	NUP93	RAD51	SPTAN1	CD28
ATR	CHD3	FANCA	HIST1H1E	MAP3K1	PALB2	RAD51B	SRC	CDC25C
ATRX	CHD4	FANCC	HIST1H2BD	MAP3K4	PARP1	RAD51C	SRSF2	CUX1
AURKA	CHD8	FANCD2	HIST1H3B	MAP3K13	PAX5	RAD51D	STAG1	CXCR4
AURKB	CHEK1	FANCE	HIST1H3C	MAPK1	PAX8	RAD52	STAG2	DCK
AXIN1	CHEK2*	FANCF	HNF1A	MAX	PBRM1	RAD54L	STAT3	DDX41
AXIN2	CIC	FANCG	HOXB13	MCL1	PDCD1 (PD-1)	RAF1	STK11	DHX15
AXL	COL5A1	FANCI	HRAS	MDM2	PDCD1LG2 (PD-L2)	RARA	SUFU	ETNK1
B2M	CREBBP	FANCL	HUWE1	MDM4	PDGFRA*	RASA1	SYK	ETV6
BAP1	CRKL	FAT1	IDH1	MECOM	PDGFRB	RB1	TAF1	ID3
BARD1	CSDE1	FBXW7	IDH2	MED12	PGR	RBM10	TBL1XR1	KLF2
BCL2	CSF1R	FGF1	IGF1R	MEF2B	PHF6	RECQL4	TBX3	LUC7L2
BCL2L1	CSF3R	FGF2	IKBKE	MEN1	PIK3C2B	REL	TCEB1	NFKBIE
BCL2L11	CTCF	FGF3	IKZF1	MET*	PIK3C2G	RET*	TCF3	PLCG2
BCL6	CTLA4	FGF4	IL6ST	MGA	PIK3CA	RHEB	TCF7L2	RBBP6
BCOR	CTNNA1	FGF7	IL7R	MGMT	PIK3CB	RHOA	TERT	RPS14
BCORL1	CTNNB1*	FGF8	INPP4B	MITF	PIK3CD	RICTOR	TET1	SAMD9
BIRC3	CUL3	FGF9	INPPL1	MLH1	PIK3CG	RIT1	TET2	SAMD9L
BLM	CYSLTR2	FGF19	IRF2	MLL2	PIK3R1*	RNF43	TFRC	SH2B3
BMPR1A	DAXX	FGFR1	IRF4	MPL	PIK3R2	ROS1	TGFBR1	SLC29A1
BRAF*	DDR2	FGFR2	IRF6	MRE11A	PIK3R3	RPL5	TGFBR2	STAT5B
BRCA1*	DDX3X	FGFR3	IRS2	MSH2	PIM1	RPS6KB1	TGIF1	STAT6
BRCA2*	DICER1	FGFR4	JAK1	MSH3	PLCB4	RPTOR	THRAP3	TRAF2
BRIP1	DMD	FH	JAK2	MSH6	PLCG1	RRAS2	TLR4	U2AF2
BTG1	DNMT3A	FLCN	JAK3	MST1R	PLXNB2	RUNX1	TMSB4X	UBA1
BTG2	DOT1L	FLNA	JUN	MTOR	PMS1	RUNX1T1	TNFAIP3	ZRSR2
BTK	EEF1A1	FLT1	KANSL1	MUC6	PMS2	RXRA	TNFRSF14	
CACNA1A	EEF2	FLT3	KDM5A	MUC16	POLD1	SCAF4	TP53	
CARD11	EGFR	FLT4	KDM5C	MUTYH	POLE	SDHA	TP63	

<i>CASP8</i>	<i>EGR3</i>	<i>FOXA1</i>	<i>KDM6A</i>	<i>MYC</i>	<i>POLQ</i>	<i>SDHB</i>	<i>TRAF3</i>
<i>CBFB</i>	<i>EIF1AX</i>	<i>FOXA2</i>	<i>KDR</i>	<i>MYCL</i> (<i>MYCL1</i>)	<i>POLRMT</i>	<i>SDHC</i>	<i>TRAF7</i>
<i>CBL</i>	<i>ELF3</i>	<i>FOXL2</i>	<i>KEAP1</i>	<i>MYCN</i>	<i>PPARG</i>	<i>SDHD</i>	<i>TSC1</i>
<i>CCND1</i>	<i>EP300</i>	<i>FOXO1</i>	<i>KEL</i>	<i>MYD88</i>	<i>PPM1D</i>	<i>SETBP1</i>	<i>TSC2</i>
<i>CCND2</i>	<i>EPAS1</i>	<i>FOXP1</i>	<i>KIF1A</i>	<i>MYH9</i>	<i>PPP2R1A</i>	<i>SETD2</i>	<i>TSHR</i>

†Contact archer-tech@idtdna.com for the panel target file to view complete list of targeted regions.

*Indicates that select regions of this gene are enabled for Structural Variant analysis in the GTF.

SNP sites targeted for sample tracking

rs560681	rs430046	rs987640	rs10776839	rs12393891
rs740598	rs8078417	rs6444724	rs6530357	chrX:4429309
rs1498553	rs9951171	rs6811238	rs5971553	chrX:11314433
rs10773760	rs576261	rs13182883	rs5953060	chrY:6738552
rs1058083	rs1109037	rs214955	rs6524626	chrY:19490214
rs4530059	rs1523537	rs321198	rs5940270	
rs1821380	rs221956	rs4606077	rs722847	

SNPs may be used in combination to uniquely tag and track samples over time. Contact archer-tech@idtdna.com for further details.

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Revision History

Document Number	Date	Description of change
RA-DOC-480/REV01	August 2024	Initial release.