



## Molecular In My Pocket...

# Solid Tumors

Prepared by the Association for Molecular Pathology Training and Education Committee

## Abbreviations

AMP	Anchored Multiplex PCR	ISH	In Situ Hybridization	Multiple Endocrine Neoplasia	NGS	Next-Generation Sequencing
CA	Cancer	FISH	Fluorescence ISH	Loss of Function	SGA	Single Gene Assay
CGX	Cytogenetics	GI	Gastrointestinal	Mismatch Repair	TKI	Tyrosine Kinase Inhibitor
IHC	Immunohistochemistry	HBOC	Hereditary Breast and Ovarian CA Syndrome	MSI	Microsatellite Instability	

Tissue Type	Disease	Gene	Common Genomic Variant(s)	Significance	Primary Assays
<b>Adrenal</b>	Hereditary Paranglioma / Pheochromocytoma	SDHA, SDHB, SDHC, SDHD	LOF		IHC, SGAs, NGS
	Neuroblastoma	MYCN	Amplification	prognosis	FISH
		IDH1 / IDH2	IDH1 R132, IDH2 R172, R140		IHC, SGAs, NGS
<b>Brain</b>	Glioma / Glioblastoma	TERT	promoter mutations	prognosis	SGAs, NGS
		1p / 19q codeletion		oligodendroglioma phenotype	FISH, CGX, NGS
		ATRX	LOF		IHC, SGAs, NGS
		MGMT	Promoter methylation	response to alkylating agents	Methylation assay
		EGFR	amplification; vlll transcript		FISH, CGX, SGAs, NGS
	Pediatric Glioma	H3F3A, HIST1H3B	K27		
	Medulloblastoma	WNT pathway genes SHH pathway genes	CTNNB1 PTCH1 LOF		SGAs, NGS SGAs, NGS
<b>Breast</b>	Breast CA	CTNNB1 (Beta-Catenin)	Exon 3 mutations	aberrant nuclear expression	IHC, SGAs, NGS
		Estrogen Receptor (ER or ESR1)		response to anti-estrogen	IHC, SGAs, NGS
		Progesterone Receptor (PR or PGR)		response to anti-estrogen	IHC, SGAs, NGS
		ERBB2 (HER2)	Amplification / Overexpression; S310; L755	responsiveness to trastuzumab	FISH, IHC
		Breast CA (Lobular)	CDH1 (E-Cadherin)	LOF	IHC, SGAs, NGS
<b>Cervix</b>	HBOC / Sporadic CA	BRCA1 / BRCA2	LOF	Risk of hereditary cancer; response to PARP inhibitors	SGAs, NGS
	Squamous cell CA	High-risk HPV			IHC (p16), ISH, Viral Typing
	Endometrial CA (sporadic)	MLH1	LOF and promoter hypermethylation	somatic etiology (not Lynch Syndrome)	promoter methylation
<b>Endometrium</b>	Lynch Syndrome	MLH1, MSH2, MSH6, PMS2	LOF	40-60% lifetime risk of endometrial cancer	IHC, SGAs, NGS, MSI
	Cowden Syndrome	PTEN (germline)	LOF	25% risk of endometrial cancer development	SGA, NGS
<b>Eye</b>	Retinoblastoma	RB1	LOF		SGAs, NGS
	Uveal Melanoma	GNAQ, GNA11	GNAQ / GNA11 Q209L		SGAs, NGS
<b>GI Tract</b>	Gastrointestinal Stromal Tumor (GIST)	exon 9 variants		Responsiveness to TKIs, associated with 1' resistance	
		exon 11 in-frame indels; D820; less frequently exons 8, 17		Responsiveness to TKIs	SGAs, NGS
		exon 14		Resistance to first-line TKIs	**IHC not a proxy**
		exon 13		Responsiveness to TKIs as 1' variant; as 2' variant confers resistance to TKIs	
		exon 12, 14, 18 variants		Responsiveness to TKIs	SGAs, NGS
		PDGFRA	exon 18 D842V	Resistance to TKIs	
		SDHA, SDHB, SDHC, SDHD	LOF	Resistance to TKIs	SGAs, NGS, IHC for SDHB
		KRAS, NRAS	G12; G13; Q61	decreased response to panitumumab or cetuximab	SGAs, NGS
		BRAF (therapeutic setting)	exon 15, V600E	decreased response to panitumumab or cetuximab	SGAs, NGS
		Colorectal CA	MSI testing	MSI-high	MMR IHC, MSI
<b>GI Tract (Lower)</b>	Colorectal CA (Lynch Syndrome)	Most commonly MLH1, MSH2, MSH6, PMS2, EPCAM	LOF	MSI-high better prognosis in general, do not respond well to 5-FU based therapies; response to immuno-therapy	MSI, MMR IHC, NGS
		MLH1	promoter methylation	somatic etiology (not Lynch Syndrome)	methylation studies
		BRAF	exon 15, V600E	somatic etiology when MLH1 deficient/MSI-high	
		Colorectal CA (FAP)	APC	LOF	SGAs, NGS
<b>Heart</b>	Hirschprung disease	RET		SGAs, NGS	
	Cardiomyopathy	MYH7		SGAs, NGS	
	Birt-Hogg-Dube syndrome	FLCN	LOF	Both dilated and hypertrophic	SGAs, NGS
<b>Kidney</b>	Clear Cell Renal CA	VHL	3p deletion		SGAs, NGS CGX
	Polycystic Kidney Disease (Dominant / Recessive)	PKD1 / PKHD1	LOF		SGAs, NGS

<b>Kidney (continued)</b>	Hereditary leiomyomatosis and Renal Cell Carcinoma Syndrome	FH	LOF		Adverse Prognosis	IHC, SGAs, NGS
	Renal CA; Von Hippel Lindau Syndrome	VHL	LOF			SGAs, NGS
	Rhabdoid Sarcoma	SMARCB1 (INI)	LOF			IHC, NGS
	Translocation-Associated Renal CA	TFE3 rearrangement	translocations involving Xp11			FISH, CGX
	Wilms' Tumor	WT1				SGAs, NGS
<b>Liver</b>	Alagille Syndrome	JAG1	LOF	haploinsufficiency		SGAs, NGS
	Hepatic Adenoma	CTNNB1 (Beta-Catenin)	Exon 3 mutations	aberrant nuclear expression		IHC, SGAs, NGS
	Hereditary Hemochromatosis	HFE	C282Y, H63D			SGAs, NGS
	Wilson Disease	ATP7B				SGAs
		BRAF	V600E	Predicts response to BRAF inhibitors (dabrafenib-trametinib)		SGAs, NGS
<b>Lung</b>	Non-Small Cell CA (NSCLC, adenocarcinoma)	ROS1 rearrangement	Multiple fusion partners	Responsiveness to TKIs		FISH
		ALK rearrangement	EML4-ALK; inv(2) exons 18-21; L858R	Responsiveness to TKIs		FISH, IHC, NGS
		EGFR	exon 20; T790M	Resistance to first line therapy; de novo and acquired		IHC, SGAs, NGS
		SERPINA1		PIMM / PIZZ alleles		SGAs
		PTEN	LOF			SGAs, NGS
<b>Multi-System</b>	Li-Fraumeni syndrome	TP53	LOF			SGAs, NGS
	McCune Albright Syndrome	GNAS				SGAs, NGS
	MEN Types 1 / 2A & 2B / 4	MEN1 / RET / CDKN1B				SGAs, NGS
	Neurofibromatosis	NF1 / NF2	LOF			SGAs, NGS
	Peutz-Jeghers Syndrome	STK11	LOF			SGAs, NGS
<b>Oropharynx</b>	Tuberous Sclerosis	TSC1 / TSC2				SGAs, NGS
	Squamous CA	High-risk HPV	high risk virus, type 16, 18, other	Improved prognosis		IHC (p16), ISH, Viral Typing
		CTNNB1 (Beta-Catenin)	Exon 3 mutations	aberrant nuclear expression		IHC, SGAs, NGS
	Ovarian CA	PIK3CA	Amplification	Responsiveness to PI3K inhibitors		SGAs, NGS
		PTEN	LOF			SGAs, NGS
<b>Ovary</b>	Small cell CA, hypercalcemic type	SMARCA4	LOF			SGAs, NGS
	High grade serous carcinoma	BRCA1 / BRCA2, germline and somatic	SNVs, indels	diagnostic		SGAs, NGS
	Low-grade serous carcinoma	KRAS, BRAF	SNVs	Response to Platinum-based chemotherapy & PARPi		IHC, SGAs, NGS
	HBOC / Sporadic	BRCA1 / BRCA2	LOF	diagnostic, potential response to MEK-inhibitors		SGA, NGS
	Prostate CA	AR	Amplification	Response to PARP inhibitors		SGAs, NGS
<b>Prostate</b>	Merkel Cell Carcinoma	TMPRSS2 rearrangement	TMPRSS2-ERG; t(21;21)			IHC, SGAs, NGS, FISH
		Merkel Cell Polyoma Virus	Virus			FISH, NGS
		BRAF	V600			ISH, Viral Test
		KIT (CD117)	exon 9, 11, 13, 17 variants	Responsiveness to vemurafenib, dabrafenib		IHC, SGAs, NGS
		PDGFRA	exon 12, 14, 18 variants	Responsiveness to TKIs		SGAs, NGS
<b>Skin</b>	Melanoma	NRAS	G12; G13; Q61	Responsiveness to TKIs		SGAs, NGS
		NFI	LOF			SGAs, NGS
		TERT	promoter mutations			SGAs, NGS
	Alveolar Rhabdomyosarcoma	FOXO1 rearrangement	FOXO1-FLI1; t(11;22) (most common)			FISH, NGS, AMP
	Ewing sarcoma	EWSR1 rearrangement	PAX3-FOXO1; t(2;13)			FISH, NGS, AMP
<b>Soft Tissue</b>	Fibrosarcoma	ETV6 rearrangements	ETV6-NTRK3; t(12;15)	Responsiveness to TKIs		FISH, NGS, AMP
	Inflammatory Myofibroblastic Tumor	ALK rearrangement	TPM3-ALK; t(2;5)			FISH, IHC, NGS, AMP
	Solitary Fibrous Tumor	STAT6 rearrangement	NAB2-STAT6; inv(12)			IHC, SGAs, NGS, AMP
	Synovial Sarcoma	SS18 rearrangement	SS18-SSX1; t(X;18)			FISH, NGS, AMP
	Gastric CA	ERBB2 (HER2)	Amplification / Overexpression; S310; L755	Responsiveness to Trastuzumab		FISH, IHC
<b>Stomach</b>	Hereditary Diffuse Gastric CA	CDH1 (E-Cadherin)	LOF			SGAs, NGS
	Thymic carcinoma	KIT (CD117)	exon 11 variants; D820	Responsiveness to TKIs		IHC, SGAs, NGS
<b>Thymus</b>	Follicular Thyroid CA	PAX8-PPARG rearrangement	PAX8-PPARG; t(2;3)			FISH
	Papillary Thyroid CA	RET rearrangement	PTC1, PTC2, PTC3 common partners	Diagnostic		IHC, SGAs, NGS
<b>Thyroid</b>		BRAF	V600			FISH, NGS, AMP
		RET	promoter mutations	Prognosis		SGAs, NGS

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