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STUDY NAME	PHASE	MOLECULE	COHORT TYPE			PREVIOUS LINE
			COHORT NUMBER	TUMOR TYPE	SPECIFICITY	
BGB-58067-101	I	BGB-58067 + Tislelizumab+platinum based chemotherapy	Phase 1b – part C dose expansion and optimization	NSCLC	Evidence of homozygous loss of MTAP gene in ctDNA or tumor tissue,	One line
BI-1479-0009	II	BI 1810631	5	NSCLC	Her2 overexpressed/amplified	
			18	Non squamous NSCLC	Her2 amplified	
EP0031-101	I/II	EP0031-101	2B	NSCLC	RET fusion-positive NSCLC, chemotherapy combination	treatment naive
GO45416	I/II	GDC-7035 (RO778493)	Expansion phase /A3	NSCLC	KRAS G12D mutation	1 line minimum

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STUDY NAME	PHASE	MOLECULE	COHORT TYPE			PREVIOUS LINE
			COHORT NUMBER	TUMOR TYPE	SPECIFICITY	
GSK-223054	I	GSK5764227+ Atezolizumab	Part 1b dose optimization	ES-SCLC		For treatment-naïve participants, no prior systemic therapies in the extensive-stage setting
		GSK5764227	Part 1b dose optimization	ES-SCLC		Disease progression after PD-1/PD-L1 inhibitor + platinum+ etoposide therapy is required for participants treated with first-line treatment in extensive-stage SCLC + patient must have received tarlatamab
		GSK5764227 + Dostarlimab	Part 1b dose optimization	Nsq NSCLC		2 or 3 prior lines of treatment
KN-8701	Ib	KIN-2787	B1	NSCLC	BRAF class II fusion	Received prior locally approved standard of care appropriate for their tumor type and stage of disease, RAFi naïve
RMC-6291	I	RMC6291 + RMC6236	extension	NSCLC	KRAS G12C	Patients who have received no more than 3 lines of treatment
TEADCO	I/II	Sotorasib + ODM-212	Part1 bras C	NSCLC	KRAS G12C	previously treated with a KRASG12C (OFF) inhibitor
UNLOCK ASP 3082	II	ASP3082	1	NSCLC	KRAS G12D	At least 1 line of prior therapy including a platinum

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STUDY NAME	PHASE	MOLECULE	COHORT TYPE			PREVIOUS LINE
			COHORT NUMBER	TUMOR TYPE	SPECIFICITY	
BI-1479-0012	Ib	Trastuzumab + Capecitabine + Zongertinib	G	Metastatic breast cancer	HER2+	
		Zanidatamab + Zongertinib	O	Metastatic breast cancer	HER2+	
	II	T-DXd 120 mg + Zongertinib 240 mg	E	Metastatic breast cancer	HER2+	1 prior line (palliative), Prior taxane + trastuzumab (± pertuzumab) required, No prior T-DXd or small-molecule HER2 inhibitor
		T-DM1 120 mg + Zongertinib 240 mg	D	Metastatic breast cancer	HER2+	Up to 2 prior lines (some patients may have 3), Prior T-DXd required, No prior T-DM1, fluoropyrimidine, or small-molecule HER2 inhibitor
		Zongertinib monotherapy (I) or Zongertinib 180 mg + Trastuzumab 360 mg (J)	I/J	Metastatic breast cancer	HER2+	Up to 5 prior lines, Prior taxane + trastuzumab + T-DXd required, No prohibited pretreatment
DSB2455-001	II	DSB2455	Part B cohort C	Breast cancer	one or more of the following: BRCA1, BRCA2, PALB2, RAD51C, RAD51D	3 lines of treatment maximum

Gynecology

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			COHORT NUMBER	TUMOR TYPE	SPECIFICITY	
BI-1479-0009	II	BI 1810631	4	Cervical cancer	HER2 overexpressed / amplified with or without HER 2 mutations	
			16	Uterine cancer	HER2 amplified	
			17	Cervical cancer	HER 2 amplified	
DSB2455-001	Ia/Ib	DSB2455	Part B cohorte E	Ovarian cancer	One or more alteration in the following genes : BRCA1, BRCA2, PALB2, RAD51C, RAD51D	
PanSOHO 22752	II	BAY 2927088	4 and 5	Cervical cancer	HER 2 activating mutation	No prior HER2 treatment

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STUDY NAME	PHASE	MOLECULE	COHORT TYPE			PREVIOUS LINE
			COHORT NUMBER	TUMOR TYPE	SPECIFICITY	
BI-1479-0009	II	BI 1810631	7	Urothelial	HER2 mutated	
			14	Urothelial	HER2 amplified	
CPI0209	II	CPI0209 + Enzalutamide	M8	Prostate adenocarcinoma		Patient must have received at least one prior ARPi (except enzalutamide) treatment in mCRPC or hormone sensitive prostate cancer (HSPC) disease stage
D926UC0001	II	Dato-DXd + Rilvegostomig	6E	Urothelial carcinoma		No prior systemic therapy in the advanced or metastatic setting; Progression >12 months after platinum based neoadjuvant or adjuvant therapy is eligible
DS7300-203	I/Ib	Ifinatamab Deruxtecan (I-DXd)	UC	Urothelial cancer		Relapse or progression after at least 1 prior line of ICI-containing systemic therapy and 1 prior line of systemic chemotherapy, given in combination with other anticancer therapy or separately, with a maximum of 3 prior therapy lines
DSB2455-001	Ia/Ib	DSB2455	Part B cohort A	mCRPC	one or more alteration in the following genes: BRCA1, BRCA2, PALB2, RAD51C, RAD51D	1 line max (PARP-inhibitor based)
GSK-223054	I	GSK5764227	Part 1b dose expansion	Urothelial cancer		Progression or intolerance to 1 or more lines of systemic therapy including enfortumab vedotin-ejfv [PADCEV] in the advanced/metastatic setting
LOXO-FG3-22001	I	monotherapy	B1	Urothelial cancer		Individual has progressed on or discontinued erdafitinib and Enfortumab vedotin individuals must also have received all other standard therapies
			B4	Urothelial cancer		
SNV1521-101	I	SNV1521	Part 3a	Metastatic Castration-resistant Prostate Cancer	Participants harboring known deleterious or suspected deleterious germline or somatic mutations in BRCA1/2, PALB2, and/or RAD51B/C/D by local assay.	May have received up to two lines of cytotoxic chemotherapy in the advanced/metastatic setting. Participants must not have received a prior PARPi

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STUDY NAMET	PHASE	MOLECULE	COHORT TYPE			PREVIOUS LINE
			COHORT NUMBER	TUMOR TYPE	SPECIFICITY	
ART0380C001		ART0380 + irinotécan	B5	CRC	ATM neg	2 lines maximum / must have previously received fluoropyrimidine-, oxaliplatin- and irinotecan-based chemotherapy /must NOT have received: Fruquintinib, regorafenib or trifluridine/tipiracil
			B6	pancreas	ATM neg/ATM low	Participant must have mPDAC that has not been previously treated with chemotherapy
BI-1479-0009	II	BI1810631	15	Biliary tract	HER2 amplified	
			19			
			20			
BI-1479-0012	Ib	Trastuzumab + mFOLFOX + Zongertinib	N	metastatic CRC	HER2+	
		mFOLFOX + Zongertinib	M	metastatic CRC	HER2+	
	II	Trastuzumab + Zongertinib	L	metastatic CRC	HER2+	Patients must not be likely to tolerate or derive clinical benefit from additional standard therapy. Prior treatment with a small-molecule HER2 inhibitor, a HER2-targeted monoclonal antibody (mAb), or a HER2-targeted antibody-drug conjugate is not permitted.
BreAK CRC 001	I	1st line induction COMBO SoC(mFOLFOX6 ± bevacizumab) + STC 1010	Escalation	Stage IIIC, T4b or Stage IV	No BRAFm, MSS/pMMR	Naif : eligible for 1st line treatment with SOC mFOLFOX6 ± bevacizumab
DSB2455-001	Ia/Ib	DSB2455	Part B cohort F	pancreas	one or more alteration in the following genes: BRCA1, BRCA2, PALB2, RAD51C, RAD51D	
SRP-22C102	I	ADU-1805	Cohort A	Colorectal cancer	MSS	Patients who have received no more than 2 lines of treatment
UNLOCK ASP3082	II	ASP3082	2	PDAC	KRASG12D	Only one prior line of chemotherapy therapy for a minimum of 5 months

Head and Neck

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STUDY NAME	PHASE	MOLECULE	COHORT TYPE			PREVIOUS LINE
			COHORT NUMBER	TUMOR TYPE	SPECIFICITY	
BI-1479-0009	II	BI 1810631	11	Salivary gland	HER2 overexpressed/amplified	
GSK-223054	I	GSK5764227	Part Ib dose expansion	HNSCC		Progression on or intolerance to 1 or 2 lines of therapy (including platinum-based chemotherapy and an anti-PD-(L)1 antibody) in the recurrent, unresectable, or metastatic setting

SNC

STUDY NAME	PHASE	MOLECULE	COHORT TYPE			PREVIOUS LINE
			COHORT NUMBER	TUMOR TYPE	SPECIFICITY	
F8394-201	II	FORE8394 900mg	Subproto A	SNC +/- metastasis or progressive SNC tumor with BRAF fusion	BRAF fusion (tumor tissue or liquid biopsy)	At least one standard line
			Subproto B	Recurring SNC tumor	BRAF V600E mutation	At least one prior line, radiotherapy included

Dermatology

STUDY NAME	PHASE	MOLECULE	COHORT TYPE			PREVIOUS LINE
			COHORT NUMBER	TUMOR TYPE	SPECIFICITY	
KN-8701	Ib	KIN-2787	B1	melanoma	BRAF class II fusion	Received prior locally approved standard of care appropriate for their tumor type and stage of disease, RAFi naïve

Solid tumors

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STUDY NAMEC	PHASE	MOLECULE	COHORT TYPE			PREVIOUS LINE
			COHORT NUMBER	TUMOR TYPE	SPECIFICITY	
BI-1479-0009	II	BI 1810631	21	Agnostic	HER2 amplified	
DSB2455-001	Ia/Ib	DSB2455	Part B cohort D	Brain metastasis	one or more of the following: BRCA1, BRCA2, PALB2, RAD51C, RAD51D	
F8394-201	II	FORE8394 (900mg) +/-Cobicistat (150mg)	Subproto C	Rare solid tumors (except SNC)	BRAF V600 mutated	Must have received standard therapy Or intolerant to available treatment
KN-8701	Ib	KIN-2787	B1	Solid tumors except CRC	BRAF class II fusion	Must have received prior locally approved standard of care appropriate for their tumor type and stage of disease, RAFi naïve
LOXO-FG3-22001	I	monothrapy	C1	non-urothelial solid tumor	FGRF3 alteration	Participants must be FGFR inhibitor naïve Participants have received all standard therapies

Solid tumors

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			COHORT NUMBER	TUMOR TYPE	SPECIFICITY	
MegaMOST	II	Avapritinib	G	Solid tumors	Activating mutations of KIT exon 17 or PDGFRA exon 18 associated or not to mutation on KIT exon 11 or PDGFRA exon 12/14	At least one line for metastatic disease
		Trametinib QD + Dabrafenib BID	F	Solid tumors	BRAF V600 mutation, tumor tissus or liquid biopsy Except melanoma, lung and CRC	
MOST PLUS	II	Nilotinib		PVNS	ABL1, KIT, PDGFRA, PDGFRB, DDR1, DDR2, CSF1R mutations	At least one line
MS201460	I	AntiGD2	I	Sarcomas, glioblastomas uresecable		2 prior lines
ODM-212	I/II	ODM-212	Part 2	Malignant pleural mesothelioma (MPM), Epithelioid hemangioendothelioma (EHE), Cholangiocarcinoma (CCA), HNSCC, NSCLC, CRC, Hepatocellular cancer (HCC), CRPC. Any other solid tumours with available local data for loss-of-function genetic alterations (truncating mutations or gene deletion) in NF2/LATS1/LATS2 or YAP/TAZ fusions. Any other solid tumour based on emerging scientific data as per sponsor's decision		