Oncology Clinical Trials - Current As Of May 2026

TUMOUR	TRIAL	Line	KEY POINTS - TRIAL/AGENT	PI	SC
Breast	INAVO	1st	A ph III study evaluating Inavolisib in combo with Phesgo vs placebo in combi with phesgo as maintenance therapy after 1st line induction tx in pts w PIK3CA mutated HER2 positive locally advanced or metastatic breast cancer	MW	JG
Breast	CAMBRIA-2	Adj	A Ph III, study to assess Camizestrant (AZD9833) vs Standard Endocrine Therapy for pts With ER+/HER2- Early Breast Cancer and an Intermediate-High/High Risk of Recurrence Who Have Completed Definitive Locoregional Treatment and Have No Evidence of Disease	AM	RC
Breast	OLIO	Neoadj.	A ph II, randomised, non-comparative two-arm trial of neoadjuvant chemo plus either olaparib or olaparib + durvalumab in young, pre-menopausal women with HRD-enriched, HR-positive, HER2-negative early breast cancer	AM	RC
Breast	MONITOR HER2	N/A	Frequency of Cardiac Monitoring in Patients Treated with HER2-Directed Therapies for Breast Cancer	DD	JG
Breast	CECI830A12101	2nd	Ph I/II study of ECI830 as a single agent and in combination with ribociclib and endocrine therapy in patients with advanced hormone receptor positive, HER2-negative breast cancer and advanced solid tumors	MW	JG
Breast	DESTINY BREAST 15	2nd	Ph 3b, Multicenter, Global, Interventional, Open-label Study of T-DXd, an Anti-HER2-ADC, in Subjects who Have Unresectable and/or Metastatic HER2-low or HER2 IHC 0 Breast Cancer	MW	JG
Breast	OPERA-01	2nd +	Ph 3 study of OP-1250 monotherapy VS SOC for the tx of ER+, HER2- advanced or metastatic breast cancer following endocrine and CDK 4/6 inhibitor therapy	DD	RC
Breast	Ascent-05	1st	A Ph 3 study of adjuvant sacituzumab govitecan and pembrolizumab vs treatment of physician's choice in patients with TNBC who have residual invasive disease after surgery and neoadjuvant therapy	MW	JG
Breast	STML-ELA-0422	Adj	Elacestrant versus Standard Endocrine Therapy in Women and Men with Node-positive, Estrogen Receptor-positive, HER2-negative, Early Breast Cancer with High Risk of Recurrence—A Global, Multicenter, Randomized, Open-label Phase 3 Study (ELEGANT)	MW	JG
Breast	CLEE011O12001	Adj	A phase IIIb study to characterize the efficacy and safety of Adjuvant ribociclib plus endocrine therapy in a close-to-clinical practice patient population with HR+ HER2− early breast cancer (Adjuvant WIDER)	MW	JG
Breast	OP1250-302 (Opera-02)	1st	A Phase 3 Randomized, Double-Blind, Active-Controlled Study of Palazestrant with Ribociclib Versus Letrozole with Ribociclib for the First-Line Treatment of ER+, HER2– Advanced Breast Cancer (OPERA-02)	AM	JG
Breast	MK2870-032	1st	A Phase 3, Randomized, Open-label Study to Evaluate the Efficacy and Safety of sac-TMT (Sacituzumab Tirumotecan, MK-2870) Followed by Carboplatin/Paclitaxel vs Chemotherapy, Both in Combination With Pembrolizumab as Neoadjuvant Therapy for High-Risk, Early-Stage, Triple-Negative Breast Cancer or Hormone Receptor-low Positive/Human Epidermal Growth Factor Receptor-2 Negative Breast Cancer	MW	RC
All tumour types	KN-587	Any	Ph 3, Long term safety and efficacy in patients currently on treatment or followup from a Pembrolizumab trial	DL	KB
Biliary Tract	DESTINY-Biliary-01	1st	Ph 3 Study of Trastuzumab Deruxtecan (T-DXd) and Rilvegostomig vs Standard-of-Care Gemcitabine, Cisplatin, and Durvalumab for First Line Locally Advanced or Metastatic HER2-expressing Biliary Tract Cancer	ZW	JP
Solid Tumours	HERTHENA	2nd +	A Phase 2, Multicenter, Multicohort, Open-Label, Proof of Concept Study of Patritumab Deruxtecan (HER3-DXd; U3-1402) in Subjects with Locally Advanced or Metastatic Solid Tumors	SF	SH
Solid Tumours	Genescreen	Any	GeneScreen 5-FU: DPYD Genotype-guided dose Personalisation for Fluoropyrimidine prescribing in Solid Organ Cancer Patients	ZW	MJ
Unknown primary	SUPER-ED	NA	Solving Unknown Primary cancER Earlier Diagnosis (SUPER-ED): A stepped wedge cluster randomised controlled trial implementing a Model of Care to support earlier diagnosis	ZW	KB
Pancreas	NEOFOL-R	1st	Efficacy of Neoadjuvant FOLFIRINOX in Resectable pancreatic cancer: An international multicenter Randomized, controlled trial	DL	MJ
Biliary Tract	BIL-PPP	2nd +	Phase II study of the combination of durvalumab (MEDI4736) (PDL1 inhibitor) and olaparib (PARP inhibitor) in advanced cholangiocarcinoma after initial chemotherapy and durvalumab (BIL-PPP)	DL	SH
GOJ	Destiny gastric 05	1st	A Multicenter, Randomized, Open-Label, Phase 3 Trial of Trastuzumab Deruxtecan (Enhertu®) Plus Chemotherapy Plus or Minus Pembrolizumab versus Chemotherapy Plus Trastuzumab Plus or Minus Pembrolizumab as First-line Treatment in Participants with Unresectable, Locally Advanced or Metastatic HER2-positive Gastric or Gastroesophageal Junction (GEJ) Cancer (DESTINY-Gastric05)	DL	MJ
Gastric/GOJ	Lucerna	1st	A Phase 3, Double-blind, Randomized Study of Zolbetuximab in Combination With Pembrolizumab and Chemotherapy (CAPOX or mFOLFOX6) in First-line Treatment of Locally Advanced Unresectable or Metastatic Gastric or Gastroesophageal Junction Adenocarcinoma in Participants Whose Tumors Are HER2-negative, Claudin (CLDN) 18.2-positive and Programmed Death-ligand 1 (PD-L1)-Positive	ZW	KB
Colorectal	KANDLELIT-012	1st	A Phase 3, Randomized, Open-label, Multicenter Clinical Study to Evaluate the Safety and Efficacy of MK-1084, Cetuximab, and mFOLFOX6 versus mFOLFOX6 With or Without Bevacizumab as First-line Treatment of Participants With KRAS G12C-mutant, Locally Advanced Unresectable or Metastatic Colorectal Cancer (KANDLELIT-012)	ZW	JP
Pancreas	XL092-311	Any	A Phase 2/3, multicenter, randomized open-label study of zanzalintinib vs everolimus in participants with previously treated, unresectable, locally advanced or metastatic neuroendocrine tumors	DL	JP

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Colorectal	ABT-301-103	3rd	An Open-label, Multicenter, Phase 1/2 Study Exploring the Safety and Efficacy of ABT-301 in Combination with Tislelizumab and Bevacizumab in Participants with Proficient Mismatch Repair (pMMR)/non-Microsatellite Instability-High (non-MSI-H) Locally Advanced or Metastatic Colorectal Cancer (mCRC)	DL	SH
Gastric/GOJ	CA266-0004	1st	A Blinded, Randomized, Phase 2/3 Study of Pumitamig in Combination with Chemotherapy Versus Nivolumab in Combination with Chemotherapy in Participants with Previously Untreated Advanced or Metastatic Gastric, Gastroesophageal Junction, or Esophageal Adenocarcinoma	DL	JP
Colorectal	DSP107-003	3rd	A Randomized, Open-label, Phase 2b Study to Compare the Efficacy of DSP107 in Combination with Atezolizumab Versus Fruquintinib in Patients with Advanced Microsatellite Stable Colorectal Cancer	DL	JP
Pancreas	PemOla	1sts	A phase II study combining pembrolizumab with olaparib in metastatic pancreatic adenocarcinoma patients with mismatch repair deficiency or tumour mutation burden > 4 mutations/Mb	AS	JP
Prostate	AMG 509	3rd	Ph I, AMG 509 in Subjects with Metastatic Castration-resistant Prostate Cancer	HH	LD
Prostate	MK5684-004	2nd +	Ph 3, Randomized, Open-label Study of MK-5684 vs Alternative Abiraterone Acetate or Enzalutamide in pts w mCRPC That Progressed On or After Prior Treatment with One Nextgeneration Hormonal Agent (NHA)	HH	SM
Prostate	GO44537	2nd +	Ph I, Open-Label, Multicenter, Dose-Escalation And Expansion Study Evaluating The Safety, Pharmacokinetics, And Activity Of RO7656594 In pts w Advanced Or Metastatic Prostate Cancer	EL	LD
Prostate	CJSB462C12201	1st	A Phase II, randomized, open-label, multi-center study of JSB462 (luxdegalutamide) in combination with abiraterone in adult male patients with metastatic hormone-sensitive prostate cancer (mHSPC)	HH	HG
Endocrinology	I-FIRST	Any	A Prospective, Multi-Centre Trial of TKI Redifferentiation Therapy in Patients with RAIR Thyroid Cancer (I-FIRST study)	MM	LD
HNSCC	GS-US-699-7184	2nd +	A Phase 2 Platform Study of Novel Combination Therapies in Participants With Head and Neck Squamous Cell Carcinoma	MA	EQ
Neuro	LUMOS2	2ND	Phase 2, multi-centre, open-label, biomarker-directed, umbrella, clinical trial for recurrent IDH mutant, grade 2/3 glioma.	EA	SL

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TUMOUR	TRIAL	Line	KEY POINTS - TRIAL/AGENT	PI	SC
NSCLC	OCEANiC	Adjuvant	Phase II, Open-label, Multi-centre Clinical Trial of Osimertinib With or Without Adjuvant Chemotherapy Guided by Tumour NGS Co-mutation Status and ctDNA Detection in Patients With Stage IIA-IIIA EGFR-Mutant NSCLC Following Complete Surgical Resection	SA	BN
NSCLC	Tropion-10	1s	Ph III Study of Dato-DXd in Combo w AZD2936 or AZD2936 Monotherapy vs Pembro Monotherapy for First-line tx of Pts w Locally-advanced or Metastatic Non-squamous NSCLC With High PD-L1 Expression and Without Actionable Genomic Alterations	SA	BN
NSCLC	Tropion-14	1st	Ph III Study of Osimertinib With or Without Dato-DXd as First-line Treatment in Participants w/ EGFR Mutation-positive, Locally Advanced or Metastatic NSCLC	SA	HS
NSCLC	BNT327-06	1st	A Phase II/III, multisite, randomized master protocol for a global trial of BNT327 in combination with chemotherapy and other investigational agents in first-line non-small cell lung cancer	SA	AF
NSCLC	Rosetta-02 (BNT327-06)	1st	A Phase II/III, multisite, randomized master protocol for a global trial of BNT327 in combination with chemotherapy and other investigational agents in first-line non-small cell lung cancer	SA	AF
NSCLC	RAC-020		A Phase 1 Dose Escalation and Expansion Study to Investigate the Safety, Tolerability, Pharmacokinetics and Preliminary Efficacy of Intravenous RC220 in Combination with Osimertinib in Patients with EGFR-mutant Non-Small Cell Lung Cancer	SA	ZF
NSCLC	RMC Lung 101 (sub-D)	2nd	A Phase 2, Open-Label, Multicenter Study of Zoldonrasib (RMC-9805) in Previously Treated Patients with RAS G12D-Mutant Non-Small Cell Lung Cancer (NSCLC) – Subprotocol D	SA	HS
SCLC	DeLLphi-309	2nd	A Phase 2, Open-label, Randomized, Multicenter Study of Tarlatamab Dosing Regimens in Subjects with SCLC	SA	ZF
SCLC	MK 6070-002	2nd	A Phase 1b/2 Open-Label Clinical Study to Evaluate the Safety and Efficacy of MK-6070 and Ifinatamab Deruxtecan (I-DXd) in Participants With Relapsed/Refractory Extensive-Stage Small Cell Lung Cancer	SA	TI
Meso	Evolve-02 Meso (D798MC00002)	1st	A Phase II, Multi-Center, Master Protocol to Evaluate the Efficacy and Safety of Volrustomig as Monotherapy or in Combination with Anti-cancer Agents in Participants with Advanced/Metastatic Solid Tumors	SA	AF
Ovarian	IMGN853-0425	2nd +	A randomized Phase 2, open-label study of mirvetuximab soravtansine in patients with platinum-resistant advanced high-grade epithelial ovarian, primary peritoneal, or fallopian tube cancers with high folate receptor-alpha expression testing 2 schedules of administration for dose optimization, with a separate cohort to determine starting dose in patients with moderate hepatic impairment	KW	ED
Ovarian	FLORENZA		A Phase 2, Open-Label, Randomized, Master Protocol Dose Optimization Study to Evaluate Safety and Efficacy of Multiple Treatment Combinations with Mirvetuximab Soravtansine in Subjects with Ovarian Cancer (FLORENZA)	KW	ED

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Ovarian	Catalina-2	2nd +	A Phase 2 Study Evaluating the Efficacy and Safety of TORL-1-23 in Women With Advanced Platinum-Resistant Epithelial Ovarian Cancer (Including Primary Peritoneal and Fallopian Tube Cancers) Expressing Claudin 6	GH	DS
Ovarian	DOVE	2nd +	A three-arm randomized phase II study of Dostarlimab alone or with Bevacizumab versus nonplatinum chemotherapy in recurrent gynecological clear cell carcinoma: DOVE (APGOT-OV7/ ENGOT-ov80 study)	SF	DS
Endometrial	BNT323-01	2nd +	A Phase III, randomized, multi-site, open-label trial of BNT323/DB-1303 versus investigator's choice of chemotherapy in previously treated patients with HER2-expressing recurrent endometrial cancer	SF	HT

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Solid Tumour	SR854A-001	2nd +	Phase 1, Dose Escalation, Safety, Tolerability, and Pharmacokinetic Study of SR-8541A (ENPP1 inhibitor) Administered Orally as Monotherapy in Subjects with Advanced/Metastatic Solid Tumors	CO	LS
Solid Tumour	VT3989-001 - Vivace	2nd +	Phase I/II, Multi-Center, Open-Label Study of VT3989, Alone or in Combination, in pts with Locally Advanced or Metastatic Solid Tumors	EA	ID
Solid Tumour	CP-AU-007-01	2nd +	Ph 1 first in human dose escalation of AU-007 monoclonal antibody to IL2 in advanced solid tumours	SF	OC
Solid Tumour	MK-1084	2nd +	Ph 1, MK-1084 (KRAS G12C inhibitor) as monotherapy in advanced solid tumours or in combination with pembrolizumab (NSCLC)	EA	HA
Solid Tumour	AXA-042-FIH-01	2nd +	Ph1, A Phase 1a/1b, first-in-human, open-label, non-randomized, multicenter, dose-escalation and dose-expansion study to evaluate the safety, tolerability, pharmacokinetics, and pharmacodynamics of AXA-042 as monotherapy and in combination with checkpoint inhibitors in subjects with advanced solid tumours.	SF	HA
Solid Tumour	Genentech GO43860	2nd +	A Phase 1a/1b open label, multicentre, dose-escalation study to evaluate the safety, pharmacokinetics, and activity of RO7502175 as a single agent and in combination with atezolizumab in patients with locally advanced or metastatic solid tumours	SF	HA
Solid Tumour	HM-EZHI-101	2nd +	A Phase I, Open-Label, Multicenter, Dose Escalation and Expansion Study of HM97662 as a Single Agent in Patients With Advanced or Metastatic Solid Tumours	CO	LM
Solid Tumour	GS-US-570-6015	2nd +	Phase 1 study of GS-1811 as monotherapy and in combination with a checkpoint inhibitor in patients with advanced solid tumours	DD	PM
Solid Tumour	AWT020	2nd +	Ph 1/2, FIH, Open-label Study of Single-agent AWT020 in pts w Progressive Locally Advanced or Metastatic Cancer	SF	LS
Solid Tumour	DCSZ11-101	2nd +	Ph 1, Multicenter, Open-Label, Dose Escalation, and Dose Expansion Study to DCSZ11 as a Monotherapy and in Combo in pts with Advanced or Metastatic Solid Tumors	DD	HA
Solid Tumour	BG-68501-101	2nd +	A Phase 1a/1b Study of BG-68501 a Selective CDK2 Inhibitor, in Participants With Advanced Solid Tumors	DD	KD
Solid Tumour	BG-C9074-101	2nd +	Ph 1a/1b Study of BG-C9074, an ADC Targeting B7H4, as Monotherapy and in combo w/ Tislelizumab in pts With Advanced Solid Tumors	DD	OC
Solid Tumour	PMV-586-101	2nd +	A Ph 1/2 Open-label, Multicenter Study to Assess PC14586 in pts w/ Locally Advanced or Metastatic Solid Tumors Harboring a TP53 Y220C Mutation	CO	KD
Solid Tumour	YL211-INT-101-01	2nd +	A Phase 1, Multicenter, Open-Label, First-in-Human Study to Evaluate the Safety, Tolerability, Pharmacokinetics, and Efficacy of YL211 in Patients with Advanced Solid Tumors	SF	OC
Solid Tumour	BGB-53038-101	2nd +	A Ph 1a/1b Study to Investigate BGB-53038, a Pan-KRAS Inhibitor, as Monotherapy or in Combo in Patients With Advanced or Metastatic Solid Tumors With KRAS Mutations or Amplification	SF	HA
Solid Tumour	BG-C137-101	2nd +	A Phase 1a/b, Open-label, Multicenter Study to Investigate BG-C137, an Antibody-Drug Conjugate Targeting FGFR2b, in Patients With Advanced Solid Tumors	CO	LM
Solid Tumour	PAUF-I	2nd +	A FIH, Phase 1/2a, Multicentre, Open-label Study Evaluating PBP1510 in Patients with Advanced/Metastatic Pancreatic Cancer	DD	ID
Solid Tumour	BGB-58067-101	2nd +	A Ph 1a/b Study Investigating BGB-58067, an MTA-Cooperative PRMT5 Inhibitor in Patients With Advanced Solid Tumors	EA	HG
Solid Tumour	CP-IVX037	2nd +	A ph 1 study of intratumoral IVX037 as monotherapy or in combo w an immune checkpoint inhibitor in pts w advanced/metastatic solid tumors	SF	LM
Solid Tumour	BNT317	2nd +	Ph I, FIH, open-label, dose escalation study of BNT317 in pts w advanced solid tumors	SF	HG
Solid Tumour	CS2009-101	2nd +	Ph I, Dose-Escalation and Dose-Expansion Study to Evaluate CS2009, a tri-specific antibody targeting PD-1/VEGFA/CTLA-4, in Participants with Advanced Solid Tumors	SF	KD
Solid Tumour	BM230-01	2nd +	Ph I, Multicenter, Non-randomized, Open-label Study to Evaluate the Safety, Tolerability, Pharmacokinetics, and Efficacy of BM230 in Patients with Advanced Solid Tumors	EA	LS
Solid Tumour	SNV1521-101	2nd +	Ph I, open-label dose escalation and expansion study of SNV1521 in pts with advanced solid tumors	SF	LS
Solid Tumour	SNV4818-101	2nd +	Ph I, Open-Label Dose Escalation and Expansion Study of SNV4818 as Monotherapy or in combo w other Anticancer Agents in pts with Advanced Solid Tumors.	SF	OC
Solid Tumour	MOMA-341-001	2nd +	A Phase 1 Study of MOMA-341 as Monotherapy or Combination Therapy in Participants with Advanced or Metastatic Solid Tumors	EA	LM
Solid Tumour	BBI-4182-101	2nd +	A Phase 1/2, First-in-Human, Dose Escalation and Expansion Study of BDC-4182 as a Single Agent in Patients with Advanced Gastric and Gastroesophageal Cancer.	SF	PM

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Solid Tumour	BG-C0902-101	2nd +	A Phase 1a/b Study to Investigate the Safety, Tolerability, Pharmacokinetics, Pharmacodynamics, and Preliminary Antitumor Activity of BG-C0902, an Antibody-Drug Conjugate Targeting EGFR × MET, Alone and in Combination With Other Therapeutic Agents in Patients With Advanced Solid Tumors.	DD	HG
Solid Tumour	BNT329-01	2nd +	First-in-human, open-label, multi-site, Phase I/IIa, dose escalation trial with expansion cohorts to evaluate safety and preliminary efficacy of BNT329 in participants with advanced solid tumors known to express CA19-9	SF	LM
Solid Tumour	MK-4716-001	2nd +	A Phase 1, Open-Label, Multicenter Study to Assess Safety, Tolerability, Pharmacokinetics, and Efficacy of MK-4716 as Monotherapy and as Part of Combination Therapy in Participants with KRAS-Altered Advanced or Metastatic Solid Tumors	SF	ID

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Stream Contact Numbers

Breast – 0459 302 205
Gynaecological – 0491 299 561
GI – 0436 386 758
Lung – 0417 607 416
GU/neuro/Head & Neck/skin – 0436 387 664
Phase 1 – 0474 769 510

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