

Pancreatic cancer trials open to recruitment – November 2025

This list includes trials targeted at pancreatic cancer alone and trials for many solid tumours which include pancreatic cancer. If you are a patient with pancreatic cancer and wish to discuss whether a clinical trial is right for you, please discuss with your treating Medical Oncologist.

NOTE: Trials listed are available for patients with pancreatic cancer to be referred for consideration. Several trials may have issues with slot availability or waitlists.

Descriptive stages of pancreatic cancer

Medical Term	What it means
Resectable	This is an early stage of pancreatic cancer where the cancer can still be removed by surgery
Unresectable or locally advanced	The pancreatic cancer has not spread around the body but unfortunately the cancer cannot be removed by surgery due to the tumour size, location, or involvement of surrounding structures
Metastatic	The pancreatic cancer has spread to other parts of the body



PANCREATIC CANCER TRIALS IN VICTORIA (Trials with specific cohorts for pancreatic cancer)

Trial Title	Targets (Genomic)	Target population (Trial suitable for patients with this stage of pancreatic cancer)	Treatment + Further Information (What the study involves)	Site (Where the study is being offered)	Contact Details (Email the contact person listed with any enquiries)
NeoFOL-R Efficacy of Neoadjuvant FOLFIRINOX in Resectable pancreatic cancer: An international multicentre Randomized, controlled trial	N/A	Exclusion: Arm A Borderline resectable chemo Locally advanced Previous treatment for pancreatic Arm B	mFOLFIRINOX chemotherapy Arm A: Surgery then 12 cycles of chemotherapy Arm B: 6 cycles of chemotherapy before surgery followed by 6 of	Epworth Health Monash Health	EHJreissatiCentre@epworth.org.au gi.oncresearch@monashhealth.org
(NeoFOL-R) - Australian protocol			cycles after surgery Further information: NeoFOL-R - Victorian Cancer Trials Link	Alfred Health	act-m@alfred.org.au
Investigation of the safety and efficacy of irreversible electroporation (IRE) using the NanoKnife® System in patients	N/A	Inclusion: 3 months of chemotherapy (FOLFIRINOX or Gemcitabine based – 1 line only)	NanoKnife System for Irreversible Electroporation (IRE) IRE is a technique using non- thermal energy to create permanent nanopores in the cell	Peter MacCallum Cancer Centre	SurgicalResearchteam@petermac.or g
with unresectable stage 3 pancreatic cancer who have received 3 months of chemotherapy			membrane in order to disrupt cellular homeostasis. Further information: DIRECT/InspIRE Australia - Victorian Cancer Trials Link	Alfred Health Epworth Health	charles.pilgrim@monash.edu EHJreissatiCentre@epworth.org.au



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	(Trials with specific cohorts for pancreatic cancer)						
Trial Title	Targets (Genomic)	Target population (Trial suitable for patients with this stage of pancreatic cancer)	Treatment + Further Information (What the study involves)	Site (Where the study is being offered)	Contact Details (Email the contact person listed with any enquiries)		
AMG193 20230223 A Phase 1b Study Evaluating the Safety, Tolerability,	MTAP deletion	METASTATIC/LOCALLY ADVANCED (Second line)	AMG 193 AMG 193 is a PMRT5 inhibitor which is administered orally.	Peter MacCallum Cancer Centre	PCCTU.MoncC@petermac.org		
Pharmacokinetics, and Efficacy of AMG 193 in Combination with Other Therapies in Subjects with Advanced Gastrointestinal, Biliary Tract, or Pancreatic Cancers with		Inclusion: Pancreatic cancer Homozygous MTAP-deletion Exclusion:	Further information: https://trials.cancervic.org.au/det ails/vctl nct06360354	Austin Health	samantha.chakar@austin.org.au		
Homozygous MTAP-deletion AMG20230223		Prior MAT2A inhibitor or PRMT5 inhibitor		Epworth Health	EHjreissaticentre@epworth.org.au		
Clarity-PT01 A Phase II, Open-label, Multicentre Study to Evaluate Safety, Tolerability, Efficacy, PK, and Immunogenicity of AZD0901 as Monotherapy and in Combination With Anti-cancer Agents in Participants With Advanced Solid Tumours Expressing Claudin 18.2.	CLDN18.2	METASTATIC/LOCALLY ADVANCED (First line) Inclusion: Pancreatic cancer Treatment naïve CLDN18.2 positive Exclusion: Exposure to prior CLDN18.2 targeted agents except anti- CLDN18.2 monoclonal antibody	AZD0901 (antibody-drug conjugate) AZD0901 targets and binds to CLDN18.2, which is a protein on tumour cells, and then delivers cytotoxic agents which damage these cancer cells. Further information: https://trials.cancervic.org.au/det ails/vctl_nct06219941	Peter MacCallum Cancer Centre	PCCTU.MoncC@petermac.org		
AMPLICITY (AMP945-202) A Phase 1b/2a, Multicenter, Open Label Study of the Safety, Efficacy and Pharmacokinetics of Narmafotinib in Combination With Modified FOLFIRINOX in Pancreatic Cancer Patients	FAK	METASTATIC/LOCALLY ADVANCED (First line) Inclusion: Treatment naïve for metastatic disease	Narmafotinib Narmafotinib is an oral FAK inhibitor. Narmafotinib will be administered in combination with mFOLFIRINOX chemotherapy Further information:	Epworth	PancreaticCentre@epworth.org.au		



PANCREATIC CANCER TRIALS IN VICTORIA (Trials with specific cohorts for pancreatic cancer) **Trial Title** Targets **Target population** Treatment + Further Site Contact Details (Trial suitable for patients with (Where the study (Email the contact person listed (Genomic) Information this stage of pancreatic cancer) is being offered) with any enquiries) (What the study involves) https://pancreaticcentre.org.au/t reatment/clinical-trials-andresearch/amplicity METASTATIC/LOCALLY N/A Fecal Microbiota Transplantation **FMT** Epworth EH-ADVANCED **Fecal Microbiota Transplantation** (First line) Further information: PancreaticCentre@epworth.org.au Inclusion: to improve pain, symptom https://www.pancreaticcentre.or management and treatment Pancreatic cancer g.au/treatment/clinical-trialsefficacy in patients with Treatment naïve (first line) and-research/fmt pancreatic cancer **Exclusion:** Antibiotic use within 8 weeks of randomisation (1 dose with ERCP allowed) dMMR/MS METASTATIC/LOCALLY Gi.oncresearch@monashhealth.org **PemOla** Pembrolizumab and Olaparib Monash Health I-high **ADVANCED** A phase II study combining or (First and second line) Pembrolizumab in anti-PD1 pembrolizumab with olaparib in TMB > 4 antibody Olaparib is a PARP inhibitor metastatic pancreatic mutations Inclusion: adenocarcinoma patients with / Mb Pancreatic cancer

Further information:

Expected to open imminently

Contact trials unit

Awaiting opening date

mismatch repair deficiency or

tumour mutation burden > 4

mutations/ Mb



Peter MacCallum

Cancer Centre

Epworth

Monash Health

EH-

Ith.org

Contact Details

with any enquiries)

act-m@alfred.org.au

(Email the contact person listed

PCCTU.MoncC@petermac.org

PancreaticCentre@epworth.org.au

earlyphase.oncresearch@monashhea

PANCREATIC CANCER TRIALS IN VICTORIA (Trials with specific cohorts for pancreatic cancer) Targets **Target population** Treatment + Further Site (Trial suitable for patients with (Where the study (Genomic) Information this stage of pancreatic cancer) is being offered) (What the study involves) FΔP METASTATIC/LOCALLY Alfred Hospital [177Lu]Lu-FAP-2286 **ADVANCED** A Phase 1/2. Multicentre. Open-(First line) [177Lu]Lu-FAP-2286 is a label. Non-randomized Study to radiopharmaceutical that targets

fibroblast activation protein

(FAP). This treatment works by

binding to the FAP to allow the

directly to FAP-expressing cancer

https://trials.cancervic.org.au/det

Zamaporvint is an oral porcupine

https://www.pancreaticcentre.or

g.au/treatment/clinical-trialsand-research/porcupine-p2ea

PBP1510 is an IgG1 monoclonal

antibody (mAb) that targets and

inhibitor. Treatment will be

administered with denosumab

targeted delivery of radiation

[177Lu]Lu-FAP-2286 with

chemotherapy

Zamaporvint

PBP1510

neutralises PAUF.

Further information:

ails/vctl nct04939610

Further information:

cells

Inclusion:

Pancreatic ductal

Combination cohort:

advanced disease

Chemotherapy naïve for

METASTATIC/LOCALLY

Progression on first line

METASTATIC/LOCALLY

treatment for incurable disease

Mandatory biopsy at enrolment

ADVANCED

(Second line)

Inclusion:

ADVANCED

Inclusion:

(Second line +)

Porcupine

PAUF

Exclusion: Active CNS disease

adenocarcinoma + other cancers

Trial Title

LuMIERE

Tumour

Porcupine P2EA

Investigate Safety and

Tolerability, Pharmacokinetics,

Activity of 177Lu-FAP-2286 in

Patients With an Advanced Solid

An open-label study to assess the

preliminary efficacy and safety of

progressed following therapy with

RXC004, (Zamaporvint), in

pancreatic cancer who have

A First in Human, Phase 1/2a,

Multicentre, Open-label Study

Evaluating the Safety, Tolerability,

patients with advanced

current standard of care.

PAUF-I

Dosimetry, and Preliminary



PANCREATIC CANCER TRIALS IN VICTORIA (Trials with specific cohorts for pancreatic cancer)

Trial Title	Targets	Target population	Treatment + Further	Site	Contact Details		
	(Genomic)	(Trial suitable for patients with	Information	(Where the study	(Email the contact person listed		
		this stage of pancreatic cancer)	(What the study involves)	is being offered)	with any enquiries)		
Pharmacokinetics, and Efficacy of		Received at least 1 line of					
PBP1510 in Patients with		chemotherapy and progressed	Further information:				
Advanced/Metastatic Pancreatic			https://trials.cancervic.org.au/det				
Cancer			ails/vctl nct05141149				
NCB161734	KRAS G12D	ADVANCED OR METASATIC SOLID CANCERS	<u>INCB161734</u>	Alfred Health	act-m@alfred.org.au		
A Phase 1, Open-Label, Multicenter		(Second line +)	INCB161734 in a KRAS G12D				
Study of INCB161734 in Participants		(Second line 1)	inhibitor				
With Advanced or Metastatic Solid		Inclusion:					
Tumors With KRAS G12D Mutation		Second line and beyond	Further information:				
			https://trials.cancervic.org.au/det				
			ails/vctl nct06179160				
AMG 410	KRAS	ADVANCED OR METASATIC SOLID CANCERS	AMG 410	Peter	PCCTU.EDD@petermac.org		
A Phase 1/1b Study Evaluating the		(Exhausted standard of care)	AMG 410 is an oral pan-KRAS	MacCallum			
Safety, Tolerability,		,,	inhibitor and will be administered	MacCallulli			
Pharmacokinetics,		Inclusion:	alone or in combination with	C C			
Pharmacodynamics, and Efficacy of		KRAS mutation or amplification	pembrolizumab and panitumumab	Cancer Centre			
AMG 410 Alone and in Combination		Exhausted conventional					
With Other Agents in Participants		treatment options	Further information:				
With KRAS Altered Advanced or		Pancreatic cohort specified in	https://trials.cancervic.org.au/det				
Metastatic Solid Tumors		part 2.	ails/vctl_nct07094113				
LK1004-101	HRR	METASTATIC/LOCALLY	EIK1004	Peninsula and	ag@paso.com.au		
	deficiency	ADVANCED					
A Phase 1/2, Open-label,		(Second line +)	EIK1004 is an oral PARP1 inhibitor	South Eastern			
Multicenter, Dose-escalation, and			and will be administered as				
Pose-Optimization Study to		Inclusion:	monotherapy	Haematology and			
valuate the Safety, Tolerability,		Suspected deleterious mutation					
and Activity of EIK1004 (IMP1707)		of select HRR genes	Further information:	Oncology Group			
s Monotherapy in Participants			https://trials.cancervic.org.au/det				
With Advanced Solid Tumors		Exclusion:	ails/vctl nct06907043				
		Prior PARP1 inhibitor					



PANCREATIC CANCER TRIALS IN VICTORIA (Trials with specific cohorts for pancreatic cancer)

Trial Title	Targets (Genomic)	Target population (Trial suitable for patients with this stage of pancreatic cancer)	Treatment + Further Information (What the study involves)	Site (Where the study is being offered)	Contact Details (Email the contact person listed with any enquiries)
ENG19	EGFR	METASTATIC/LOCALLY ADVANCED	E-EDV-D682/GC	Peninsula and	ag@paso.com.au
An open-label, multicenter, Phase I/IIa study assessing the safety and		(Third line +)	E-EDV-D682/GC is a combination of a EnGelC Dream Vector (EDV)	South Eastern	
efficacy of EGFR targeted EDVsTM carrying cytotoxic drug PNU-159682		Inclusion: Progressed on second line or	transporting the cytotoxic drug PNU-159682 to cells expressing	Haematology and	
plus concurrent immunomodulatory adjuvant non-targeted EDVs		treatment exhausted EGFR expression on local IHC or	EGFR and an EDV carrying alphagalactoslyceramide (EDV-GC).	Oncology Group	
carrying a-galactosyl ceramide in subjects with advanced EGFR-		liquid biopsy	Further information:		
expressing cancers who have failed second-line therapy or where first-		**STUDY ON HOLD**	https://trials.cancervic.org.au/det ails/vctl actrn12625000203459		
and/or second-line therapy is not appropriate (EGFR EDV-D682/GC Trial)					



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ADCE-T02-001 First-in-Human, Phase 1 Study of AMT-754, a Targeting Tissue Factor Antibody-Drug Conjugate, in Patients With Advanced Solid Tumors	Tissue Factor (TF)	ADVANCED OR METASTATIC SOLID CANCERS Inclusion: Pancreatic cancer Received ≥1 prior line of therapy No further standard therapy available Exclusion: Active CNS disease	AMT-754 AMT-754 is a targeting tissue factor antibody-drug conjugate (ADC). ADCs bind to a specific part of the surface of a cancer cell and then deliver targeted treatment directly into the cell. Further information: https://trials.cancervic.org.au/det ails/vctl nct06597721	Peninsula and South Eastern Haematology and Oncology Group	clinicaltrials@cabrini.com.au ag@paso.com.au
HM-EZHI A Phase I, Open-Label, Multicenter, Dose Escalation and Expansion Study of HM97662 as a Single Agent in Patients With Advanced or Metastatic Solid Tumors	SWI/SNF Complex aberration (ARID1A SMARCA4 SMARCA2)	ADVANCED OR METASTATIC SOLID CANCERS Inclusion: SW/SNF Complex aberration (ARID1A, SMARCA4, SMARCA2) Exclusion: Prior valemetostat or other EZH1/2 inhibitor use	HM97662 HM97662 is an oral medication (EZH1/2 dual inhibitor). Further information: https://trials.cancervic.org.au/det ails/feed-cta-trial541	Peninsula and Southeast Oncology Grampians Health (Ballarat)	ag@paso.com.au clinicaltrials@gh.org.au
BGB 58067 A Phase 1a/b Study Investigating the Safety, Tolerability, Pharmacokinetics, Pharmacodynamics, and Preliminary Antitumor Activity of BGB-58067, an MTA-Cooperative PRMT5 Inhibitor in Patients With Advanced Solid Tumors	MTAP loss	ADVANCED OR METASTATIC SOLID CANCERS Inclusion: MTAP loss (pre-screening available) Exclusion: Prior treatment with PRMT5 or MAT2A inhibitor	BGB-58067 BGB-58067 is an MTA- Cooperative PRMT5 inhibitor. Further information: NCT06589596 - Victorian Cancer Trials Link	Monash Health (Note: Austin Health is only recruiting GBM)	earlyphase.oncresearch@monashhealth.org



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A Phase 1a/1b Study to Investigate the Safety, Tolerability, Pharmacokinetics, Pharmacodynamics, and Preliminary Antitumor Activity of BGB-53038, a Pan-KRAS Inhibitor,	KRAS mutation or amplificati on (excluding KRAS	ADVANCED OR METASTATIC SOLID CANCERS Inclusion: Evidence of KRAS mutation or wild-type amplification Exclusion:	BGB-53038 BGB-53038 is a pan-KRAS inhibitor.	Monash Health	earlyphase.oncresearch@monashhea lth.org
As Monotherapy or in Combinations in Patients with Advanced or Metastatic Solid Tumors with KRAS Mutations or Amplifications	G12R)	KRAS G12R mutation Prior treatment with other RAS targeting treatment	Further information: NCT06585488 - Victorian Cancer Trials Link	Peter MacCallum Cancer Centre	PCCTU.EDD@petermac.org
YL211-INT-101-01: 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	смет	ADVANCED OR METASTATIC SOLID CANCERS Inclusion: No further standard treatment options available	YL211 YL211 is a C-MET targeted antibody-drug conjugate (ADC) available for all tumour types for dose escalation where there is either a C-met aberration or where there is a biological rationale for C-met directed therapy. Further information: YL211-INT-101-01 - Victorian Cancer Trials Link	Monash Health	earlyphase.oncresearch@monashhea lth.org
MK-1084 A Phase 1, Open-Label, Multicenter Study to Assess Safety, Tolerability, PK, and	KRAS G12C	ADVANCED OR METASTATIC SOLID CANCERS Inclusion: KRAS G12C mutation	MK1084 MK1084 is an oral KRAS G12C inhibitor.	Monash Health	earlyphase.oncresearch@monashhea lth.org



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Trial Title	Targets	Target population	Treatment + Further	Site	Contact Details
	(Genomic)	(Trial suitable for patients with	Information	(Where the study	(Email the contact person listed
		this stage of pancreatic cancer)	(What the study involves)	is being offered)	with any enquiries)
Efficacy of MK-1084 as					
Monotherapy and in Combination			Further information:		
With Pembrolizumab in Subjects			MK-1084 - Victorian Cancer Trials		
With KRASG12C Mutant			<u>Link</u>		
Advanced Solid Tumors					
<u>PRIMROSE</u>	MTAP	ADVANCED OR METASTATIC	AZD3470	Peter MacCallum	PCCTU.EDD@petermac.org
	deficient	SOLID CANCERS		Cancer Centre	
A Modular Phase I/IIa, Multi-			AZD3470 is a novel, potent and		
centre, Dose Escalation, and		Inclusion:	selective second-generation PRT5		
Expansion Study of AZD3470, a		At least 1 prior line of treatment	inhibitor.		
MTA Cooperative PRMT5		and exhausted treatment options			
Inhibitor, as Monotherapy and in		Archival or baseline tumour			
Combination With Anticancer		sample for MTAP testing	Further information:		
Agents in Patients With			PRIMROSE - Victorian Cancer		
Advanced/Metastatic Solid		Exclusion:	<u>Trials Link</u>		
Tumours That Are MTAP Deficient		Prior PRMT5 inhibitor			
<u>CS5001</u>	ROR1	ADVANCED OR METASTATIC SOLID CANCERS	<u>CS5001</u>	Epworth	connie.barlas@epworth.org.au or
A Phase I, Dose-Escalation and			CS5001 in an antibody drug		EH-
Dose-Expansion Study to Evaluate		Inclusion:	conjugate targeting ROR1		
the Safety, Tolerability,		Progression on at least 1 prior			PancreaticCentre@epworth.org.au
Pharmacokinetics, and Antitumor		line of systemic therapy	Further information:		
Activities of CS5001, an Anti-		ECOG 0-1	https://www.pancreaticcentre.or		
ROR1 Antibody-Drug Conjugate,			g.au/treatment/clinical-trials-		
Used as A Single Agent and in			and-research/cs5001		
Combination with Systemic					
Therapies in Patients with					
Advanced Solid Tumors and					
Lymphomas.					



Trial Title	Targets	Target population	Treatment + Further	Site	Contact Details
That fille	(Genomic)	(Trial suitable for patients with this stage of pancreatic cancer)	Information (What the study involves)	(Where the study is being offered)	(Email the contact person listed with any enquiries)
MarkV A Phase 1a/1b, First-in-Human, Open Label Study to Assess the Safety, Tolerability, and Pharmacokinetics of PMC-309 (Anti-VISTA), as Monotherapy and Combined With Pembrolizumab, in Patients With Advanced or	VISTA	ADVANCED OR METASATIC SOLID CANCERS Inclusion: Progressed on available prior lines of therapy Progressed on PD-1 or PD-L1 inhibitor immunotherapy	PMC-309 PMC-309 is an anti-VISTA monoclonal antibody PMC-309 will be administered alone or in combination with pembrolizumab	Grampians Health (Ballarat)	clinicaltrials@gh.org.au clinicaltrials@cabrini.com.au
Metastatic Solid Tumors		Exclusion: Prior anti-VISTA therapy	Further information: https://trials.cancervic.org.au/det ails/vctl nct05957081		
AT-0174-001 A Phase I Study to Evaluate the Safety, Tolerability, Pharmacology, and Preliminary	IDO1/TDO 2	ADVANCED OR METASATIC SOLID CANCERS Inclusion: Progressed on available prior	AT-0174 AT-0174 is a novel dual inhibitor of IDO1/TDO2	Grampians Health (Ballarat)	clinicaltrials@gh.org.au
Efficacy of AT-0174 in Subjects with Advanced Solid Malignancies		lines of therapy	Further information: https://trials.cancervic.org.au/det-ails/vctl_actrn12623000956606	St Vincent's Hospital Melbourne	OncologyTrialCoordinators@svha.org .au
AKTive-001 A Phase 1/1b Multiple Cohort Trial of ALTA2618 in Patients with Advanced Solid Tumors with AKT1 E17K Mutation	AKT E17K mutation	ADVANCED OR METASATIC SOLID CANCERS Inclusion: Exhausted standard of care therapies Exclusion:	ALTA2618 ALTA2618 is an oral AKT E17K inhibitor Further information: https://trials.cancervic.org.au/det	Cabrini	clinicaltrials@cabrini.com.au
	THE STATE OF THE S	Prior mTOR or PI3K inhibitors Presence of KRAS, NRAS, HRAS or BRAF genomic alterations	ails/vctl nct06533059		



Trial Title	Targets (Genomic)	Target population (Trial suitable for patients with this stage of pancreatic cancer)	Treatment + Further Information (What the study involves)	Site (Where the study is being offered)	Contact Details (Email the contact person listed with any enquiries)
AMT-676-01 First-in-Human, Phase 1 Study of AMT-676, an Anti-CDH17 Antibody- Drug Conjugate, in Patients with Advanced Solid Tumors	CDH17	ADVANCED OR METASATIC SOLID CANCERS Inclusion: Second or later line therapy	AMT-676 AMT-676 is an anti-CDH17 antibody drug conjugate. Further information: https://trials.cancervic.org.au/details/vctl_nct06400485	Cabrini	clinicaltrials@cabrini.com.au
D3S-001-100 A Phase 1, Open Label, Dose Escalation and Dose Expansion Study Evaluating the Safety, Tolerability, Pharmacokinetics, Pharmacodynamics, and Preliminary Efficacy of D3S 001 Monotherapy in Subjects with Advanced Solid Tumors with a KRAS p.G12C Mutation	KRAS G12C	ADVANCED OR METASATIC SOLID CANCERS Inclusion: KRAS G12C mutation *Note pancreatic specific cohort has closed. Pantumour cohort remains open with limited slots	D3S 001 D3S 001 is a KRAS G12C inhibitor Further information: https://trials.cancervic.org.au/det ails/feed-cta-trial449	Peter MacCallum Cancer Centre	clinicaltrials@cabrini.com.au PCCTU.EDD@petermac.org
A Phase 1, Open-label, Multicenter Clinical Trial of S095035 (MAT2A Inhibitor) in Adult Participants With Advanced or Metastatic Solid Tumors With Homozygous Deletion of MTAP	MAT2A MTAP	ADVANCED OR METASATIC SOLID CANCERS Inclusion: Homozygous deletion of MTAP Second line and beyond	S095035 S095035 is an oral MAT2A inhibitor. Further information: https://trials.cancervic.org.au/details/vctl_nct06188702	Alfred Health	act-m@alfred.org.au



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	(Genomic)	(Trial suitable for patients with	Information	(Where the study	(Email the contact person listed
		this stage of pancreatic cancer)	(What the study involves)	is being offered)	with any enquiries)
KEYNOTE-F49	LILRB1	ADVANCED OR METASATIC	IOS-1002 +/- Pembrolizumab	Bendigo Health	cancerresearch@bendigohealth.org.a
	LILRB2	SOLID CANCERS			
A Phase 1a/1b, First-in-human,	KIR3DL1		IOS-1002 binds to LILRB1 (ILT2),		<u>u</u>
Open-label, Non-randomized,	PD-1		LILRB2 (ILT4), and KIR3DL1		
Multicenter, Dose-escalation and		Inclusion:	receptors on innate and adaptive		
Cohort Expansion Study to		Second or later line	immune cells that suppress		
Evaluate the Safety, Tolerability,			immune responses when		
Efficacy, Immunogenicity,			activated.	Austin Health	samantha.chakar@austin.org.au
Pharmacokinetics, and					
Pharmacodynamics of IOS-1002			Pembrolizumab in anti-PD1		
Administered Alone and in			antibody.		
Combination With a PD-1					
Monoclonal Antibody in			Further information:		
Advanced Solid Tumors			https://trials.cancervic.org.au/det		
			ails/feed-cta-trial528		
GDC-7035	KRAS	ADVANCED OR METASATIC	GDC-7035	Peter MacCallum	PCCTU.EDD@petermac.org
	G12D	SOLID CANCERS		Cancer Centre	
A Phase I/II Dose-Escalation and			GD-7035 is a KRAS G12D		
Expansion Study Evaluating the			inhibitor. Treatment will be		
Safety, Pharmacokinetics, and		Inclusion:	monotherapy or in combination		
Activity of GDC-7035 as a Single		KRAS G12D mutation	with other anti-cancer		
Agent and in Combination With			treatments.		
Other Anti-Cancer Therapies in					
Patients With Advanced Solid			Further information:		
Tumors With a KRAS G12D			https://trials.cancervic.org.au/det		
Mutation			ails/vctl_nct06619587		
RO7566802	ανβ8	ADVANCED OR METASATIC	RO7566802	Peter MacCallum	PCCTU.EDD@petermac.org
	integrin	SOLID CANCERS		Cancer Centre	
A Phase I, Open-Label,			RO7566802 is a αvβ8 integrin		
Multicenter, Dose-Escalation			inhibitor delivered intravenously.		
Study Evaluating the Safety,		Inclusion:			
Pharmacokinetics, and Activity of		Second or later line	Further information:		
RO7566802 as a Single Agent and			https://trials.cancervic.org.au/det		



Trial Title	Targets	Target population	Treatment + Further	Site	Contact Details
	(Genomic)	(Trial suitable for patients with	Information	(Where the study	(Email the contact person listed
		this stage of pancreatic cancer)	(What the study involves)	is being offered)	with any enquiries)
in Combination With			ails/vctl_nct06031441		
Atezolizumab in Patients With					
Locally Advanced or Metastatic					
Solid Tumors					
MK-6837-001	TROP2	ADVANCED OR METASATIC	MK-6837 +/- pembrolizumab	Alfred Health	act-m@alfred.org.au
		SOLID CANCERS			
A Phase 1 Open-label, Multicenter			MK-6837 is a TROP2-MMAE		
Study of MK-6837 as		Exclusion:	antibody-drug conjugate		
Monotherapy and Combination		Uncontrolled HIV, Hepatitis B or C	delivered as monotherapy or in		
Therapy in Participants With Advanced/Metastatic Solid			combination with PD-1 inhibitor		
Tumors			pembrolizumab.		
Tuttiots			Further information:		
			https://trials.cancervic.org.au/det		
			ails/vctl nct06460961		
BG-C477	CEACAM5	ADVANCED OR METASATIC	BG-C477	Alfred Health	act-m@alfred.org.au
		SOLID CANCERS		, and a reduction	
A Multicenter, Open-Label, Phase			BG-C477 is an antibody-drug		
1a/b First-in-Human Study to		Inclusion:	conjugate targeting CEACAM5.		
Investigate the Safety,		CEA >5	BG-C477 will be delivered as		
Tolerability, Pharmacokinetics,			monotherapy or in combination		
Pharmacodynamics, and			with capecitabine and		
Preliminary Antitumor Activity of			bevacizumab.		
BG-C477 in Patients With					
Selected Advanced Solid Tumors			Further information:		
			https://trials.cancervic.org.au/det		
			ails/vctl nct06596473		
BP44474	dMMR/ MSI-H	ADVANCED OR METASATIC SOLID CANCERS	RO7589831	Alfred Health	act-m@alfred.org.au
A Phase I, Open-Label Study to			RO7589831 is an orally		
Evaluate the Safety, Tolerability,		Inclusion:	administered WRN inhibitor		
Pharmacokinetics,		Progressed on at least one prior			
Pharmacodynamics, and Anti-		line of therapy in the advanced	Further information:		



Trial Title	Targets	Target population	Treatment + Further	Site	Contact Details
	(Genomic)	(Trial suitable for patients with this stage of pancreatic cancer)	Information (What the study involves)	(Where the study is being offered)	(Email the contact person listed with any enquiries)
Tumor Activity of RO7589831 in Participants With Advanced Solid Tumors Harboring Microsatellite Instability (MSI) and/or Deficient Mismatch Repair (dMMR)		setting Tumour known to be d-MMR or MSI-H	https://trials.cancervic.org.au/det ails/vctl_nct06004245		
AMT-562-01	HER3	ADVANCED OR METASATIC SOLID CANCERS	AMT-562	Cabrini	clinicaltrials@cabrini.com.au
First-in-Human, Phase 1 Study of AMT-562, an Anti HER3 Antibody- Drug Conjugate, in Patients with Advanced Solid Tumors		Inclusion: Second line and beyond	AMT-562 is a novel HER3 targeting antibody drug conjugate.		
			Further information: https://trials.cancervic.org.au/det ails/vctl nct06199908		
IKSUDA A Phase 1 Dose Escalation Trial to Determine the Safety, Tolerance, Maximum Tolerated Dose, and Preliminary Antineoplastic Activity of IKSO14, a HER2-Targeting Antibody Drug Conjugate (ADC), in Participants With Advanced HER2+ Solid Tumors	HER2	ADVANCED OR METASATIC SOLID CANCERS Inclusion: HER2 IHC 1-3+ HER2 ISH negative and positive	IKS014 IKS014 is a HER2 targeting antibody drug conjugate. Further information: https://trials.cancervic.org.au/det ails/vctl nct05872295	Peninsula and Southeast Oncology	ag@paso.com.au
LOXO-RAS-200001	KRAS G12C	ADVANCED OR METASATIC SOLID CANCERS	LY3537982	Peninsula and Southeast	ag@paso.com.au
A Phase 1a/1b Study of LY3537982 in Patients With KRAS G12C-Mutant Advanced Solid Tumors		Inclusion: KRAS G12C mutation	LY3537982 is a KRAS G12C inhibitor and will be administered as monotherapy or in combination with other systemic anticancer medications.	Oncology	



Trial Title	Targets (Genomic)	Target population (Trial suitable for patients with this stage of pancreatic cancer)	Treatment + Further Information (What the study involves)	Site (Where the study is being offered)	Contact Details (Email the contact person listed with any enquiries)
			Further information: https://trials.cancervic.org.au/det_ails/vctl_nct04956640		
Stingray SR-8541A-001 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	ENPP1	ADVANCED OR METASATIC SOLID CANCERS Inclusion: Exhausted conventional treatment	SR-8541A SR-8541A is an ENPP1 inhibitor. Further information: https://trials.cancervic.org.au/det-ails/vctl-nct06063681	Peninsula and Southeast Oncology	ag@paso.com.au
BAY3498264 Phase 1 Study of a SOS1 Inhibitor, BAY 3498264, in Combination in Participants With Advanced KRASG12C-mutated Solid Tumors	SOS1 KRAS	ADVANCED OR METASATIC SOLID CANCERS Inclusion: KRAS G12C mutation	BAY3498264 BAY3498264 is a SOS1 inhibitor. Treatment will be in combination with sotorasib. Further information: https://trials.cancervic.org.au/details/vctl/nct06659341	Peninsula and Southeast Oncology	ag@paso.com.au
AK138D1 A First-in-human, Phase I Study of Evaluating Safety, Tolerability, Pharmacokinetics and Preliminary Efficacy of AK138D1 in the Treatment of Advanced Solid Tumors	HER3	ADVANCED OR METASATIC SOLID CANCERS Inclusion: Exhausted conventional treatment	Patritumab Deruxtecan Patritumab Deruxtecan an anti- HER3 Antibody drug conjugate Further information: https://trials.cancervic.org.au/details/vctl/nct06730386	Peninsula and Southeast Oncology	ag@paso.com.au



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Trial Title	Targets	Target population	Treatment + Further	Site	Contact Details
	(Genomic)	(Trial suitable for patients with	Information	(Where the study	(Email the contact person listed
		this stage of pancreatic cancer)	(What the study involves)	is being offered)	with any enquiries)
DT-7012-CLI-001	CCR8	ADVANCED OR METASATIC SOLID	<u>DT-7012</u>	Peninsula and	ag@paso.com.au
		CANCERS		Southeast	
Study of DT-7012 as a Single				Oncology	
Agent and in Combination With		Inclusion:	DT-7012 is an anti-CCR8 antibody.		
an Immune Checkpoint Inhibitor		Second or later line			
in Participants With Advanced			Further information:		
Solid Tumors (DOMISOL)			https://clinicaltrials.gov/study/NC		
			T06819735		
<u>SNV4818</u>	PIK3CA	ADVANCED OR METASATIC SOLID	<u>SNV4818</u>	Monash Health	earlyphase.oncresearch@monashheal
		CANCERS			
A Phase 1, Open-Label Dose			SNV4818 is an oral PI3Kα		th.org
Escalation and Expansion Study of		Inclusion:	inhibitor. SNV4818 will be		
SNV4818 as Monotherapy or in		Activating PIK3CA mutation	delivered with or without		
Combination With Other		Exhausted conventional treatment	fulvestrant		
Anticancer Agents in Participants		Nata Dantum and a dan			
With Advanced Solid Tumors		Note: Pantumour for dose escalation cohorts	Further information:		
			NCT06736704 - Victorian Cancer		
<u>BM230</u>	HER2	ADVANCED OR METASATIC SOLID CANCERS	<u>BM230</u>	Monash Health	earlyphase.oncresearch@monashheal
A Phase I, Multicenter, Non-			BM230 is delivered		th.org
randomized, Open-label Study to		Inclusion:	subcutaneously each week for 3		
Evaluate the Safety, Tolerability,		HER2 1+ expression	weeks followed by fortnightly		
Pharmacokinetics, and Efficacy of			administration		
BM230 in Patients With Advanced					
Solid Tumors			Further information:		
			BM230 - Victorian Cancer Trials		
<u>BT317</u>		ADVANCED OR METASATIC SOLID	<u>BT317</u>	Monash Health	earlyphase.oncresearch@monashheal
		CANCERS			
A Phase I, First-in-human, Open-			BT317 is administered		th.org
label, Dose Escalation Study of		Inclusion:	intravenously		
the Safety, Tolerability,		Exhausted conventional treatment			
Pharmacokinetics, and			Further information:		
Immunogenicity of BNT317 in			NCT06750185 - Victorian Cancer		



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Trial Title	Targets	Target population	Treatment + Further	Site	Contact Details
	(Genomic)	(Trial suitable for patients with	Information	(Where the study	(Email the contact person listed
		this stage of pancreatic cancer)	(What the study involves)	is being offered)	with any enquiries)
Patients with Advanced Solid			<u>Trials Link</u>		
Tumors					
07 001 101	67470	ADVANCED OR METACATIC COLID	07 004	5	
<u>OZ-001-101</u>	STAT3	ADVANCED OR METASATIC SOLID CANCERS	<u>OZ-001</u>	Peninsula and South Eastern	ag@paso.com.au
A Phase 1, Open-label, First-in		CANCERS	OZ-001 is a small molecule dual	Haematology and	
Human Study to Examine the		Inclusion:	inhibitor of the STAT3 and T-type	Oncology Group	
Safety, Tolerability,		Exhausted conventional treatment	calcium channels	Oncology Group	
Pharmacokinetic Profile, and		Exhausted conventional treatment	calciant charmers		
Preliminary Efficacy of OZ-001		Confirmed accepting PDAC for	Further information:		
when Administered Orally in		phase 1a	https://trials.cancervic.org.au/det		
Adults with Solid Tumours with a			ails/vctl actrn12625000163404		
Focus on Triple Negative Breast					
Cancer					
INI-4001-101	TLR7/8	ADVANCED OR METASATIC SOLID	INI-4001	Cabrini	clinicaltrials@cabrini.com.au
		CANCERS			
An Open-label, Multiple-			INI-4001 is TLR7/8 agonist.		
Ascending Dose, Two-Part Dose		Inclusion:			
Ranging and Cohort Expansion		Exhausted conventional treatment	Further information:		
Study of INI-4001 in Patients with		Pre-screening component (review	https://trials.cancervic.org.au/det		
Advanced Solid Tumours		of medical history)	ails/vctl nct06302426		
		*Minimal slots available			



Trial Title	Targets	Target population	Treatment + Further	Site	Contact Details
	(Genomic)	(Trial suitable for patients with	Information	(Where the study	(Email the contact person listed
		this stage of pancreatic cancer)	(What the study involves)	is being offered)	with any enquiries)
LM350-01-10	CDH17	ADVANCED OR METASATIC SOLID	<u>LM350</u>	Peninsula and	ag@paso.com.au
		CANCERS		South Eastern	
A Phase I/II, First-in-Human (FIH),			LM350 is a CDH17 targeted	Haematology and	
Open-Label, Multiple Centre		Inclusion:	antibody drug conjugate	Oncology Group	
Clinical Study to Evaluate the		Exhausted conventional treatment			
Safety, Tolerability,			Further information:		
Pharmacokinetics,			https://clinicaltrials.gov/study/NC		
Immunogenicity and Preliminary			T07112222?aggFilters=status:not		
Efficacy of LM-350 in Patients					
with Advanced Solid Tumors					
RO7673396	RAS	ADVANCED OR METASATIC SOLID	RO7673396	Peter MacCallum	PCCTU.EDD@petermac.org
		CANCERS		Cancer Centre	
A Phase I Dose Escalation and			RO7673396 is an oral RAS		
Expansion Study to Evaluate the		Inclusion:	inhibitor		
Safety, Tolerability,		Exhausted conventional treatment			
Pharmacokinetics and Preliminary		Confirmed presence of RAS	Further information:		
Clinical Activity of RO7673396 as		mutation	https://trials.cancervic.org.au/det		
a Single Agent and in			ails/vctl nct06884618		
Combination With Other					
Anticancer Therapies in Patients					
With Advanced Solid Tumors					
Harboring RAS Mutation(s)					
AK146D1-102	TROP2	ADVANCED OR METASATIC SOLID	AK146D1	Austin Health	samantha.chakar@austin.org.au
	NECTIN4	CANCERS			
A Phase Ia Clinical Study to			AK146D1 is a an anti-		
Evaluate the Safety, Tolerability,		Exclusion:	TROP2/NECTIN4 bispecific		
Pharmacokinetics and Anti-tumor		Prior TROP2 or NECTIN4 targeting	antibody drug conjugate		
Efficacy of AK146D1 for Injection,		treatments			
an Anti-Trop2/Nectin4 Bispecific		Prior topoisomerase 1 inhibitor	Further information:		
Antibody-drug Conjugate, in		treatment	https://trials.cancervic.org.au/det		
Patients With Advanced Solid			ails/vctl nct06929663		
Tumors					



Trial Title	Targets (Genomic)	Target population (Trial suitable for patients with	Treatment + Further Information	Site (Where the study	Contact Details (Email the contact person listed
		this stage of pancreatic cancer)	(What the study involves)	is being offered)	with any enquiries)
IDOV-Immune for Advanced	N/A	ADVANCED OR METASATIC SOLID	<u>IDOV-Immune</u>	Alfred Health	act-m@alfred.org.au
Solid Tumors		CANCERS			
			IDOV-immune is an oncolytic		
A First-in-human, Phase I, Multi-		Inclusion:	virus therapy administered		
center, Open-label, Dose-		Exhausted conventional treatment	intravenously. Patients will		
escalation Study to Evaluate the			receive one dose of therapy.		
Safety, Tolerability,					
Pharmacokinetics,			Further information:		
Pharmacodynamics and			https://trials.cancervic.org.au/det		
Preliminary Evidence of			ails/vctl nct06910657?utm sourc		
Antitumor Activity of IDOV-			e=ccv-		
Immune in Adult Participants			website&utm_medium=website&		
With Advanced Solid Tumors			utm_campaign=api		



SUPPORTIVE CARE TRIALS FOR PATIENTS WITH PANCREATIC CANCER						
Trial Title	Targets (Genomic)	Target population (Trial suitable for patients with this stage of pancreatic cancer)	Treatment + Further Information (What the study involves)	Site (Where the study is being offered)	Contact Details (Email the contact person listed with any enquiries)	
DINE-PC	N/A	ALL STAGES OF PANCREATIC	<u>Dietetics counselling</u>	Epworth	EH-	
		CANCER		Victoria	PancreaticCentre@epworth.org.au	
Dietetics Intervention and			Further information:			
Nutritional Evaluation in			https://www.pancreaticcentr			
Pancreatic Cancer Care (DINE-PC)			e.org.au/treatment/clinical-			
			trials-and-research/dine-pc			
SuperQol:	N/A	ALL STAGES OF PANCREATIC	<u>Dietician</u>	Latrobe Regional	clinicaltrials@lrh.com.au	
		CANCEER	counselling/intervention	Hospital		
Assessing the impact of an				Victoria		
intensive dietitian-led			Patients will be randomised			
telehealth intervention			to receive usual care or			
focusing on nutritional			intensive dietetics			
adequacy and symptom			intervention			
control, on quality of life in						
patients with pancreatic			Further intervention:			
cancer: a randomised			https://trials.cancervic.org.au			
controlled trial			/details/vctl_actrn126240000			
			84583			



Phase 1 Trials are offered in Victoria at the following locations:

Study Site	Contact	Email	Phone
Alfred Hospital (Prahran VIC 3004)	Clinical Trial Team	$\frac{moncACT1@alfredhealthconnect.onmicrosoft.co}{\underline{m}}$	ТВА
Austin Hospital (Heidelberg VIC 3084)	Samantha Chakar	samantha.chakar@austin.org.au	03 9496 3088
Barwon Health (Geelong VIC 3220)	Karen Aitken	cstu.manager@barwonhealth.org.au	03 4215 2758
Cabrini Malvern (Malvern VIC 3144)	Rochelle Woods	rwoods@cabrini.com.au	95083437
Epworth Hospital (Richmond VIC 3121)	Clinical Research Coordinator	ehjreissaticentre@epworth.org.au	0448 842 680 or 03 9426 8880
PASO Medical (Frankston VIC 3199)	Albert Goikman Clinical Trials Manager	ag@paso.com.au	03 91131307
Monash Health (Clayton VIC 3168)	Early Phase Research Study Coordinator	earlyphase.oncresearch@monashhealth.org	0474 769 510
Peter MacCallum Cancer Centre (Parkville VIC 3052)	Enquiries Line Coordinator	clinicaltrials.enquiries@petermac.org	03 8559 7456 (9am-2pm, Mon-Fri)
St Vincent's Hospital (Fitzroy VIC 3065)	Nadia Ranieri	oncology.research@svha.org.au	03 9231 3167
Western Health (St Albans VIC 3021)	Heike Raunow	CancerClinicalTrials@wh.org.au	03 83959136 or 0434915739

For patients not eligible for any of the above trials, and who have pancreatic cancer that has progressed, are intolerant or are ineligible for standard of care treatments, please consider a phase 1 trial. Clinicians can email or phone the contact listed with any enquiries. Patients, please contact your treating Medical Oncologist if you wish to discuss phase 1 trials.



Trial Title	Targets (Genomic)	Target population (Trial suitable for patients with this stage of pancreatic cancer)	Treatment + Further Information (What the study involves)	Site (Where the study is being offered)	Contact Details (Email the contact person listed with any enquiries)
Porcupine P2EA An open-label study to assess the preliminary efficacy and safety of RXC004, (Zamaporvint), in patients with advanced pancreatic cancer who have progressed following therapy with current standard of care.	Porcupine	PANCREATIC CANCER - METASTATIC/LOCALLY ADVANCED (Second line) Inclusion: Progression on first line treatment for incurable disease Mandatory biopsy at enrolment	Zamaporvint Zamaporvint is an oral porcupine inhibitor. Treatment will be administered with denosumab Further information: https://anzctr.org.au/Trial/Registration/TrialReview.aspx?A https://anzctr.org.au/Trial/Registration/TrialReview.aspx?A CTRN=12624000588594p	Gold Coast University Hospital	Susan.Caird@health.qld.gov.au
A Randomized, Phase 2/3 Study Comparing BMS-986504 in Combination With Nab-paclitaxel and Gemcitabine Versus Placebo in Combination With Nab- paclitaxel and Gemcitabine in Participants With Untreated Metastatic Pancreatic Ductal Adenocarcinoma Harboring Homozygous MTAP Deletion	MTAP deletion	PANCREATIC CANCER - METASTATIC/LOCALLY ADVANCED Inclusion: Untreated advanced pancreatic cancer Homozygous MTAP deletion	Patients will receive gemcitabine and nabpaclitaxel combined with BMS-986504 (a novel PRMT5 inhibitor) or placebo. Further information: https://www.clinicaltrials.go v/study/NCT07076121	Royal Brisbane & Women's Hospital	amy.ives@health.qld.gov.au



Trial Title	Targets (Genomic)	Target population (Trial suitable for patients with this stage of pancreatic cancer)	Treatment + Further Information (What the study involves)	Site (Where the study is being offered)	Contact Details (Email the contact person listed with any enquiries)
Clarity-PT01 A Phase II, Open-label, Multicentre Study to Evaluate Safety, Tolerability, Efficacy, PK, and Immunogenicity of AZD0901 as Monotherapy and in Combination With Anti-cancer Agents in Participants With Advanced Solid	CLDN18.2	PANCREATIC CANCER - METASTATIC/LOCALLY ADVANCED Inclusion: Pancreatic cancer Treatment naïve (first line) CLDN18.2 positive	AZD0901 (antibody-drug conjugate) AZD0901 targets and binds to CLDN18.2, which is a protein on tumour cells, and then delivers cytotoxic agents which damage these cancer cells.	Prince of Wales Hospital (NSW)	SESLHD-POW- CTRUreferrals@health.nsw.gov.au
Tumours Expressing Claudin 18.2.		Exclusion: Exposure to prior CLDN18.2 targeted agents except anti- CLDN18.2 monoclonal antibody	Further information: https://www.anzctr.org.au/Tr ialSearch.aspx#&&searchTxt NCT06219941	Fiona Stanley Hospital (WA)	Chia.Tan@health.wa.gov.au
AMG193 20230223 A Phase 1b Study Evaluating the Safety, Tolerability, Pharmacokinetics, and Efficacy of AMG 193 in Combination with Other Therapies in Subjects with Advanced Gastrointestinal, Biliary	MTAP deletion	PANCREATIC CANCER - METASTATIC/LOCALLY ADVANCED Inclusion: Pancreatic cancer Homozygous MTAP-deletion	AMG 193 AMG 193 is a PMRT5 inhibitor which is administered orally. AMG 193 will be administered with chemotherapy.	Genesis Care Norths Shore (NSW)	admin.northshore@genesiscare.com



Trial Title	Targets (Genomic)	Target population (Trial suitable for patients with this stage of pancreatic cancer)	Treatment + Further Information (What the study involves)	Site (Where the study is being offered)	Contact Details (Email the contact person listed with any enquiries)
Tract, or Pancreatic Cancers with Homozygous MTAP-deletion AMG20230223		Exclusion: Prior MAT2A inhibitor or PRMT5 inhibitor	Further information: https://www.cancer.nsw.gov.au/r esearch-and-data/cancer-clinical- trials-in-nsw/find-a-cancer- clinical-trial-in-nsw/item?r=13988	Chris O'Brien Lifehouse	jasmine.sell@lh.org.au
AMPLICITY (AMP945-202) A Phase 1b/2a, Multicenter, Open Label Study of the Safety, Efficacy and Pharmacokinetics of Narmafotinib in Combination With Modified FOLFIRINOX in Pancreatic Cancer Patients	FAK	PANCREATIC CANCER - METASTATIC/LOCALLY ADVANCED (First line) Inclusion: Treatment naïve for metastatic disease	Narmafotinib Narmafotinib is an oral FAK inhibitor. Narmafotinib will be administered in combination with mFOLFIRINOX chemotherapy Further information: AMPLICITY Trial	Genesis Care Norths Shore (NSW)	admin.northshore@genesiscare.com
A Randomized, Phase 2/3 Study Comparing BMS-986504 in Combination With Nab-paclitaxel and Gemcitabine Versus Placebo in Combination With Nab- paclitaxel and Gemcitabine in Participants With Untreated Metastatic Pancreatic Ductal Adenocarcinoma Harboring Homozygous MTAP Deletion	МТАР	PANCREATIC CANCER - METASTATIC/LOCALLY ADVANCED (First line) Inclusion First line treatment of metastatic pancreatic cancer Homozygous MTAP deletion	BMS-986504 BMS-986504 is a selective MTA-cooperative PRMT5 inhibitor. Patients receive gemcitabine and nab- paclitaxel with or without BMS-986504 Further information: https://cancertrialswa.zepli.com. au/clinical-trials/90123	St John Of God Murdoch Hospital	Mel.Hernandez@sjog.org.au



Trial Title	Targets (Genomic)	Target population (Trial suitable for patients with this stage of pancreatic cancer)	Treatment + Further Information (What the study involves)	Site (Where the study is being offered)	Contact Details (Email the contact person listed with any enquiries)
CA233-0000/ BMS-986484 A Study of BMS-986484 Alone and Combination Therapy in Participants With Advanced Solid Tumors ALKOVE-1	CD40/FAP	ADVANCED OR METASTATIC SOLID CANCERS Exclusion: History of ILD	BMS-986484 BMS-986484 (a CD40/FAP bispecific agonist) is delivered as monotherapy or in combination with nivolumab Further information: https://www.cancer.nsw.gov.au/research-and-data/cancer-clinical-trials-in-nsw/find-a-cancer-clinical-trial-in-nsw/item?r=14232 NVL655	St Vincent's Hospital Darlinghurst (NSW) Lyell McEwin Hospital (SA)	Svhs.research@svha.org.au Health.NALHNCancerResearch@sa.sov.au PI: malinda.itchins@sydney.edu.au
A Phase 1/2 Study of the Selective Anaplastic Lymphoma Kinase (ALK) Inhibitor NVL-to655 in Patients with Advanced NSCLC and Other Solid Tumors (ALKOVE- to1)	ALK	ADVANCED OR METASTATIC SOLID CANCERS Inclusion: ALK rearrangement or activating ALK mutation	NVL655 NVL655 (neladalkib) is an oral selective ALK inhibitor. Further information: https://www.cancer.nsw.gov.au/research-and-data/cancer-clinical-trials-in-nsw/find-a-cancer-clinical-trial-in-nsw/item?r=13595	Royal North Shore Hospital NSW	Trial coordinator: shirley.liang@health.nsw.gov.au
AT-0174-001 A Phase I Study to Evaluate the Safety, Tolerability, Pharmacology, and Preliminary Efficacy of AT-0174 in Subjects with Advanced Solid Malignancies	IDO1/TDO 2	ADVANCED OR METASATIC SOLID CANCERS Inclusion: Progressed on available prior lines of therapy	AT-0174 AT-0174 is a novel dual inhibitor of IDO1/TDO2 Further information: https://www.anzctr.org.au/Trial	Royal North Shore Hospital (NSW)	PI: helen.wheeler@health.nsw.gov.au Trial Coordinator: wenqiong.yu@health.nsw.gov.au



Trial Title	Targets (Genomic)	Target population (Trial suitable for patients with this stage of pancreatic cancer)	Treatment + Further Information (What the study involves)	Site (Where the study is being offered)	Contact Details (Email the contact person listed with any enquiries)
BGB-58067 A Phase 1a/b Study Investigating the Safety, Tolerability, Pharmacokinetics, Pharmacodynamics, and Preliminary Antitumor Activity of BGB-58067, an MTA-Cooperative PRMT5 Inhibitor in Patients With Advanced Solid Tumors	MTAP deficiency	ADVANCED OR METASATIC SOLID CANCERS Inclusion: Any treatment line	BGB-58067 BGB-58067 is a PMRT5 inhibitor Further information: https://www.cancer.nsw.gov.au/r esearch-and-data/cancer-clinical-trials-in-nsw/find-a-cancer-clinical-trial-in-nsw/item?r=14130	Blacktown Cancer & Haematology Centre (NSW)	William.dAvigdor@health.nsw.gov. au
GeneScreen 5FU DPYD Genotype-guided dose Personalisation for Fluoropyrimidine prescribing in Cancer	DPYD	Inclusion: Intention to treat with Fluoropyrimidine (FP) containing chemotherapy Exclusion: Prior FP containing chemotherapy prior to study entry.	DPYD genotyping Pre-emptive DPYD genotyping prior to commencing Fluoropyrimidine chemotherapy Further information: ANZCTR - Registration	Lake Macquarie Private Hospital (NSW) Fiona Stanley Hospital (WA)	clinicalTrialsUnit.LMP@ramsayhealth .com.au audrey.margery- Muir@health.wa.gov.au
INCB161734 A Phase 1, Open-Label, Multicenter Study of INCB161734 in Participants With Advanced or Metastatic Solid Tumors With KRAS G12D Mutation	KRAS G12D	ADVANCED OR METASATIC SOLID CANCERS (Second line +) Inclusion: Second line and beyond ***Long waitlist***	INCB161734 INCB161734 in a KRAS G12D inhibitor Further information: https://www.cancer.nsw.gov.au/research-and-data/cancer-clinical-trials-in-nsw/find-a-cancer-clinical-trial-in-nsw/item?r=14020	St Vincent's Hospital Darlinghurst (NSW) Chris O'Brien Lifehouse	robert.kent@svha.org.au sarah.hing@lh.org.au
BAY3713372	MTAP loss	ADVANCED OR METASATIC SOLID CANCERS	BAY 3713372	Chris O'Brien Lifehouse	teresa.nicholls@lh.org.au



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Trial Title	Targets (Genomic)	Target population (Trial suitable for patients with this stage of pancreatic cancer)	Treatment + Further Information (What the study involves)	Site (Where the study is being offered)	Contact Details (Email the contact person listed with any enquiries)
A first in human study to evaluate the safety, tolerability, and pharmacokinetics, pharmacodynamics, and preliminary clinical activity of BAY 3713372, a novel 2 nd generation PRMT5 inhibitor in participants with MTAP deleted solid tumors		Inclusion Systemic Treatment naive or treatment exposed advanced pancreatic adenocarcinoma Homozygous MTAP deletion	BAY 3713372 is a novel 2 nd generation PRMT5 inhibitor. Further information: https://www.cancer.nsw.gov.au/research-and-data/cancer-clinical-trials-in-nsw/find-a-cancer-clinical-trial-in-nsw/item?r=14234		
BNT326-01 A Phase I/II, Open-label, Adaptive Two-part Trial to Evaluate the Safety, Efficacy, Optimal Dose and Pharmacokinetics of BNT326 as Monotherapy and in Combination With Cancer Immunotherapies in Participants With Advanced Solid Tumors	HER3	ADVANCED OR METASATIC SOLID CANCERS Inclusion Second or third line	BNT326 BNT326 is a HER3 targeted ADC with a topoisomerase I inhibitor. Further information: https://cancertrialswa.zepli.com.au/clinical-trials/80123	One Clinical Research, Hollywood Medical Centre, Nedlands WA	scott.mcgregor@oneclinicalresearch.c om.au
KANDLELIT-014 A Phase 2, Open-Label, Multicenter, Tumor-agnostic Study of MK-1084 as Monotherapy and in Combination With Cetuximab, in Participants With KRAS G12C-Mutant, Advanced Solid Tumors (KANDLELIT-014)	KRAS G12C	ADVANCED OR METASATIC SOLID CANCERS Inclusion Progression on standard of care therapy	MK-1084 MK-1084 is an oral KRAS G12C inhibitor. Patients will be treated alone or in combination with cetuximab. Further information: https://clinicaltrials.gov/stud y/NCT07209111	Prince of Wales Hospital, NSW	SESLHD-POW- CTRUreferrals@health.nsw.gov.au



PANCREATIC CANCER TRIALS IN SINGAPORE					
Trial Title	Targets (Genomic)	Target population (Trial suitable for patients with this stage of pancreatic cancer)	Treatment + Further Information (What the study involves)	Site (Where the study is being offered)	Contact Details (Email the contact person listed with any enquiries)
A phase II study of metronomic capecitabine, oxaliplatin and UGT1A1 genotype-directed irinotecan in metastatic pancreatic cancer patients.	NA	METASTATIC/LOCALLY ADVANCED Inclusion: Systemic Treatment naive or treatment exposed advanced pancreatic adenocarcinoma	Metronomic capecitabine, oxaliplatin and UGT1A1 genotype directed irinotecan Further information: https://clinicaltrials.gov/study/NC T05929885?term=NCT05929885 &rank=1b	National Cancer Centre Singapore	honey.shwe.sin@nccs.com.sg
A first in human study to evaluate the safety, tolerability, and pharmacokinetics, pharmacodynamics, and preliminary clinical activity of BAY 3713372, a novel 2 nd generation PRMT5 inhibitor in participants with MTAP deleted solid tumors	MTAP loss	ADVANCED OR METASATIC SOLID CANCERS Inclusion Systemic Treatment naive or treatment exposed advanced pancreatic adenocarcinoma Homozygous MTAP deletion	BAY 3713372 BAY 3713372 is a novel 2 nd generation PRMT5 inhibitor. Further information: https://clinicaltrials.gov/stud y/NCT06914128	National Cancer Centre Singapore	Wang.jue.lynn@nccs.com.sg
A phase I/II Dose-Escalation and expansion study evaluating the safety, pharmacokinetics, and activity of GDC-7035 as a single agent and in combination with other anti-cancer therapies in patients with advanced solid tumors with a KRAS G12D mutation	KRAS G12D	METASTATIC/LOCALLY ADVANCED Inclusion: Systemic Treatment refractory KRAS G12D pancreatic adenocarcinoma	GDC-7035 GDC-7035 is a KRAS G12D inhibitor	National Cancer Centre Singapore	Ye.xin@nccs.com.sg



PANCREATIC CANCER TRIALS IN SINGAPORE						
Trial Title	Targets (Genomic)	Target population (Trial suitable for patients with this stage of pancreatic cancer)	Treatment + Further Information (What the study involves)	Site (Where the study is being offered)	Contact Details (Email the contact person listed with any enquiries)	
PAUF-I A first in human phase I/2A, multicentre, open label study of evaluating the safety, tolerability, pharmacokinetics, and efficacy of PBP1510 in patients with advanced/metastatic pancreatic cancer	PAUF	METASTATIC/LOCALLY ADVANCED Inclusion: Systemic Treatment exposed pancreatic adenocarcinoma	PBP 1510 PBP 1510 is an anti-PAUF antibody Further information: https://clinicaltrials.gov/stud y/NCT05141149	National Cancer Centre Singapore	Goh.mui.leng@singhealth.com.sg	

PANCREATIC CANCER PREVENTION					
Trial Title	Targets (Genomic)	Target population (Trial suitable for patients with this stage of pancreatic cancer)	Treatment + Further Information (What the study involves)	Site (Where the study is being offered)	Contact Details (Email the contact person listed with any enquiries)
APRISE Assess the effectiveness of the Australian Pancreatic High-RIsk ScrEening Program in identifying early-stage pancreatic cancer among high-risk individuals due to familial or genetic risk factors	NA	FAMILY HISTORY OF PANCREATIC CANCER Inclusion: High risk features for pancreatic cancer development	Screening Screening with endoscopic ultrasound or MRI for a period of 10 years Further information:	Epworth Health	EHJreissatiCentre@epworth.org.au
			https://trials.cancervic.org.au/det ails/vctl actrn12624000421538_		