Anschutz | Cancer Center

Open Clinical Trials

November 2026

To refer a patient, please contact the disease specific nurse navigator directly to schedule a visit with one of our providers.

Please note that slot status may vary over time.

For more information regarding trials at the University of Colorado, please visit:
https://researchstudies.cuanschutz.edu/

Jump to Cancer Types

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- Phase 1, Expansion, Molecular Studies (POEMs)
- Radiation Oncology
- Sarcoma
- Thoracic (Lung)
- Cancer Prevention and Survivorship

Common Eligibility Criteria for Clinical Trials

Location Key	
AMC	Anschutz Medical Campus, Anschutz Cancer Pavilion, 1665 Aurora Ct, Aurora, CO 80045
CCMC	Cherry Creek Medical Center, 100 Cook St, Denver, CO 80206
HRH	Highlands Ranch Hospital, 1500 Park Central Dr, Highlands Ranch, CO 80129
LT	Lone Tree Medical Center, 9548 Park Meadows Dr, Lone Tree, CO 80124

Satellite Site Contacts

Breast Cancer Medical Oncology Anschutz Medical Campus (AMC)		
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Metastatic Breast Cancer			
Study Name	**Patients**	**NCT#**	**Sites**
PRE-I-SPY Phase 1/1b Platform, PRE-Investigation of Serial Studies (24-1242)	Previously treated metastatic breast cancer	NCT05868226	AMC
Phase 3 trial of saruparib (PARP1i) plus camizestrant (SERD) vs MD choice of CDK4/6 plus endocrine therapy (24-1711)	ER+, HER2-, BRCA1, BRCA2 or PALB2 mutations	NCT06380751	AMC
Phase 3 study of zanidatamab (HER2 bispecific ab) + standard therapy v. trastuzumab + standard therapy (24-1635)	Previously treated metastatic HER2+ breast cancer	NCT06435429	AMC, HRH, CCMC
CDK4/6 Inhibitor dosing knowledge (CDK) study (24-1798)	Metastatic breast cancer in older adults	NCT06377852	AMC
Phase 1/1b Study of IAM1363 (HER2/HER2 mutant TKI, CNS penetrant) (25-0333)	Advanced Cancers Harboring HER2 tyrosine kinase domain mutations or HER2+	NCT06253871	AMC (POEMs)
A Phase 2 study of alisertib in combination with endocrine therapy in patients with HR+, HER2-negative recurrent or metastatic breast cancer (24-1686)	HR+, HER2 – metastatic adenocarcinoma of breast not amenable to curative therapy. Progression with at least 2 prior lines of endocrine therapy, no prior chemotherapy	NCT06369285	AMC, HRH, CCMC
A Phase 1 First-in-Human, Open-Label, Multicenter Study of OP-3136 (KAT6A/KAT6B inhibitor) in Adult Participants with Advanced or Metastatic Solid Tumors (25-2398)	Advanced/metastatic ER+ HER2- breast cancer	NCT06784193	AMC
A Phase 3 Randomized, Double-Blind, Active-Controlled Study of Palazestrant (Oral SERD and complete estrogen receptor antagonist) with Ribociclib (CDK4/6) Versus Letrozole (Aromatase inhibitor) with Ribociclib for the First-Line Treatment of ER+HER2 Advanced Breast Cancer (OPERA-02) (25-2524)	1L ER-positive, HER2-negative locally advanced or metastatic breast cancer	NCT07085767	AMC, LT, HRH, CCMC
A Phase 1b/2 Multicenter, Open-label, Randomized Study Evaluating the Safety, Pharmacokinetics, and Activity of	Advanced/Metastatic ER+/HER2- breast cancer	NCT07100106	AMC

GDC-4198 Alone and in Combination with Giredestrant in Comparison with Abemaciclib and Giredestrant in Participants with Locally Advanced or Metastatic ER Positive, HER2 Negative Breast Cancer who have Previously Progressed During or After a CKD4/6 Inhibitor (MoonROSE) (25-1535)			
A Phase 1b/2, Multicenter, Open-label, Dose-Finding Study to Evaluate the Safety and Antitumor Activity of Patritumab Deruxtecan in Participants with HER2 Positive Unresectable Locally Advanced Breast Cancer or Metastatic Breast Cancer (HERTHENA-Breast-01) (25-1063)	Advanced/Metastatic HER2+ breast cancer	NCT06686394	AMC
Early-Stage Breast Cancer			
ISPY2.2 neoadjuvant study (10-0374)	Stage II-III Breast cancer	NCT01042379	AMC, LT, HRH, CCMC
OFSET: Phase 3 study of chemotherapy v. ovarian suppression (23-2461)	Premenopausal HR+HER2- breast cancer	NCT05879926	AMC, LT, HRH, CCMC
OptimICE-PCR: De-escalation after pCR (23-1255)	TNBC with pCR to neoadjuvant therapy	NCT05812807	AMC, LT, HRH
S2206: Phase 3 trial of neoadjuvant durvalumab (PD-L1i) plus chemo vs chemo alone (24-0401)	MammaPrint ultrahigh ER/PR+, HER2-	NCT06058377	AMC, LT, HRH, CCMC
S2212 SCARLET: Shorter Anthracycline-Free Chemo Immunotherapy in Early TNBC (23-2438)	Early Stage TNBC	NCT05929768	AMC, CCMC LT HRH
A012303: ShortStop-HER2: Shortened Duration of Adjuvant Therapy in Patients with Early-Stage HER2+ Breast Cancer Who Achieve pCR After Neoadjuvant Chemotherapy with HER2 Blockade (25-2075)	Clinical Stage T1c-T3 N0-N1 (except T3N1) with pCR after neoadjuvant therapy (HER2+)	NCT06876714	AMC, CCMC LT HRH

Cutaneous Oncology Anschutz Medical Campus (AMC)		
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High-Risk Resected Melanoma No current trials available			
Advanced Melanoma			
Study Name	**Patients**	**NCT#**	**Sites**
Phase 2/3 study of LNS8801 (GPER agonist) +/- without pembrolizumab (25-0336)	Treatment refractory, unresectable melanoma	NCT06624644	AMC
TILVANCE-301: Phase 3 trial of Lifileucel (LN-144 tumor infiltrating lymphocyte) + pembro vs pembro alone (24-0982)	Untreated, unresectable or metastatic melanoma	NCT05727904	AMC
Phase 1 study of mRNA-4359 + pembrolizumab in advanced solid tumors (22-1490)	1L NSCLC PD-L1 TPS $\geq$ 50% 1L PD-L1+ melanoma 2L PD-L1+ melanoma	NCT05533697	AMC (POEMs)
Phase 3 trial of fianlimab + cemiplimab vs relatlimab (LAG-3i) and nivolumab (24-1713)	Unresectable or metastatic melanoma	NCT06246916	AMC
SUPRAME: Phase 3 study of IMA203 (autologous T-cell receptor-engineered T cell therapy targeting PRAME) versus investigator's choice in patients with melanoma (25-0844)	Unresectable or previously treated cutaneous melanome; HLA-A*02:01 positive, Disease progression (resistance or toxicity) on at least 1 PD-1 inhibitor	NCT06743126	AMC
Lipid lowering therapy for atherosclerotic disease in melanoma patients receiving immune checkpoint inhibitors (25-1921)	Melanoma diagnosis without prior Immune Checkpoint Inhibitor use, going on Immune Checkpoint Inhibitor as SOC therapy	NCT07704918	AMC
A First-in-human Study of RLY-8161 (NRAS Inhibitor) for Treatment of Advanced NRAS-Mutant Melanoma and Other Solid Tumors (26-0306)	Unresectable Stage III or IV melanoma that is refractory to standard/ intolerant or declined therapy	NCT07584226	AMC
Neoadjuvant or Resectable Melanoma/			
Comparing the Effectiveness of Less Invasive Procedures With the Current Standard of Care for the Treatment of Older Adults With Early-Stage Melanoma	Newly diagnosed cutaneous melanoma with WLE and SNL biopsy indicated pending surgical management or risk of SNL metastasis of at least 5%. Pts must at least 75 yo	NCT07068074	AMC
Uveal Melanoma			
OptimUM-10: Phase 3 study of neoadjuvant darovasertib (PKCi) in patients with non-metastatic uveal melanoma (25-2370)	Non-metastatic uveal melanoma	NCT07015190	AMC
TARGET-Tebe: A Phase II Study of KIMMTRAK (bispecific T -cell engager) in HLA-A*0201 positive untreated metastatic uveal melanoma (25-2068)	Untreated metastatic uveal melanoma positive for HLA-A*0201	NCT06070012	AMC

REVEAL: Phase 2/3 Study of RP2 (oncolytic virus) + nivolumab v. ipilimumab + nivolumab (25-0004)	Immune checkpoint inhibitor-naïve metastatic uveal melanoma; no more than 2 lines of systemic therapy	NCT06581406	AMC
Merkel Cell Carcinoma, Squamous Cell Carcinoma			
Clinical trial of IFx-Hu2.0 (cancer vaccine) added to pembro in IO naïve merkel cell carcinoma (25-0580)	Advanced or metastatic merkel cell carcinoma	NCT06947928	AMC
Phase 1 study of BL-M14D1 (DLL3-ADC) in metastatic small cell lung cancer and other neuroendocrine neoplasms (26-0037)	SCLC, LCNEC, NEPC, GEP-NEC, EP-NECs, MCC or other poorly differentiated and/or high-grade neuroendocrine neoplasms with evidence of DLL3 expression who have failed at least 1 line of standard therapy	NCT07080242	AMC (POEMs)
NRG14-HN014: Phase 3 trial of neoadjuvant IO with response adapted treatment v. SOC for resectable Stage III/IV cutaneous squamous cell carcinoma (CSCC) (25-1924)	Invasive CSCC or regional lymph node or in-transit metastasis of CSCC	NCT06568172	AMC (RT)
A Randomized Phase II Study of Amivantamab (JNJ-61186372) (Bispecific anitbody targeting EGFR and MET) and hyaluronidase (rHuPH20) versus Cetuximab in Immunocompromised Participants with Recurrent Inoperable or Metastatic Cutaneous Squamous Cell Carcinoma (26-1077)	Metastatic Cutaneous Squamous Cell Carcinoma	NCT07042295	AMC
Advanced Cutaneous Malignancies in Transplant Recipients			
No current trials available			

Gastrointestinal Oncology Anschutz Medical Campus (AMC)		
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Gastric, Gastroesophageal, and Gastroesophageal Junction (GEJ) Cancers			
Study Name	**Patients**	**NCT#**	**Sites**
Phase 3 study of rilvegostomig (bispecific ab) + fluoropyrimidine + trastuzumab deruxtecan v. trastuzumab, chemotherapy + pembrolizumab (ARTEMIDE-Gastric01/25-0646) (25-0646)	1L HER2 gastric cancer	NCT06764875	AMC
Phase 1 study of SGN-CEACAM5C (CEACAM5 ADC Topoi) advanced solid tumors including CRC, GC/GEJ, pancreatic cancer (23-2329)	2L+ advanced CRC, GC/GEJ or pancreatic cancer	NCT06131840	AMC
Phase 1 study of TJ033721 (Givastomig, Claudin 18.2 x 4-1BB bispecific ab) in metastatic gastric, GE and esophageal adenocarcinoma (21-3254)	1L metastatic gastric, GE, and esophageal adeno ca positive for CLDN18.2	NCT04900818	AMC
CHS-114-102: Phase 1b study of CHS-114 (anti-CCR8ab) + toripalimab +/- other treatments (25-1556)	2L Gastric, GEJ and Esophageal Adenocarcinoma and 2L esophageal squamous cell carcinoma	NCT06657144	AMC
Phase 1/1b Study of IAM1363 (HER2/HER2 mutant TKI, CNS penetrant) (25-0333)	Advanced Cancers Harboring HER2 tyrosine kinase domain mutations or HER2+	NCT06253871	AMC (POEMs)
Phase I study of DPTX3186 (Wnt/B-catenin inhibitor) Wnt-Pathway Activated Solid Tumors (25-2381)	Solid Tumors known to be Wnt-Pathway activated (Gastric, TNBC, CRC) with no other approved treatment options available	NCT07312903	AMC (POEMs)
BL-M05D1-ST-101: A Phase 1 study of BL-M05D1 (Claudin 18.2 ADC Topoi) in advanced solid tumors (25-1878)	2L + Gastric, GEJ or Esophageal Adenocarcinoma, 2L+ biliary cancers	NCT07021066	AMC
EA2234: A Randomized Phase II/III Trial of Intraperitoneal Paclitaxel Plus Systemic Treatment vs Systemic Treatment Alone in Gastric Carcinomatosis - STOPGAP II (25-2369)	MSS gastric or GEJ Adenocarcinoma (cyt+) or peritoneal carcinomatosis. Must have received min 3 months and max 6 months of 1L systemic treatment	NCT07001748	AMC

GCT1184-09 (RAINFOL-09): A Phase 2, Open-label, Multicohort, Study of Rinatabart Sesutecan (Rina-S) (ADC) in Participants with Advanced Gastrointestinal (GI) Cancers (26-0771)	Advanced or metastatic gastric or GEJ adenocarcinoma that received at least 1 line but no more than 3 prior lines of therapy in advanced/metastatic setting or Esophageal squamous cell cancer that has recieved at least 1 but not more than 2 lines including platinum- based chemo and an immune checkpoint inhibitor	NCT07539311	AMC
Early-Stage Colorectal Cancer (CRC)			
CIRCULATE-US: Colon adjuvant chemotherapy based on evaluation of residual disease (22-0916)	Stage IIIA/B MSS colon adenocarcinoma following curative resection	NCT05174169	AMC, LT, HRH, CCMC
Phase 2 Single-Arm Rectal Cancer brachytherapy for patients with low-lying residual adenocarcinoma after total neoadjuvant therapy to improve organ preservation rates (25-1202)	MMR-Proficient low-mid rectal adenocarcinoma cT1-4, N0-2, M0	NCT07292298	AMC (RT Group)
M24-534: Colorectal Cancer: A Phase 2, Randomized, Open-Label Study Comparing Telisotuzumab Adizutecan (ABBV-400) (ADC targeting CMET) Monotherapy to the Current Standard of Care in Subjects with Post Adjuvant ctDNA Positive Colorectal Cancer and No Radiographic Evidence of Disease (NED) (25-1723)	Post Adjuvant (at least 3 months of platinum-based chemotherapy in adjuvant or neoadjuvant setting) ctDNA Positive Colorectal Cancer and No Radiographic Evidence of Disease	NCT07023289	AMC
Metastatic Colorectal Cancer (CRC)			
SMT112-3005: A Randomized, Active-Controlled, Double-blind, Multicenter, Phase 3 Clinical Study of Ivonescimab (PD1 and VEGF) in Combination with FOLFOX versus Bevacizumab in Combination with FOLFOX for the First-line Treatment of Metastatic Colorectal Cancer (HARMONi-GI3) (25-2600)	1L metastatic CRC	NCT07228832	AMC, HRH, LT, CCMC
Phase I trial of AWN003694 + cetuximab + encorafenib (25-0300)	Refractory BRAF V600E metastatic CRC	NCT06102902	AMC
Study of RAS (ON) inhibitors in GI tumors in combination with standard therapies (24-1326)	1-3L metastatic RAS-mutated CRC, 1-2L metastatic RAS-mutated pancreatic cancer	NCT06445062	AMC, LT
OrigAMI-3: Phase 3 study of FOLFIRI + amavantamab v cetuximab/bevacizumab in metastatic CRC (24-2380)	2L metastatic CRC KRAS/NRAS and BRAF WT	NCT06750094	AMC

Phase 1 study of INBRX-109 (DR5 ab) in metastatic solid tumors (18-1672)	Ewings sarcoma/ 2nd Line CRC	NCT03715933	AMC (POEMs)
INTRINSIC: Phase 1/1b umbrella study of targeted therapies in metastatic CRC (21-3774)	1L (untreated) metastatic CRC with a KRAS G12C mutation	NCT04929223	AMC
CARABiNER: Phase 1 study of GCC19CART in metastatic CRC. *This trial requires hospitalization and inpatient delivery of study drug for some cycles* (22-0697)	Previously treated metastatic CRC, GCC+, MSS	NCT05319314	AMC
Phase 1 study of SGN-CEACAM5C (CEACAM5 ADC Topoi) advanced solid tumors including CRC, GC/GEJ, pancreatic cancer (23-2329)	2L+ advanced CRC, GC/GEJ or pancreatic cancer	NCT06131840	AMC
AE2222: PUMP: Phase 3 study of systemic therapy +/- hepatic arterial infusion for unresectable CRC liver mets (24-0029)	Unresectable liver metastatic CRC	NCT05863195	AMC
Phase 1 study of JNJ-89402638 (bispecific ab) for metastatic CRC (24-1600)	3L+ unresectable metastatic CRC	NCT06663319	AMC
Phase 1 study of BI1831169 (oncolytic virus) +/- ezabenlimab (anti-PD-1) in advanced solid GI tumors (23-2048)	Advanced or metastatic GI tumors with no standard therapies	NCT05155332	AMC
Phase 1 study of PF-07934040 (KRASi) in patients (24-1423)	Select advanced solid tumors with KRAS mutations	NCT06447662	AMC (POEMs)
CHS-114-102: Phase 1b study of CHS-114 (anti-CCR8) + toripalimab with or without other treatments in participants with advanced or metastatic solid tumors (25-1556)	4L + CRC	NCT06657144	AMC
Phase I study of DPTX3186 (Wnt/B-catenin inhibitor) Wnt-Pathway Activated Solid Tumors (25-2381)	Solid Tumors known to be Wnt-Pathway activated (Gastric, TNBC, CRC) with no other approved treatment options available	NCT07312903	AMC (POEMs)
DSP107_003: Phase 2b study of DSP107 (bi-specific 4-1BB T-cell engager) + atezolizumab v. fruquintinib in advanced microsatellite stable CRC (26-0039)	MSS or pMMR CRC progressed or intolerant to 5FU, oxaliplatin, irinotecan, anti-VEGF and anti-EGFR (if RAS WT). No more than 3 prior lines of therapy	NCT07235293	AMC, HRH
MRG007-001/ARR-217-101: Phase 1 study of MRG007 (ARR-217, CDH17 ADC) in advanced solid tumors (25-2495)	Advanced or metastatic solid tumor that has failed or intolerant to standard therapy. Phase 1b is CRC Only	NCT07066657	AMC
DSP107_003: A Randomized, Open-label, Phase 2b Study to Compare the Efficacy of DSP107 (CD47 x CD137) in Combination with Atezolizumab (PD-L1) Versus Fruquintinib (VEGFR TKI) in Patients with Advanced Microsatellite Stable Colorectal Cancer (26-0039)	Advanced or metastatic MSS/pMMR CRC that has failed or intolerant to standard therapy including 5FU, Irinotecan, Oxaliplatin, Lonsurf, Bev and mutational targeting therapies if applicable	NCT07235293	AMC, HRH

A Phase 1 First-in-human Open-label Study Evaluating the Safety, Tolerability, Pharmacokinetics, Pharmacodynamics and Preliminary Efficacy of ERAS-0015 (pan-RAS) Monotherapy and in Combination in Patients with Advanced Solid Tumors (26-0780)	Locally Advanced or metastatic NSCLC, BTC, CRC or PDAC who have progressed on standard therapy. No prior RAS therapy allowed	NCT06983743	AMC (POEMs)
Protocol 10716, A Phase 1 and Randomized Phase 2 Trial of Pidnarulex (CX-5461) and Cemiplimab (REGN2810) in Refractory Microsatellite Stable Colorectal Cancer (26-1297)	Advanced or metastatic MSS/pMMR CRC that has failed or intolerant to standard therapy	NCT07147231	AMC
Liver Cancer			
Phase Ib/2 study of multiple immunotherapy based treatment combinations (24-1787)	Locally advanced/metastatic /unresectable HCC; no priory treatment	NCT04524871	AMC, HRH, CCMC
Phase 1b/2 study of TTI-101 (STAT3i) in advanced or metastatic HCC (22-1621)	1-3L Unresectable, locally advanced or metastatic HCC	NCT05440708	AMC
Biliary Tract Cancer			
Protocol 10608, A phase II trial of durvalumab with gemcitabine and cisplatin as neoadjuvant therapy for high-risk resectable intrahepatic cholangiocarcinoma (24-0573)	Resectable intrahepatic cholangiocarcinoma	NCT06050252	AMC
D702NC00001: Phase III, Randomized, Open-label, Global, Multicenter Study of Rilvegostomig (PD-1/TIGIT) or Durvalumab (PD-L1) in Combination With Chemotherapy as a First-line Treatment for Patients With Advanced Biliary Tract Cancer (ARTEMIDE-Biliary02) (26-0190)	Previously untreated advanced/ unresectable or metastatic biliary tract cancer	NCT07221253	AMC
A Phase 1/2 Study to Evaluate the Safety and Efficacy of MK-2870 Monotherapy or in Combination With Other Anticancer Agents in Gastrointestinal Cancers (24-1250)	Cohort 3 – 2L Biliary Tract Cancer Cohort 4 – 1L Biliary Tract Cancer	NCT06428409	AMC, HRH, CCMC
A Phase 1 First-in-human Open-label Study Evaluating the Safety, Tolerability, Pharmacokinetics, Pharmacodynamics and Preliminary Efficacy of ERAS-0015 (pan-RAS) Monotherapy and in Combination in Patients with Advanced Solid Tumors (26-0780)	Locally Advanced or metastatic NSCLC, BTC, CRC or PDAC who have progressed on standard therapy. No prior RAS therapy allowed	NCT06983743	AMC (POEMs)
Protocol 10716, A Phase 1 and Randomized Phase 2 Trial of Pidnarulex (CX-5461) and Cemiplimab (REGN2810) in Refractory Microsatellite Stable Colorectal Cancer (26-1297)	Unresectable and or metastatic MSI-S or pMMR colorecctal adenocarcinoma for which standard curative or palliative measures do not exist or no longer effective	NCT07147231	AMC
Pancreatic Cancer			

Intraperitoneal paclitaxel for pancreatic ductal adenocarcinoma (24-2111)	Pancreatic Ductal Adenocarcinoma with Peritoneal Carcinamatosis	NCT07030283	AMC
Study of RAS (ON) inhibitors in GI tumors in combination with standard therapies (24-1326)	1-3L metastatic RAS-mutated CRC, 1-2L metastatic RAS-mutated pancreatic cancer	NCT06445062	AMC, LT
Phase 1 study of SGN-CEACAM5C (CEACAM5 ADC Topoi) advanced solid tumors including CRC, GC/GEJ, pancreatic cancer (23-2329)	2L+ advanced CRC, GC/GEJ or pancreatic cancer	NCT06131840	AMC
Phase 1 study of CA-4948 (IRAK4/FLT3i) + gemcitabine+ nab-paclitaxel in metastatic pancreatic cancer (23-0613)	2-3L metastatic or unresectable pancreatic ductal adeno	NCT05685602	AMC
A Phase 1A/B Study to Evaluate the Safety and Tolerability Of EBC-129 (CEACAM5/6 ADC) As A Single Agent and In Combination with Pembrolizumab in Advanced Solid Tumors (23-0580)	Metastatic Pancreatic Adenocarcinoma	NCT05701527	AMC (POEMs)
MRG007-001/ARR-217-101: Phase 1 study of MRG007 (ARR-217, CDH17 ADC) in advanced solid tumors (25-2495)	Advanced or metastatic solid tumor that has failed or intolerant to standard therapy.	NCT07066657	AMC
223675: A Phase1b/2 study of GSK5764227 (B7-H3 ADC Topoi) alone and in combination in previously treated advanced unresectable or metastatic gastrointestinal solid tumors (25-1442)	2L Pancreatic adenocarcinoma (irinotecan naive)	NCT06885034	AMC
BL-M05D1-ST-101: A Phase 1 study of BL-M05D1 (Claudin 18.2 ADC Topoi) in advanced solid tumors (25-1878)	2L+ Pancreas Adenocarcinoma	NCT07021066	AMC
A Phase 1 First-in-human Open-label Study Evaluating the Safety, Tolerability, Pharmacokinetics, Pharmacodynamics and Preliminary Efficacy of ERAS-0015 (pan-RAS) Monotherapy and in Combination in Patients with Advanced Solid Tumors (26-0780)	Locally Advanced or metastatic NSCLC, BTC, CRC or PDAC who have progressed on standard therapy. No prior RAS therapy allowed	NCT06983743	AMC (POEMs)
GCT1184-09 (RAINFOL-09): A Phase 2, Open-label, Multicohort, Study of Rinatabart Sesutecan (Rina-S) (ADC) in Participants with Advanced Gastrointestinal (GI) Cancers (26-0771)	Pancreatic adenocarcinoma that has received at least 1 but no more than 2 lines with documented disease progression after gemcitabine and/or irinotecan based therapy	NCT07539311	AMC
A first-in-human study to evaluate the safety, tolerability and pharmacokinetics, pharmacodynamics and preliminary clinical activity of BAY 3713372, a novel 2nd generation PRMT5 inhibitor, in participants with MTAP-deleted solid tumors. (26-0950)	Locally advanced unrectable or metatatic pancreatic adenocarcinoma that has	NCT06914128	AMC

	progressed on at lease 1 line of systemic therapy		
Neuroendocrine Cancers			
COMPOSE: Study of peptide receptor radionuclide therapy with lutetium edotreotide v standard therapy for SSTR+ GI neuroendocrine tumors (21-3552)	Well-differentiated aggressive, grade 2/3, SSTR+ neuroendocrine tumors of gastroenteric or pancreatic origin	NCT04919226	AMC
NET Retreat: Phase 2 study of 177Lutetium retreatment v. everolimus in metastatic midgut neuroendocrine (23-1917)	Metastatic grade 1/2 midgut neuroendocrine tumors	NCT05773274	AMC
Phase 1 study of BL-M14D1 (DLL3-ADC) in metastatic small cell lung cancer and other neuroendocrine neoplasms (26-0037)	SCLC, LCNEC, NEPC, GEP-NEC, EP-NECs, MCC or other poorly differentiated and/or high-grade neuroendocrine neoplasms with evidence of DLL3 expression who have failed at least 1 line of standard therapy	NCT07080242	AMC (POEMs)

Genitourinary Oncology Anschutz Medical Campus (AMC)		
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Localized Prostate Cancer			
Study Name	**Patients**	**NCT#**	**Sites**
S2210: Neoadjuvant carboplatin for patients with localized prostate cancer and germline BRCA1/2 mutations (23-2420)	Localized prostate cancer with germline BRCA1/2 mutation	NCT05806515	AMC
Pragmatic study of metformin v. lifestyle mods (19-1536)	Localized prostate cancer	NCT05515978	AMC
The Phase III `High Five Trial' Five Fraction Radiation For High-Risk Prostate Cancer (25-1121)	High Risk Prostate Adenocarcinoma	NCT05946213	AMC, HRH (RT)
Metastatic Prostate Cancer			
S1802: SST +/- surgery or radiation (18-2252)	Metastatic hormone sensitive prostate cancer	NCT03678025	AMC
KEYNOTE-365: Study of pembrolizumab combinations COHORT I: Carbo/Etop +/- pembro (21-3724)	De novo and treatment emergent neuroendocrine/small cell prostate cancer	NCT02861573	AMC
MK-5684-003: Phase 3 study of opevesostat v. abiraterone/enzalutamide (24-0350)	Metastatic castration-resistant prostate cancer previously treated with hormonal agent and taxane	NCT06136624	AMC, HRH, LT
MK-5684-004: Phase 3 study of opevesostat v. abiraterone/enzalutamide (24-0742)	Metastatic castration-resistant prostate cancer previously treated with hormonal agent	NCT06136650	AMC, HRH, LT
JNJ-87189401 and JNJ-78278343 (23-1990)	Metastatic castration-resistant prostate cancer	NCT06095089	AMC
Phase 1b/2a study of pocenbrodib alone or in combination with abiraterone acetate, olaparib, or 177Lu-PSMA-617 (24-2278)	Metastatic castration-resistant Prostate Cancer	NCT06785636	AMC
CCTG-PR26 - Randomized phase 3 clinical trial for the addition of docetaxel to androgen receptor pathway inhibitors (TRIPLE-SWITCH) (25-0680)	Metastatic castration-sensitive prostate cancer and suboptimal PSA response	NCT06592924	AMC CCMC LT HRH
Phase 3 study of pasritamig (JNJ-78278343), a T-cell-redirecting agent targeting human kallikrein 2 v best supportive care for metastatic castration-resistant prostate cancer (25-1528)	Metastatic castration-resistant Prostate Cancer	NCT07164443	AMC
Phase 1 study of BL-M14D1 (DLL3-ADC) in metastatic small cell lung cancer and other neuroendocrine neoplasms (26-0037)	SCLC, LCNEC, NEPC, GEP-NEC, EP-NECs, MCC or other poorly differentiated and/or high-grade neuroendocrine neoplasms with evidence of	NCT07080242	AMC (POEMs)

	DLL3 expression who have failed at least 1 line of standard therapy		
Alliance A032304 Radioligand Efficacy comparison by initial PSA-Response Outcome in Metastatic CRPC with Lutetium 177Lu PSMA RLT (RECIPROCAL) (26-0904)	Metastatic castration-resistant Prostate Cancer that has received at least one androgen receptor	NCT07200830	AMC
Bladder Cancer			
Phase 2 study of intravesicular instillation of TARA-002 (24-1264)	High grade non-muscle invasive bladder cancer	NCT05085990	AMC
EA8192 Preoperative durvalumab and/or chemo (21-3855)	Upper tract urothelial cancer	NCT04628767	AMC
MODERN: A032103 risk adapted adjuvant therapy - ctDNA testing and nivo+/- relatlimab or surveillance (24-0700)	Muscle invasive bladder cancer or upper tract urothelial cancer after surgery	NCT05987241	AMC
S1937: Eribulin + gemcitabine v. standard therapy (22-0927)	3L+ metastatic bladder cancer post PD1i	NCT04579224	AMC
NRG-GU015: Phase 3 trial of adaptive RT + chemo for muscle invasive bladder cancer (ARCHER) (25-2502)	CT2-T3,N0M0 Urothelial carcinoma of bladder	NCT07097142	AMC (RT)
S2427: Phase 2 study of bladder preservation with immunoradiotherapy after clinically meaningful response to neoadj therapy in muscle invasive bladder cancer (BRIGHT) (25-2505)	CT2-T4aN0M0 muscle invasive urothelial carcinoma of bladder	NCT07061964	AMC (RT)
A phase 1/2, first-in-human study to assess the safety, tolerability, pharmacokinetics, pharmacodynamics, and antitumor activity of AVZO-103, (Nectin4/Trop2 ADC), as a single agent and in combination therapy in patients with locally advanced or metastatic urothelial cancer or other solid tumors (26-0180)	Locally advanced or metastatic urothelial carcinoma and selected advanced/metastatic solid tumors – Urothelial Carcinoma, TNBC, SCCHN and Cervical Cancer	NCT07193511	AMC, LT (POEMs)
Kidney Cancer			
An interventional phase 1b/2 study to evaluate the safety and efficacy of pf-08634404 (pd-1 and vegf) monotherapy and in combination with other anticancer agents in adult participants with locally advanced or metastatic renal cell carcinoma(26-0406)	Locally advanced or metastatic RCC not amenable to curative surgery or radiation therapy that has not received prior systemic therapy. Immunotherapy after surgery is allowed if >12 months prior	NCT07227415	AMC, HRH, LT, CCMC
SAMURAI: Phase 2 study of SABR in patients receiving immunotherapy (24-2480)	Metastatic unresectable RCC	NCT05327686	AMC LT HRH CCMC
NEOPAX: Pembrolizumab + axitinib for RCC with IVC tumor thrombus (22-1669)	Neoadjuvant therapy for RCC with tumor thrombus	NCT05969496	AMC

HC-7366 + belzutifan for locally advanced or metastatic clear cell RCC (24-0745)	2L+ metastatic clear cell RCC	NCT06234605	AMC
STRIKE/A032201: Study of short term intensified pembrolizumab + tivozanib for high-risk renal cell carcinoma (25-0682)	Clear cell RCC following complete resection	NCT06661720	AMC HRH CCMC LT
A092204: Phase 2 study of cabozantinib + cemiplimab v cabozantinib in adolescents and adults with advanced adrenocortical cancer (25-2801)	Locally advanced unresectable or recurrent/metastatic adrenocortical carcinoma	NCT06900595	AMC
A Randomized, Double-Blind, Active-Control, Multicenter Phase 3 Trial of Casdatifan and Cabozantinib Versus Placebo and Cabozantinib in Patients With Advanced Clear Cell Renal Cell Carcinoma (25-1976)	2L Unresectable locally advanced or metastatic renal cell carcinoma with a primary clear cell component that has recieved anti-PD-1 oranti-PD-L1 as part of most recent regimen	NCT07011719	AMC, HRH, LT
Penile Cancer			
EA8134 InPACT: Neoadjuvant chemo + ILND v chemo +RT+ILND v ILND (17-1515)	Penile cancer	NCT02305654	AMC

Gynecologic Oncology Anschutz Medical Campus (AMC)		
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Ovarian Cancer - Frontline			
Study	**Patients**	**NCT#**	**Sites**
GOG-3068 HOTT HIPEC: Phase 3 study of HIPEC + cisplatin v no HIPEC at time of cytoreductive surgery followed by niraparib in stage III and IV ovarian cancer (23-2189)	Newly diagnosed stage III/IV ovarian, primary peritoneal, and fallopian tube cancer treated with neoadj chemo	NCT05659381	AMC
Ovarian Cancer - Maintenance			
UPROAR: Phase 2 study of maintenance mirvetuximab soravtansine (ADC) and olaparib in recurrent platinum sensitive, ovarian cancer (22-0384)	Recurrent platinum-sensitive ovarian, peritoneal, and fallopian tube cancer	NCT05887609	AMC
NRG-GY036: Phase 3 trial of one vs. two years maintenance olaparib, +/- bevacizumab (25-0399)	BRCA1/2 mutated or HRD+ ovarian cancer following response to 1L platinum-based chemotherapy	NCT06580314	AMC, UCH North
MK-2870-021: A Phase 3, Randomized, Open-label, Multicenter Study of Sacituzumab Tirumotecan (sac-TMT, MK-2870) Maintenance Treatment With or Without Bevacizumab Versus Standard of Care in Participants With Newly Diagnosed Advanced non-HRD positive Ovarian Cancer Following First-line Platinum-based Chemotherapy (TroFuse-021/ENGOTov85/GOG-3102) (26-0408)	FIGO 2014 Stage III or Stage IV epithelial ovarian, primary peritoneal or fallopian tube carcinoma with one of the following histologies: high-grade serous, high-grade endometroid, clear cell carcinoma, or malignant mixed Mullerian tumor with high-grade serous component. Has completed 1L platinum based chemotherapy with a response of SD, PR, CR or NED	NCT07318558	AMC, HRH
Ovarian Cancer - Recurrent			
A Phase 1b study of psilocybin assisted psychotherapy to address fear of recurrence in patients diagnosed with early-stage breast cancer and ovarian cancer in remission (23-1455)	Clinical stage 1 or 2, completed primary treatment, risk of recurrence at 10 years <20%	NCT06430541	AMC
Phase I study of CLK/DYRK (Cirtuvivint) + olaparib in BRCA/HRD platinum resistant ovarian cancer (24-1749)	Platinum-resistant ovarian cancer with 1-3 prior lines of therapy. BRCA or HRD + with prior parp inhibitor.	NCT06856499	AMC
Phase 1/2 study of MBRC-101 in select advanced solid tumors (23-2430)	Metastatic refractory Ovarian Adenocarcinoma	NCT06014658	AMC (POEMs)

CORT125134-557: An Open-label, Global, Multi-Arm Study to Evaluate the Efficacy and Safety of Relacorilant (GR antagonist/modulator) in Combination with Different Treatment Regimens in Patients with Gynecological Cancers (BELLA) (26-0057)	Platinum-sensitive epithelial ovarian, peritoneal or fallopian-tube carcinoma	NCT06906341	AMC
GOG-3124/AbbVie M25-219 (ADC) - Gynecologic Oncology: A Phase 1b dose escalation and expansion trial for adult subjects with gynecologic cancers. (25-2205)	Platinum-sensitive ovarian, fallopian tube and primary peritoneal cancer with disease progression after platinum combination SOC therapy	NCT07024784	AMC
FRAmework-01: A Three -Part Phase 3 Study of Sofetabart Mipitecan (LY4170156) (ADC) versus Chemotherapy or Mirvetuximab Soravtansine (ADC) in Platinum-Resistant Ovarian Cancer, and Sofetabart Mipitecan plus Bevacizumab versus Platinum-Based Chemotherapy plus Bevacizumab in Platinum-Sensitive Ovarian Cancer (25-2695)	Platinum-sensitive ovarian, fallopian tube and primary peritoneal cancer with disease progression on or after most recent line of systemic therapy	NCT07213804	AMC, HRH
A Phase 1 Multicenter, Open-Label, Dose-Escalation and Expansion Study of NTB-928, a FR Targeting Bispecific T cell Engager, in Subjects with Relapsed/Refractory Ovarian Carcinoma (26-1183)	Relapsed or refractory ovarian cancer, including high-grade serous, high-grade endometrioid, low-grade serous, or clear cell ovarian cancer, or carcinosarcoma of the ovary that has already received standard treatments known to provide benefit	NCT07690189	AMC
Ovarian Cancer – Low Grade			
No current studies available			
Endometrial Cancer - Presurgical			
No current studies available			
Endometrial Cancer - Frontline			
NRG GY032: Phase 2 study of tailored adjuvant therapy in POLE-mutated p53 WT early-stage endometrial cancer (24-1417)	1L Stage I-III endometrial cancer POLE-mutated, completed surgery	NCT06388018	AMC
NRG GY035: Randomized, Phase 3 Trial of Carboplatin, Paclitaxel, Pembrolizumab Versus Carboplatin, Paclitaxel, Bevacizumab Versus Carboplatin, Paclitaxel, Pembrolizumab, Bevacizumab in the Treatment of pMMR, TP53 Mutated Advanced or Recurrent Endometrial Cancer (26-0636)	Stage III and Stage IVA Endometrial Cancers with Measurable Disease, Stage IVB Endometrial Cancer (with or without measurable disease), Recurrent Endometrial Cancer (with or without measurable disease. pMMR and TP53 mutation.	NCT07198074	AMC, HRH
Endometrial Cancer - Recurrent			

GOG-3082: Phase 1b/2 basket study of ACR-368 (CHK1/2i) +/-gemcitabine in platinum-resistant ovarian, endometrial and urothelial ca based on acrivon oncosignature (23-0115)	Up to 2L in recurrent setting: Recurrent, platinum-resistant ovarian cancer, endometrial adenoca, urothelial carcinoma with Acrivol Oncosignature Status	NCT05548296	AMC
Phase 1 study of BG-C9074 (B7H4 ADC) +/- tislelizumab (24-1128)	Previously treated metastatic HR+HER2- or TNBC, endometrial, or ovarian cancer	NCT06233942	AMC (POEMs)
224035 (NERT): A Phase 1/2 open-label, multicenter study of oral GSK5460025 (NER) alone or in combination with other anti-cancer agents in adult participants with Mismatch Repair-deficient (dMMR) or Microsatellite Instability-High (MSI-H) solid tumors (25-2741)	Advanced Endometrial Cancer that has received at least 1 but no more than 3 lines of systemic therapy	NCT07213609	AMC
BNT323-01 Fern-EC-01 (GOG-3105, ENGOT-EN25): 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 (26-0302)	Recurrent endometrial cancer with HER2 IHC score of 1+, 2+ or 3+ that has recurrence after platinum therapy in metastatic setting or less than 12 months after completing in adjuvant therapy and received prior ICI treatment	NCT06340568	AMC
Cervical Cancer			
NRG-GY037: A Phase III Study of Induction Pembrolizumab and Chemotherapy Followed by Chemoradiation and Pembrolizumab vs Chemoradiation and Pembrolizumab Both Followed by Pembrolizumab for High Risk Locally Advanced Cervical Cancer (25-2515)	High Risk Locally Advanced Cervical Cancer	NCT07061977	AMC, HRH
A phase 1/2, first-in-human study to assess the safety, tolerability, pharmacokinetics, pharmacodynamics, and antitumor activity of AVZO-103, (Nectin4/Trop2 ADC), as a single agent and in combination therapy in patients with locally advanced or metastatic urothelial cancer or other solid tumors (26-0180)	Locally advanced or metastatic urothelial carcinoma and selected advanced/metastatic solid tumors – Urothelial Carcinoma, SCCHN and Cervical Cancer	NCT07193511	AMC, LT (POEMs)
Uterine Leiomyosarcoma			
GOG 3088: Phase 2 study of letrozole v. observation in uterine leiomyosarcoma (24-0041)	Newly diagnosed ER+ uterine leiomyosarcoma	NCT05649956	AMC, HRH
BRCA Mutation Carriers			
SOROC: Study of salpingectomy v. salpingo-oophorectomy to reduce risk of ovarian cancer in BRCA1 carriers (20-2079)	BRCA1 germline mutation carriers with no prior cancer diagnosis age 35-50	NCT04251052	AMC, HRH

Head and Neck Cancer Anschutz Medical Campus (AMC)		
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Head and Neck Cancer			
Study Name	**Patients**	**NCT#**	**Sites**
FORTIFI-HN010: Phase 2/3 study of ficerafusp alfa (BCA101) v. placebo + pembrolizumab (25-0122)	1L PD-L1-positive, recurrent or metastatic NHSCC	NCT06788990	AMC
eVOLVE-HNSCC: Phase 3 study of volrustomig v. observation in unresected locally advanced HNSCC (23-2332)	Unresected locally advanced HNSCC who have not progressed following definitive concurrent chemoRT	NCT06129864	AMC, HRH
Phase 1b study of losartan, pembrolizumab and SBRT in locoregionally recurrent refractory or oligometastatic HNSCC (24-1630)	locoregionally recurrent refractory or oligometastatic HNSCC	NCT06211335	AMC, HRH
NRG- HN010: Phase 2 study of docetaxel + trastuzumab v. T-DM1 for HER2+ salivary gland cancer (22-2363)	Recurrent, metastatic, or treatment-naïve, unresectable HER2+ salivary gland cancer	NCT05408845	AMC, HRH
A Phase 2 Study of ONC206 in Advanced Pheochromocytoma and Paraganglioma (25-1457)	Pheochromocytoma or paraganglioma that is unresectable and failed or declined SOC	NCT07282587	AMC
A phase 1/2, first-in-human study to assess the safety, tolerability, pharmacokinetics, pharmacodynamics, and antitumor activity of AVZO-103, (Nectin4/Trop2 ADC), as a single agent and in combination therapy in patients with locally advanced or metastatic urothelial cancer or other solid tumors (26-0180)	Locally advanced or metastatic urothelial carcinoma and selected advanced/metastatic solid tumors – Urothelial Carcinoma, SCCHN and Cervical Cancer	NCT07193511	AMC, LT (POEMs)

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Leukemia			
Study	**Patients**	**NCT#**	**Sites**
Phase 1, Open Label, Dose Escalation Study to Evaluate the Safety, Expansion, Persistence, and Preliminary Clinical Activity of Autologous CD64 CAR T cells in Patients with Relapsed and/or Refractory Acute Myeloid Leukemia (AML) (PI: Mat Angelos/25-0718)	Relapsed/refractory disease. ≥5% blasts in the bone marrow or peripheral blood by morphology, flow cytometry, or immunohistochemistry after a minimum of 1 cycle of a hypomethylating agent (HMA) and venetoclax (Ven) combination (Ven/HMA)	NCT07347418	AMC
Phase 3 Randomized, Double-blind, Placebo-controlled Studies Assessing Ziftomenib in Combination with Either Standard of Care Nonintensive (Venetoclax+Azacitidine) or Intensive (7+3) Therapy in Patients with Untreated NPM1 Mutated or KMT2A Rearranged Acute Myeloid Leukemia (PI: C. McMahon/25-1278)	Nonintensive Therapy Study (Ven+Aza) • NPM1-m, >75 years of age, Age <75 and unfit for IC based on comorbidities Intensive Therapy Study (7+3) • NPM1-m or KMT2A-r, Age >18, FLT3 wild-type or ITD ratio <0.05 OR ineligible to receive FLT3-targeted therapy (medically ineligible or mutation in which FLT3 inhibition is not SOC). Lack of access to an FLT3 inhibitor is not considered “ineligible” for FLT3-targeted therapy, Fit for IC per Investigator opinion	NCT07007312	AMC
Randomized, Sequential, Open-Label Study to Evaluate the Efficacy of IDH targeted/non-targeted versus non-targeted/IDH targeted Approaches in the Treatment of Newly Diagnosed IDH	Patients with newly diagnosed IDH1 or IDH2 mutated AML. Not a candidate for or refuses intensive induction therapy. Adequate organ function.	NCT05401097	AMC

mutated AML Patients Not Candidates for Intensive Induction Therapy (I-DATA Study) (PI: Angelos/23-2138)			
Phase 1b study to evaluate eganelisib as monotherapy and in combination with cytarabine in patients with relapsed/refractory acute myeloid leukemia (PI: McMahon, C/24-2332)	Relapsed/refractory AML; ECOG of ≤2; adequate organ function	NCT06533761	AMC
Phase I study of SNDX-5613 in patients with relapsed/refractory leukemias including those harboring an MLL/KMT2A gene rearrangement or nucleophosmin1 (NPM1) Mutation (PI: McMahon C/ 22-1050)	Relapsed/refractory AML; ECOG of ≤2; adequate organ function	NCT04065399	AMC
Phase I study of CD19 directed CAR T Cells in adult patients with B-Cell Acute Lymphoblastic Leukemia (B-ALL) with minimal residual disease (MRD) positivity at first complete remission (PI: Schwartz/22-0054)	Ages ≥ 18; B-cell ALL in first complete morphologic remission; MRD positive; ECOG of ≤2	NCT05535855	AMC
Study of Mitoxantrone for venetoclax-resistant AML (PI: Kent/24-0178)	Non-APL AML by WHO criteria and have been treated with first-line venetoclax/HMA (azacitidine or decitabine)	NCT06429449	AMC
A Phase 1 Multi-Center, Open-Label Study to Evaluate the Safety, Tolerability, Pharmacokinetics, and Preliminary Anti- Leukemia or anti-MDS Activity of RPT1G in Patients with Acute Myeloid Leukemia (AML) and Higher-Risk (HR) Myelodysplastic Syndromes/Neoplasms (MDS) (PI: Pollyea/25-2789)	Histological confirmation of AML (ELN 2022 criteria) that relapsed and/or refractory AML or high-risk MDS as defined by International Consortium for MDS (icMDS) 2023 criteria that have received appropriate standard of care therapy, in the opinion of the investigator or declined receipt of these.	NCT07107126	AMC
KO-MEN-007: Phase 1 study of venetoclax/azacitidine or venetoclax + ziftomenib (KO-539) or standard induction cytarabine/daunorubicin (7+3) + ziftomenib for AML (PI: McMahon/23-0061)	Ages ≥ 18; documented NPM1 mutation or KMT2A rearrangement and have either newly diagnosed or relapsed/refractory AML; ECOG of ≤2	NCT05735184	AMC
A First-in-Human Study of the Safety, Pharmacokinetics, and Pharmacodynamics of JNJ-88549968, a T-cell Redirecting Bispecific Antibody for CALR-mutated Myeloproliferative Neoplasms (PI: Anglos/25-0787)	Diagnosis of either ET or MF as defined by the 2022 WHO criteria. Adequate organ function	NCT06150157	AMC
Myelodysplastic Syndrome Trials			
A Phase 1 Multi-Center, Open-Label Study to Evaluate the Safety, Tolerability, Pharmacokinetics, and	Histological confirmation of AML (ELN 2022 criteria) that relapsed and/or refractory AML or high-	NCT07107126	AMC

Preliminary Anti- Leukemia or anti-MDS Activity of RPT1G in Patients with Acute Myeloid Leukemia (AML) and Higher-Risk (HR) Myelodysplastic Syndromes/Neoplasms (MDS) (PI: Pollyea/25-2789)	risk MDS as defined by International Consortium for MDS (icMDS) 2023 criteria that have received appropriate standard of care therapy, in the opinion of the investigator or declined receipt of these.		
A Global Multicenter, Double-Blind, Randomized, Registrational Phase 3 Study of Lisaftoclax (APG-2575) in Combination with Azacitidine (AZA) in Patients with Newly Diagnosed Higher Risk Myelodysplastic Syndrome (HR-MDS) (GLORA-4) (PI: Kent/26-0427)	Newly diagnosed higher-risk MDS. ECOG score of ≤2. Adequate organ function.	NCT06641414	AMC
RALLY-MF: A Phase 1b/2 Open-Label Study to Evaluate the Safety, Tolerability, Pharmacokinetics, Pharmacodynamics, and Clinical Activity of DISC-0974 in Participants with Myelofibrosis or Myelodysplastic Syndrome and Anemia (PI: McMahon, Brandon/25-1397)	DIPSS score of 3 to 4 (intermediate 2 risk) or ≥5 (high-risk) primary MF, post PV MF, and/or post ET MF, as confirmed in the most recent local bone marrow biopsy report, according to WHO 2016 criteria	NCT05320198	AMC
Research On Nicotinamide Riboside Supplement Support in MDS (ROSS Trial)	Subject must have confirmation of high risk CCUS (defined by CHRS score) or the diagnosis of MDS in the lower risk category as defined by IPSS-M less than 0; adequate organ function	NCT07579429	AMC
Lymphoma			
A Phase 3, randomized controlled trial of Rondecabtagene autoleucel, an autologous, dual-targeting CD 19/CD20 CAR-T cell product candidate, vs. Investigator's choice of CD19 CAR-T cell therapy in patients with relapsed or refractory large b-cell lymphoma in the second-line setting (PINACLE-H2H)(PI: Kamdar/25-2042)	relapsed or refractory large b-cell lymphoma in the second-line setting; adequate organ function	NCT07188558	AMC
A Study of the Efficacy and Safety of Lisocabtagene Maraleucel (Liso-cel) as First-Line Therapy in Adults with Transplant-Ineligible Primary Central Nervous System Lymphoma (PI: Haverkos/25-1099)	Histologically confirmed primary central nervous system (CNS) lymphoma (PCNSL) prior to screening, as assessed by local pathology; transplant ineligible	NCT07015242	AMC
BAF_FRontier-1, A First-in-Human, Phase 1Trial to Assess Safety, Tolerability, and Preliminary Efficacy of LY4152199, a B-cell Activation Factor Receptor (BAFF-R) T-Cell Engager Bispecific Antibody, in Adult	Must have a diagnosis of either follicular lymphoma or diffuse large B-cell lymphoma. Participants with lymphoma must have at least 1 bi-dimensionally measurable lesion or in the absence of measurable	NCT07101328	AMC

Participants with Previously Treated B-cell Malignancies (PI: Kamdar/25-2677)	lymphadenopathy, documentation of bone marrow involvement. Adequate organ function.		
Phase I study of patient-derived multi-tumor associated antigen specific T-cells (MT-601) in patients with relapsed/refractory non- hodgkin lymphoma (PI: Kamdar/23-0056)	Diagnosis of NHL, HL or CLL; ECOG or zero or 1; adequate organ function	NCT05798897	AMC
Phase 3 interim response adapted trial comparing standard therapy with immuno-oncology therapy for children and adults (PI: Milgrom/24-2065)	Newly diagnosed Stage I and II classic Hodgkins Lymphoma	NCT05675410	AMC (RT Group)
S2308: Randomized Phase 3 study of mosunetuzumab vs. rituximab for low tumor burden follicular lymphoma (PI: Kamdar/ 25-0024)	Participants must have a histologically confirmed diagnosis of Classic Follicular Lymphoma (cFL). Involvement of no more than 3 nodal or extra nodal sites with diameter greater than 3 cm.	NCT06337318	AMC
A Phase 1a/1b trial in relapsed/refractory T-cell non-hodgkin lymphoma to determine the safety profile, pharmacology, and maximum tolerated dose of ST-001, a fenretinide phospholipid suspension (12.5 mg/mL) for intravenous infusion (PI: Haverkos/24-0458)	Ages > 18; ECOG of 0-1; diagnosed CTCL or Nodal TCL	NCT04234048	AMC
Phase Ib Study of mosunetuzumab or glofitamab + CC-220 and CC-99282 in B-Cell Non-Hodgkin Lymphoma (PI: Kamdar/22-0939)	Ages ≥ 18; ECOG of ≤2	NCT05169515	AMC
Phase 1/1b study of bispecific CD19 and CD22 chimeric antigen receptorCo-expressing T cells (CD19x22 CAR T) in adolescent and adult patients with relapsed and/or refractory B-NHL (PI: Kamdar/21-2578)	Ages ≥ 16; Relapsed or refractory aggressive B-cell NHL; ECOG of 0 or 1	NCT05098613	AMC
Phase 3 Study comparing the combination of beleodaq-CHOP or folotyn-COP to the CHOP regimen alone (24-2331)	Newly Diagnosed Patients with Peripheral T-Cell Lymphoma	NCT06072131	AMC
Multiple Myeloma			
A Phase 2, open-label, randomized, dose optimization study evaluating the efficacy and safety of belantamab mafodotin administered in combination with cyclophosphamide, bortezomib, and dexamethasone in adult participants with newly diagnosed amyloid light chain amyloidosis (ALANIS) (PI:Monge/26-0510)	Has histologically confirmed newly diagnosed primary AL amyloidosis. Measurable clonal disease. Not considered candidate for high-dose chemotherapy with autologous stem cell transplantation (ASCT) as part of first line of therapy. Adequate organ function.	NCT07224672	AMC
A Phase 3 Randomized Study Comparing JNJ-79635322 versus Teclistamab in Participants with	Documented diagnosis of multiple myeloma. Received 1 to	NCT07518186	AMC

Relapsed or Refractory Multiple Myeloma after 1 to 3 Prior Lines of Therapy, Including an Anti-CD38 Antibody and Lenalidomide (PI: Sherbenou/26-0679)	3 prior lines of antimyeloma therapy, including an anti-cluster of differentiation (CD) 38 antibody and lenalidomide. Adequate organ function.		
Phase 3 Study of Teclistamab in Combination With Lenalidomide and Teclistamab Alone versus Lenalidomide Alone in Participants With Newly Diagnosed Multiple Myeloma as Maintenance Therapy Following Autologous Stem Cell Transplantation MajesTEC-4 (PI: Monge/24-2397)	Newly diagnosed MM; post Auto stem cell transplant	NCT05243797	AMC
A phase 1b/2, open-label study of GC012F (AZD0120), a CD19/BCMA dual CART-cell therapy, in adult subjects with relapsed/refractory multiple myeloma remission (PI: Monge Urrea/24-2523)	Ages ≥ 18; Received at least three prior MM treatment lines of therapy; Have received as part of their previous therapy a PI and IMiD and an antiCD38 antibody	NCT05850234	AMC
Benign Heme			
Phase 3 study of bomedemstat (MK-3543/IMG-7289) v. best available therapy (24-2052)	Essential thrombocythemia who have an inadequate response to or are intolerant of hydroxyurea	NCT06079879	AMC
RALLY-MF: A Phase 1b/2 Open-Label Study to Evaluate the Safety, Tolerability, Pharmacokinetics, Pharmacodynamics, and Clinical Activity of DISC-0974 in Participants with Myelofibrosis or Myelodysplastic Syndrome and Anemia (PI: McMahon, Brandon/25-1397)	DIPSS score of 3 to 4 (intermediate 2 risk) or ≥5 (high-risk) primary MF, post PV MF, and/or post ET MF, as confirmed in the most recent local bone marrow biopsy report, according to WHO 2016 criteria	NCT05320198	AMC
A First-in-Human Study of the Safety, Pharmacokinetics, and Pharmacodynamics of JNJ-88549968, a T-cell Redirecting Bispecific Antibody for CALR-mutated Myeloproliferative Neoplasms (PI: Mat Angelos/25-0787)	Positive for a calreticulin (CALR) driver mutation of essential thrombocythemia (ET) or myelofibrosis (MF	NCT06150157	AMC
A Randomized, Double-Blind, Placebo-Controlled Phase 2/3 Study of BLU-263 in Indolent Systemic Mastocytosis (PI: Duarte/25-1016	Participant has confirmed diagnosis of ISM, confirmed by Central Pathology Review	NCT04910685	AMC
Allo Transplant/GVHD			
No active trials			

Neuro-Oncology Anschutz Medical Campus (AMC)		
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Neuro-Oncology			
Study	**Patients**	**NCT#**	**Sites**
Phase I Study of [177Lu] Lu-NeoB in combination with radiotherapy and temozolomide in newly diagnosed glioblastoma and as a single agent in recurrent glioblastoma (24-0082)	Recurrent GMB	NCT05739942	AMC (CReST)
GBM AGILE: Phase 2/3 platform trial evaluating multiple regimens for glioblastoma (19-0744)	Newly diagnosed GBM	NCT03970447	AMC
A071401 Phase 2 study of SMO/AKT/NF2 inhibitors in progressive meningiomas (15-1747)	Progressive meningiomas with AKT1, PIK3CA or PTEN mutations	NCT02523014	AMC
Protocol 10505, A phase 1 and randomized phase 2 trial of selinexor and temozolomide in recurrent glioblastoma (23-0099)	Glioblastoma (IDH wild-type, MGMT promoter methylated) that has undergone resection or bx upon 1st recurrence. Has received 1L treatment of temozolomide + rt	NCT05432804	AMC
A Phase 2, multicenter, clinical study to evaluate the efficacy and safety of safusidenib erbumine in patients with isocitrate dehydrogenase 1 (IDH1)-mutant glioma (25-1162)	newly diagnosed astrocytoma (Grade 2-4) following standard temozolomide therapy, as well as select patients with Grade 3 oligodendroglioma who have experienced progression/recurrence after surgical resection.	NCT05303519	AMC

Phase 1, Expansion, Molecular Studies (POEMs) Anschutz Medical Campus (AMC)		
Nurse Navigator	**Email**	**Phone**
Contact Disease Site Specific Nurse Navigator (ie Breast Cancer, GI, GU, etc.)		
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POEMs Studies			
Study	**Patients**	**NCT#**	**Sites**
Phase I study of RO7673396 (pan-RASi) as single agent and in combination with other anticancer therapy (25-0860)	PDAC RAS mutated	NCT06884618	AMC (POEMs)
Phase 1 study of INBRX-109 (DR5 ab) in metastatic solid tumors (18-1672)	Ewings sarcoma/ 2nd Line CRC	NCT03715933	AMC (POEMs)
Phase 1 study of mRNA-4359 + pembrolizumab in advanced solid tumors (22-1490)	Melanoma	NCT05533697	AMC (POEMs)
Phase 1 study of BG-C9074 (B7H4 ADC Topoi) +/- tislelizumab (24-1128)	Endometrial and platinum cancer no prior Topoi ADC	NCT06233942	AMC (POEMs)
Phase 1/1b Study of IAM1363 (HER2/HER2 mutant TKI, CNS penetrant) (25-0333)	Advanced Cancers Harboring HER2 tyrosine kinase domain mutations or HER2+	NCT06253871	AMC (POEMs)
Phase 1b, Multicenter, Open-label, Dose Escalation and Dose Expansion Study of RMC-6291 in Combination with RMC-6236 in Participants with Advanced KRASG12C-Mutated Solid Tumors (24-0554)	KRASG12C mutated NSCLC	NCT06128551	AMC (POEMs)
Phase I study of DPTX3186 (Wnt/B-catenin inhibitor) Wnt-Pathway Activated Solid Tumors (25-2381)	Solid Tumors known to be Wnt-Pathway	NCT07312903	AMC (POEMs)
Phase I study of BG-C477 (CEACAM5 ADC Topoi) in Patients with Selected Advanced Solid Tumors (25-2371)	NSCLC, PDAC, requires prescreening	NCT06596473	AMC (POEMs)
Phase 1 study of BL-M14D1 (DLL3-ADC) in metastatic small cell lung cancer and other neuroendocrine neoplasms (26-0037)	SCLC	NCT07080242	AMC (POEMs)
A phase 1/2, first-in-human study to assess the safety, tolerability, pharmacokinetics, pharmacodynamics, and antitumor activity of AVZO-103, (Nectin4/Trop2 ADC), as a single agent and in combination therapy in patients with locally advanced or metastatic urothelial cancer or other solid tumors (26-0180)	Locally advanced or metastatic urothelial carcinoma and selected advanced/metastatic solid tumors – Urothelial Carcinoma, SCCHN and Cervical Cancer	NCT07193511	AMC, LT (POEMs)

A Phase 1/2, Open-Label, Dose-Escalation and Expansion First-In-Human Study of ATX-295,an Oral Inhibitor of the Kinesin Motor Protein KIF18A, in Patients with Locally Advanced or Metastatic Solid Tumors, including High-Grade Serous Ovarian Cancer (26-0522)	Histologically confirmed solid tumors who have locally recurrent or metastatic disease, prioritizing HGSOC	NCT06799065	AMC (POEMs)
A Phase 1 First-in-human Open-label Study Evaluating the Safety, Tolerability, Pharmacokinetics, Pharmacodynamics and Preliminary Efficacy of ERAS-0015 (pan-RAS) Monotherapy and in Combination in Patients with Advanced Solid Tumors (26-0780)	Locally Advanced or metastatic NSCLC, BTC, CRC or PDAC who have progressed on standard therapy. No prior RAS therapy allowed	NCT06983743	AMC (POEMs)
A Phase 1a, Dose Escalation, Safety and Tolerability Study of NX-1607, a Casitas B-lineage lymphoma proto-oncogene (CBL-B) inhibitor, in Adults with Advanced Malignancies, with Phase 1b Expansion in Select Tumor Types (23-1337)		NCT05107674	AMC (POEMs)

Radiation Oncology Anschutz Medical Campus (AMC)		
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Radiation Oncology-Breast Cancer			
No current active studies			
Radiation Oncology-Colorectal Cancer			
Study	**Patients**	**NCT#**	**Sites**
Phase 2 Single-Arm Rectal Cancer Brachytherapy for Patients with Low-Lying Residual Adenocarcinoma After Total Neoadjuvant Therapy to Improve Organ Preservation Rates (25-1202)	Residual Adenocarcinoma after total neoadjuvant therapy	NCT07292298	AMC (RT)
Radiation Oncology- Genitourinary Cancer			
NRG-GU015: Phase 3 trial of adaptive RT + chemo for muscle invasive bladder cancer (ARCHER) (25-2502)	CT2-T3,N0M0 Urothelial carcinoma of bladder	NCT07097142	AMC (RT)
S2427: Phase 2 study of bladder preservation with immunoradiotherapy after clinically meaningful response to neoadj therapy in muscle invasive bladder cancer (BRIGHT) (25-2505)	CT2-T4aN0M0 muscle invasive urothelial carcinoma of bladder	NCT07061964	AMC (RT)
The Phase III `High Five Trial' Five Fraction Radiation For High-Risk Prostate Cancer (25-1121)	High Risk Prostate Adenocarcinoma	NCT05946213	AMC, HRH (RT)
Radiation Oncology - Lymphoma			
Phase 3 interim response adapted trial comparing standard therapy with immuno-oncology therapy for children and adults (24-2065)	Newly diagnosed Stage I and II classic hodgkins lymphoma	NCT05675410	AMC
Radiation Oncology -Lung Cancer			
NRG-LU008: Phase 3 trial of primary lung SBRT followed by mediastinal chemo-RT locally advanced NSCLC (23-1258)	Locally advanced NSCLC	NCT05624996	LT, HRH
DURABLE: Delayed or upfront brain Radiotherapy (24-1615)	naïve lung cancer patients with asymptomatic or minimally symptomatic Brain metastases and ALK rearrangement	NCT05987644	AMC
Radiation Oncology-Pancreatic Cancer			
No current active studies			
Radiation Oncology –Neuro Oncology			
No current active studies			
Radiation Oncology-Multiple Cancer Types			
No current active studies			

Sarcoma Anschutz Medical Campus (AMC)		
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Sarcoma			
Study	**Patients**	**NCT#**	**Sites**
Phase 1 study of peposertib (M3814) and low-dose liposomal doxorubicin (25-0217)	Metastatic Leiomyosarcoma and Other Soft Tissue Sarcomas	NCT05711615	AMC
STRASS 2 (EA7211): Phase 3 study of neoadjuvant chemo followed by surgery v surgery alone high-risk retroperitoneal sarcoma (23-1625)	High risk retroperitoneal sarcoma (LPS/LMS)	NCT04031677	AMC
EA7222: Phase 3 trial of doxorubicin +/- pembrolizumab undifferentiated pleomorphic sarcoma (24-2043)	Undifferentiated pleomorphic sarcoma and related poorly differentiated sarcomas	NCT06422806	AMC
A Phase I/Ib Study of losartan in combination with sunitinib in the treatment of pediatric and adult patients with relapsed or refractory osteosarcoma (18-2740)	Osteosarcoma	NCT03900793	CHCO, AMC
Phase 1 study of INBRX-109 (DR5 ab) in metastatic solid tumors (18-1672)	Ewings sarcoma	NCT03715933	AMC (POEMs)
Phase I study of DPTX3186 (Wnt/B-catenin inhibitor) Wnt-Pathway Activated Solid Tumors (25-2381)	Solid Tumors known to be Wnt-Pathway-Desmoid, GIST	NCT07312903	AMC (POEMs)
A phase II study on efficacy and tolerability of weekly doxorubicin in elderly patients (24-0118)	Advanced or Metastatic Leiomyosarcoma of Soft Tissue	NCT07125183	AMC
Phase 1/2, open-label, dose escalation and expansion study of ADCE-D01, a humanized anti-human uPARAP antibody linked to a topoisomerase I inhibitor (24-2281)	Metastatic and/or Unresectable Soft Tissue Sarcoma	NCT06797999	AMC
Multi-cohort, open-label, Phase 1/2 study of DCC-3009 in participants with gastrointestinal stromal tumors (GIST) (24-2282)	GIST	NCT06630234	AMC
Phase 1 study of ziftomenib + imatinib (25-0466)	GIST after imatinib failure	NCT06655246	AMC
A Phase 1/2 Study of FOG-001 in Participants with Locally Advanced or Metastatic Solid Tumors (25-2227)	**Part 1a:** solid tumor with documented WPAM, at least 1 prior line of systemic therapy, progressed or was nonresponsive, or is unfit for	NCT05919264	AMC

	available therapies. Evaluable or measurable disease per RECIST. **Part 1d:** Confirmed desmoid tumor, ≥10% in the active component of the tumor per investigator assessment within 12 months prior to the first dose. Treatment naïve or recurrent/refractory disease following at least 1 line of therapy		
SARC046: A Phase II Trial of Nab-Sirolimus (mTOR inhibitor) in Patients With Progressing or Symptomatic Epithelioid Hemangioendothelioma (25-2550)	Epithelioid Hemangioendothelioma that is progressing or clinically symptomatic and is not a candidate for curative intent surgery	NCT07104331	AMC
Expanded access to bezuclastinib to be coadministered with sunitinib for patients with locally advanced, unresectable, or metastatic gastrointestinal stromal tumors (25-1975)	Locally advanced, metastatic and/or unreectable GIST intolerant to Imatinib or progressed on Imatinib	NCT06948955	AMC

Thoracic (Lung) Oncology Anschutz Medical Campus (AMC)		
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Early-Stage NSCLC			
Study	**Patients**	**NCT#**	**Sites**
V940-009: A Phase 3 Randomized Double-blind Study of Adjuvant Pembrolizumab With or Without V940 in Participants With Resectable Stage II to IIIB (N2) NSCLC not Achieving pCR After Receiving Neoadjuvant Pembrolizumab With Platinum-based Doublet Chemotherapy (INTerpath-009) (24-2314)	Neo-adjuvant (Stage II-IIIB(N2)) NSCLC	NCT06623422	AMC
NRG-LU008: Phase 3 trial of primary lung SBRT followed by mediastinal chemo-RT locally advanced NSCLC (23-1258)	Locally advanced NSCLC	NCT05624996	AMC, LT, HRH (RT Group)
A Phase 3, Randomized, Placebo-Controlled Study of Adjuvant Intismeran Autogene (mRNA cancer vaccine) Plus Subcutaneous Pembrolizumab and Berahyaluronidase Alfa (MK-3475A) or Intismeran Autogene (mRNA cancer vaccine) Monotherapy Versus Placebo in Participants With Completely Resected High-Risk Stage I Non-Small Cell Lung Cancer (INTerpath-014) (26-0727)	Stage 1 NSCLC with at least 1 of the following high-risk pathologic features: tumor size > 2cm, visceral pleural invasion, lymphovascular invasion or high-grade histology that has undergone complete surgical resection	NCT07513376	AMC
1st Line Metastatic NSCLC without actionable genomic alterations			
No current active studies			
PD-L1≥ 50%			
Phase 3 study of vonescimab v. pembrolizumab (25-0500)	First line Metastatic NSCLC with High PD-L1 expression	NCT06767514	AMC
Any PD-L1 Range			
M24-536: An Open-label Multi-Cohort Phase 1b/2 Study to Evaluate the Safety, Efficacy, and Optimal Dose of Telisotuzumab Adizutecan in Combination with Pembrolizumab in Advanced or Metastatic Non- Squamous NSCLC with No Prior Treatment for Advanced Disease and No Actionable Genomic Alterations (AndroMETa-Lung-536) (25-0149)	Advanced/Metastatic Non-Squamous NSCLC with No Prior Treatment for Advanced Disease and No Actionable Genomic Alteration	NCT06772623	AMC

A Phase Ib/II, multi-site, open-label, two-part trial to evaluate the efficacy, safety, pharmacokinetics, and recommended combination dose of BNT324 with BNT327 in participants with advanced lung cancer (25-1395)	1 L NSCLC	NCT06892548	AMC
Study of response of bony metastasis for patients on TKI or systemic treatment (19-0392)	NSCLC with completed molecular testing; receiving standard of care TKI or systemic therapy for actionable driver	NCT03958565	AMC, LT
Phase I study of BG-C477 (CEACAM5 ADC Topoi) in Patients with Selected Advanced Solid Tumors (25-2371)	NSCLC, PDAC, requires prescreening	NCT06596473	AMC (POEMs)
EGFR mutant NSCLC - 1st Line			
A Global, Phase 3, Randomized, Multicenter, Open-Label Study to Evaluate the Efficacy and Safety of Firmonertinib (EGRF TKI) Compared With Investigator's Choice of Osimertinib or Afatinib as First-Line Treatment in Participants Who Have Locally Advanced or Metastatic Non-Small-Cell Lung Cancer With Epidermal Growth Factor Receptor P-Loop and Alpha C-Helix Compressing (PACC) Uncommon Mutations (ALPACCA) (26-0358)	First-line EGFR PACC-mutated locally advanced or metastatic NSCLC	NCT07185997	AMC
A Phase 2/3 Randomized Study to Evaluate the Safety, Efficacy, and Optimal Dose of Telisotuzumab Adizutecan in Combination with Osimertinib as First-Line Treatment in Patients with Locally Advanced Unresectable or Metastatic EGFR-Mutated Non-Squamous Non-Small Cell Lung Cancer (25-1494)	Metastatic non-squamous NSCLC with classical EGFR mutation (Exon 19 Del or Exon 21 L858R). No prior EFGR TKI in locally advanced/metastatic setting	NCT07005102	AMC
EGFR mutant NSCLC - 2nd Line or beyond			
Systimmune: Phase 1 study of BL-B01D1 (EGFR-HER3 bispecific ADC topoisomerase payload) in metastatic NSCLC (23-2288)	Metastatic NSCLC with EGFR mutation	NCT05983432	AMC
ALK, ROS1, or RET rearranged NSCLC - 1st Line			
No current active studies			
ALK, ROS1, or RET rearranged NSCLC - 2nd Line or beyond			
A Phase 1/2, dose escalation and expansion study of TRI-611, an oral ALK molecular glue degrader in participants with advanced ALK-positive NSCLC (25-2822)	ALK positive NSCLC	NCT07491497	AMC
A Phase 1/2 Study of the Highly Selective ROS1 Inhibitor NVL-520 (ROS-1) in Patients with Advanced NSCLC and Other Solid Tumors (ARROS-1) (21-4641)	Any ROS1 + solid tumor other than NSCLC. For Adolescents older than 12 yo any ROS1 + solid tumor including NSCLC	NCT05118789	AMC

MET or HER2 altered NSCLC - 1st Line			
No current active studies			
MET or HER2 altered NSCLC - 2nd Line and Beyond			
SystImmune: Phase 1 study of BL-B01D1 (EGFR-HER3 bispecific ADC topoi payload) metastatic NSCLC (23-2288)	HER2 Exon 20 mutated metastatic NSCLC	NCT05983432	AMC
Phase 1/1b Study of IAM1363 (HER2/HER2 mutant TKI, CNS penetrant) (25-0333)	Advanced cancers harboring HER2 tyrosine kinase domain mutations or HER2+	NCT06253871	AMC (POEMs)
KRAS or BRAF altered NSCLC			
No current active studies			
KRAS G12C			
Phase 1b, Multicenter, Open-label, Dose Escalation and Dose Expansion Study of RMC-6291 in Combination with RMC-6236 in Participants with Advanced KRASG12C-Mutated Solid Tumors (24-0554)	KRASG12C mutated NSCLC	NCT06128551	AMC (POEMs)
A Phase 1a/1b Open-Label Study Evaluating the Safety, Tolerability, Pharmacokinetics, and Efficacy of BBO-8520 in Subjects with Advanced KRASG12C Mutant Non-Small Cell Lung Cancer - The ONKORAS-101 Study (25-1233)	Advanced NSCLC with KRAS G12C	NCT06343402	AMC
Allele Specific KRAS Inhibitors			
A Phase 1 First-in-human Open-label Study Evaluating the Safety, Tolerability, Pharmacokinetics, Pharmacodynamics and Preliminary Efficacy of ERAS-0015 (pan-RAS) Monotherapy and in Combination in Patients with Advanced Solid Tumors (26-0780)	Locally Advanced or metastatic NSCLC, BTC, CRC or PDAC who have progressed on standard therapy. No prior RAS therapy allowed. Must have KRAS G12X or G13X	NCT06983743	AMC (POEMs)
A Phase 2, Open-label Study of VS-7375, an Oral KRAS G12D (ON/OFF) Inhibitor, in Patients with Previously Treated, Advanced KRAS G12D-Mutated Non-Small Cell Lung Cancer (TARGET-D 202) (26-1574)	Locally advanced or metastatic NSCLC with confirmed KRAS G12D mutation. Must have received a platinum based chemo and an immune checkpoint inhibitor in advanced, non-resectable setting	NCT07659782	AMC
Pan-Ras Inhibitors			
A Phase 1 First-in-human Open-label Study Evaluating the Safety, Tolerability, Pharmacokinetics, Pharmacodynamics and Preliminary Efficacy of ERAS-0015 (pan-RAS) Monotherapy and in Combination in Patients with Advanced Solid Tumors (26-0780)	Locally Advanced or metastatic NSCLC, BTC, CRC or PDAC who have progressed on standard therapy. No prior RAS therapy allowed	NCT06983743	AMC (POEMs)

Metastatic NSCLC: ADCs, cytotoxics, radiation - 2nd Line and Beyond			
A Multicenter, Open-Label, Phase 1a/b First-in-Human Study to Investigate the Safety, Tolerability, Pharmacokinetics, Pharmacodynamics, and Preliminary Antitumor Activity of BG-C477 in Patients with Selected Advanced Solid Tumors (25-2371)	Phase 1A NSCLC with CEA high Expression treated with at least 2 prior lines	NCT06596473	AMC (POEMs)
A Phase Ib/II, multi-site, open-label, two-part trial to evaluate the efficacy, safety, pharmacokinetics, and recommended combination dose of BNT324 with BNT327 in participants with advanced lung cancer (25-1395)	2 L NSCLC	NCT06892548	AMC
Metastatic NSCLC: Immunotherapy agents - 2nd Line and Beyond			
No current active studies			
Metastatic NSCLC: Small Molecules - 2nd Line and Beyond			
No current active studies			
Small Cell Lung Cancer			
A Phase 1b/2 Open-label Study Evaluating different MK-6070 (DLL3 TriTAC) and Ifinatamab Deruxtecan (ADC) (MK-2400)-based regimens in First-line Extensive Stage Small Cell Lung Cancer (26-0320)	1L ES-SCLC	NCT07227597	AMC
MAVERICK (S1827): MRI brain surveillance +/- cranial irradiation small-cell lung (20-0359)	Small-Cell Lung Cancer	NCT04155034	AMC, HRH
MK-6070-002: 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 (25-0794)	Relapsed/Refractory Extensive-Stage SCLC	NCT06780137	AMC
SystImmune: Phase 1 study of BL-B01D1 (EGFR-HER3 bispecific ADC topoi payload) metastatic NSCLC (23-2288)	HER2 Exon 20 mutated metastatic NSCLC and SCLC	NCT05983432	AMC
Phase 1 study of BL-M14D1 (DLL3-ADC) in metastatic small cell lung cancer and other neuroendocrine neoplasms (26-0037)	SCLC, LCNEC, NEPC, GEP-NEC, EP-NECs, MCC or other poorly differentiated and/or high-grade neuroendocrine neoplasms with evidence of DLL3 expression who have failed at least 1 line of standard therapy	NCT07080242	AMC (POEMs)
A Phase Ib/II, multi-site, open-label, two-part trial to evaluate the efficacy, safety, pharmacokinetics, and recommended combination dose of BNT324 with BNT327 in participants with advanced lung cancer (25-1395)	1L and 2L+ SCLC	NCT06892548	AMC
A Randomized, Open-Label, Phase 3 Study of ZL-1310, a DLL3 Antibody-Drug Conjugate (ADC), Compared to	Metastatic or extensive-stage small cell lung cancer	NCT06179069	AMC, HRH

Investigator's Choice Therapy in Participants with Relapsed Small Cell Lung Cancer (25-2325)			
Mesothelioma and Thymic Malignancies			
No current active studies			

Cancer Prevention and Survivorship Anschutz Medical Campus (AMC)		
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Cancer Prevention and Survivorship			
Study	**Patients**	**NCT#**	**Sites**
IL-6 and IL-6R Genetic Polymorphisms in Immune-Related Adverse Events Caused by Cancer Immunotherapy (25-0049)	Metastatic solid tumor of any type; starting new regimen of anti-CTLA4, anti PD1, anti PD-L-1 or anti LAG-3 therapy alone or in combination with standard of care	**n/a**	AMC
NRG-CC015: Harnessing E-Mindfulness Approaches for Living After Breast Cancer---HEAL-ABC (25-2629)	Stage 0, 1, 2, or 3 non metastatic breast cancer who have completed all primary breast cancer treatments at least 6 months but no more than 5 years ago, Age 18-50	NCT06748222	AMC
A212102: Blinded Reference Set for Multicancer Early Detection Blood Tests (23-1087)	Various cohorts for multiple cancer types, gender and race cohorts.	NCT05334069	AMC

Satellite Site Contacts		
	Physician Leads	**Nurse Navigators**
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LT	Corbin Eule, MD (Corbin.Eule@CUAnschutz.edu)	Crystal Stribling (CrystalD.Stribling@uchealth.org)

Common Eligibility Criteria for Clinical Trials	
Performance Status	ECOG 0-1 Karnofsky Performance Status ≥ 60
Age	Age ≥ 18 years (ask for sarcoma trials)
Measurable disease	**Lesions that can be accurately measured in at least one dimension** with a minimum size requirement: • **For CT or MRI scans:** o The lesion must be **≥10 mm** in its longest diameter. o For lymph nodes, the short axis must be **≥15 mm** to be considered measurable. • **For physical exam or clinical measurement:** o The lesion must be **≥10 mm** in its longest dimension and should be measurable with a caliper or ruler. • **For ultrasound:** o Lesions can only be considered measurable if they are confirmed by CT or MRI measurements. **Non-Measurable Lesions:** • Lesions that do not meet the criteria above are classified as **non-measurable.** • Please note that bone lesions (unless there is soft tissue extension), pleural effusions, or ascites are **not** considered measurable disease.
Adequate organ and bone marrow function	**Bone marrow:** ANC ≥ 1.5, Hgb ≥ 9.0 (without recent transfusion), platelets ≥ 100 (without recent transfusion). **Hepatic:** AST and ALT < 2 x ULN, bilirubin < 2.0. **Renal:** CrCl ≥ 50 ml/min. **Cardiovascular:** No symptomatic congestive heart failure; no MI, CVA, or unstable angina in the past 6 months. **Pulmonary:** History of ILD requiring steroids (protocol dependent). **Neurologic:** If brain metastases are present, they must have been treated and not progressing.