

Blood and Bone Marrow Transplant

<u>Short Name</u>	<u>Comments</u>	<u>Type</u>	<u>Investigator</u>	<u>Contact</u>
BMT Cord Blood Access Protocol	A Multicenter Access and Distribution Protocol for Unlicensed Cryopreserved Cord Blood Units (CBUs) in Pediatric and Adult Patients with Hematologic Malignancies and Other Indications	Treatment	SHORE: Dr. Troy Quigg	Laura Paulsen laura.paulsen@spectrumhealth.org 616.486.6125
NYBB UCB	A Multicenter Safety Study of Unlicensed, Investigational Cryopreserved Cord Blood Units (CBUs) Manufactured by the National Cord Blood Program (NCBP) and Provided for Unrelated Hematopoietic Stem Cell Transplantation of Pediatric and Adult Patients	Treatment	SHORE: Dr. Troy Quigg	Laura Paulsen laura.paulsen@spectrumhealth.org 616.486.6125
2018-051 Cardiac Strain	Utility of Serial Echocardiogram Parameters to Assess Cardiotoxicity and Evaluate Cardio-Protective Therapy in Adult Blood and Marrow Transplant Patients (Adult BMT)	Other	SHORE: Dr. Sami Brake	Laura Paulsen laura.paulsen@spectrumhealth.org 616.486.6125
2019-118: Ruxolitinib EA	An Open-label, Expanded Access Program of Ruxolitinib for the Treatment of Graft vs Host Disease Following Allogeneic Hematopoietic Stem Cell Transplant	Treatment	SHORE: Dr. Sami Brake	Laura Paulsen laura.paulsen@spectrumhealth.org 616.486.6125
Kymriah EAP/MAP	Managed Access Program (MAP) to Provide Access to CTL019, for Acute Lymphoblastic Leukemia (ALL) or Large B Cell Lymphoma Patients with out of Specification Leukapheresis Product and/or Manufactured Tisagenlecleucel out of Specification for Commercial Release	Treatment	SHORE: Drs. Sami Brake, Ulrich Duffner, Troy Quigg	Laura Paulsen laura.paulsen@spectrumhealth.org 616.486.6125

GRAVITAS 309	A Phase 3 Study of Itacitinib or Placebo in Combination with Corticosteroids as Initial Treatment for Chronic Graft-Versus-Host-Disease	Treatment	SHORE: Dr. Sami Brake	Laura Paulsen laura.paulsen@spectrumhealth.org 616.486.6125
2013-024	10-CMSMDS-1 (MDS Study and MCR)	Other	SHORE: Dr. Troy Quigg	Josh Haveman josh.haveman@spectrumhealth.org 616.329.2724
2013-024	16-CMS-MF (Myelofibrosis Study and MCR)	Other	SHORE: Dr. Troy Quigg	Josh Haveman josh.haveman@spectrumhealth.org 616.329.2724
2013-024	17-CMS-SCD (Sickle Cell Study and MCR)	Other	SHORE: Dr. Troy Quigg	Josh Haveman josh.haveman@spectrumhealth.org 616.329.2724
2013-024	17-CMS-MM (Multiple Myeloma and Allo Transplant Study and MCR)	Other	SHORE: Dr. Troy Quigg	Josh Haveman josh.haveman@spectrumhealth.org 616.329.2724
	SC17-10 (ALL/PASS Study)	Treatment	SHORE: Dr. Troy Quigg	Josh Haveman josh.haveman@spectrumhealth.org 616.329.2724

SC16-040 (HL/Nivolumab)

Other

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Quigg

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Glioma Registry	Glioma Data Registry for Research	Registry	SHORE: Dr. Todd Vitaz	James Nelson james.nelson@spectrumhealth.org 616.267.8277
2015-175	NMTRC012 - PEDS-PLAN – Pediatric Precision Laboratory Advanced Neuroblastoma Therapy- A Pilot Study Using Molecular Guided Therapy with Induction Chemotherapy followed by maintenance with DFMO for Subjects with Newly Diagnosed High-Risk Neuroblastoma	Treatment	SHORE: Dr. Deanna Mitchell	Sarah Vos sarah.vos@helendevoschildrens.org 616.486.2064 Julie Steinbrecher julie.steinbrecher@helendevoschildrens.org 616.267.1163
2016-258	HEAD START 4 PROTOCOL: Newly Diagnosed Children (Less than 10 Years Old) With Medulloblastoma and Other Central Nervous System Embryonal Tumors. Clinical and Molecular Risk-Tailored Intensive and Compressed Induction Chemotherapy Followed by Consolidation with Randomization to Either Single-Cycle or to Three Sequential Cycles of Marrow-Ablative Chemotherapy with Autologous Hematopoietic Progenitor Cell Rescue	Treatment	SHORE: Dr. Albert Cornelius	Sarah Vos sarah.vos@helendevoschildrens.org 616.486.2064 Julie Steinbrecher julie.steinbrecher@helendevoschildrens.org 616.267.1163 Laura Paulsen laura.paulsen@spectrumhealth.org 616.486.6125
NMTRC 014-NMTT	NMTT- Neuroblastoma Maintenance Therapy Trial Using Difluoromethylornithine (DFMO)	Treatment	SHORE: Dr. Deanna Mitchell	Sarah Vos sarah.vos@helendevoschildrens.org 616.486.2064 Julie Steinbrecher julie.steinbrecher@helendevoschildrens.org 616.267.1163

	A Phase II Study of Metronomic and Targeted Anti-angiogenesis Therapy for Children with Recurrent/progressive Medulloblastoma	Treatment	SHORE: Dr. Rebecca Loret de Mola	<p>Sarah Vos sarah.vos@helendevoschildrens.org 616.486.2064</p> <p>Julie Steinbrecher julie.steinbrecher@helendevoschildrens.org 616.267.1163</p> <p>Laura Paulsen laura.paulsen@spectrumhealth.org 616.486.6125</p>
NRG BN003	Phase III Trial of Observation Versus Irradiation for a Gross Totally Resected Grade II Meningioma	Treatment	CRCWM Physician Investigators	<p>CRCWM (616) 391-1230 www.crcwm.org</p>
A071401	A Phase II Trial of SMO/AKT/NF2 Inhibitors in Progressive Meningiomas with SMO/AKT/NF2 Mutations	Treatment	CRCWM Physician Investigators	<p>CRCWM (616) 391-1230 www.crcwm.org</p>
ACNS1721	A Phase 2 Study of Veliparib (ABT-888, IND # 139199) and Local Irradiation, Followed by Maintenance Veliparib and Temozolomide, in Patients with Newly Diagnosed High-Grade Glioma (HGG) without H3 K27M or BRAFV600E Mutations	Treatment	CRCWM Physician Investigators	<p>CRCWM (616) 391-1230 www.crcwm.org</p>
ACNS1723	Dabrafenib Combined With Trametinib After Radiation Therapy in Treating Patients with Newly Diagnosed High Grade Glioma	Treatment	CRCWM Physician Investigators	<p>CRCWM (616) 391-1230 www.crcwm.org</p>
A071701	Genomically-guided Treatment Trial in Brain Metastases	Treatment	CRCWM Physician Investigators	<p>CRCWM (616) 391-1230 www.crcwm.org</p>
URCC 19085	A Phase II Clinical trial with Wireless Transcutaneous Electrical Nerve Stimulation (TENS) for Chemotherapy-induced Peripheral Neuropathy	Treatment	CRCWM Physician Investigators	<p>CRCWM (616) 391-1230 www.crcwm.org</p>

ACNS1422**A Phase II Study of Reduced Therapy for Newly Diagnosed
Average-risk WNT-Driven Medulloblastoma Patients**

Treatment

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Physician
InvestigatorsCRCWM
(616) 391-1230
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CCTG MA.39	Tailor RT: A Randomized Trial of Regional Radiotherapy in Biomarker Low Risk Node Positive Breast Cancer	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
NSABP B-51	A Randomized Phase III Clinical Trial Evaluating Post-Mastectomy Chestwall and Regional Nodal XRT and Post-Lumpectomy Regional Nodal XRT in Patients with Positive Axillary Nodes Before Neoadjuvant Chemotherapy Who Convert to Pathologically Negative Axillary Nodes After Neoadjuvant Chemotherapy	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
A011202	A Randomized Phase III Trial Comparing Axillary Lymph Node Dissection to Axillary Radiation in Breast Cancer Patients (cT1-3N1) Who Have Positive Sentinel Lymph Node Disease After Neoadjuvant Chemotherapy	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
NRG BR003	TRIPLE NEGATIVE; Adjuvant; HIGH RISK NODE(-)or NODE(+) / Invasive; A Randomized Phase III Trial of Adjuvant Therapy Comparing Doxorubicin Plus Cyclophosphamide Followed by Weekly Paclitaxel with or without Carboplatin	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
A231901CD	Improving Patient-Centered Communication in Breast Cancer: A RCT of A Shared Decision Engagement System (SHARES)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

S1418	TRIPLE NEGATIVE: A Randomized, Phase III Trial to Evaluate the Efficacy and Safety of MK-3475 (Pembrolizumab) as Adjuvant Therapy for Triple Receptor-Negative Breast Cancer with ≥ 1 CM Residual Invasive Cancer or Positive Lymph Nodes (ypN+) after Neoadjuvant Chemotherapy	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
S1703	Randomized Non-Inferiority Trial Comparing Overall Survival of Patients Monitored with Serum Tumor Marker Directed Disease Monitoring (STMDDM) versus Usual Care in Patients with Metastatic Hormone Receptor Positive Breast Cancer	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
AFT-38 "Patina"	A Randomized, Open Label, Phase III Trial to Evaluate the Efficacy and Safety of Palbociclib + Anti-HER2 Therapy + Endocrine Therapy vs. Anti-HER2 Therapy + Endocrine Therapy after Induction Treatment for Hormone Receptor Positive (HR+)/HER2 - Positive Metastatic Breast Cancer	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
S1501	A Phase III, Prospective Evaluation of Carvedilol in Prevention of Cardiac Toxicity in Patients with Metastatic Her-2+ Breast Cancer	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EA1151: TMIST	Tomosynthesis Mammographic Imaging Screening Trial	Other	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
NRG BR002	A Phase IIR/III Trial of Standard of Care Therapy With or Without Stereotactic Body Radiotherapy (SBRT) and/or Surgical Ablation for Newly Oligometastatic Breast Cancer	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

NRG BR004	A Randomized, Double-Blind, Phase III Trial of Paclitaxel/Trastuzumab/Pertuzumab with Atezolizumab or Placebo in First-Line HER2-Positive Metastatic Breast Cancer	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
ALLIANCE A221505	Phase III Randomized Trial of Hypofractionated Post Mastectomy Radiation with Breast Reconstruction	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
URCC 16070	Netupitant/Palonosetron Hydrochloride and Dexamethasone With or Without Prochlorperazine or Olanzapine in Improving Chemotherapy-Induced Nausea and Vomiting in Patients With Breast Cancer	Symptom Management	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
WF-97415	Understanding and Predicting Breast Cancer Events After Treatment (UPBEAT)	Prevention	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
AFT-25 "Comet"	AFT-25: A Phase III Prospective Randomized Trial Comparison of Operative to Monitoring and Endocrine Therapy (COMET) Trial for Low Risk DCIS	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
A211601	Evaluation of Mammographic Breast Density Effect of Aspirin: A Companion Study to Alliance Study A011502	Supportive Care	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

EA1181	EA1181 (CompassHER2-pCR): Preoperative THP and postoperative HP in patients who achieve a pathologic complete response	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
Bolt	BBI-20201001: A Phase I of BDC-1001 as a Single Agent and in Combination with Pembrolizumab in Patients with Advanced and HRE2-Expressing Solid Tumors	Treatment	START Midwest Investigators	START Midwest (616) 954-5554

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HAI for Pancreatic Mets	A Phase II Trial of Hepatic Artery Infusional Floxuridine with Systemic Chemotherapy in the Treatment of Pancreatic Cancer Liver Metastases	Treatment	SHORE: Dr. G. Paul Wright	Marianne Morrissey marianne.morrissey@spectrumhealth.org (616) 391-1129
EA2142	Gastroenteropancreatic Neuroendocrine Carcinomas: Advanced G3 Non-Small Cell: Randomized Phase II Study of Cisplatin and Etoposide versus Temozolomide and Capecitabine	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EA2185	Comparing the Clinical Impact of Pancreatic Cyst Surveillance Programs	Other	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EA2183	A Phase III Study of Consolidative Radiotherapy in Patients with Oligometastatic Esophageal and Gastric Adenocarcinoma	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EA2187	EA2187 - A Phase 2 Study of Pevonedistat in Combination with Carboplatin and Paclitaxel in Advanced Intrahepatic Cholangiocarcinoma	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EA8143	A Phase 3 RandOmized Study Comparing PERioperative Nivolumab vs. Observation in Patients with Renal Cell Carcinoma Undergoing Nephrectomy (PROSPER RCC)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

EA2176	A Phase III Study of Immune checkpoint inhibition with chemotherapy in treatment-naïve metastatic anal cancer patients	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EA2197	Optimal Perioperative Therapy for Incidental Gallbladder Cancer (OPT-IN)- A Randomized Phase II/III Trial	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
S2001	Randomized Phase II Clinical Trial of Olaparib + Pembrolizumab vs. Olaparib Alone as Maintenance Therapy in Metastatic Pancreatic Cancer Patients with Germline BRCA1 or BRCA2 Mutations.	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
A021806	A Phase III Trial of Perioperative Versus Adjuvant Chemotherapy for Resectable Pancreatic Cancer	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
AHEP1531	Cisplatin and Combination Chemotherapy in Treating Children and Young Adults With Hepatoblastoma or Liver Cancer After Surgery	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EA2186 (GIANT)	A Randomized Phase II Study of Gemcitabine and Nab-Paclitaxel Compared with 5-Fluorouracil, Leucovorin, and Liposomal Irinotecan in Older Patients with Treatment Naïve Metastatic Pancreatic Cancer	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

Novocure EF-27	PANOVA 3: Effect of Tumor Treating Fields (TTFields, 150 kHz) as Front-line Treatment of Locally-advanced Pancreatic Adenocarcinoma Concomitant with Gemcitabine and Nab-paclitaxel	Treatment	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com
Lexicon (Tele-ABC)	A Phase II, Multicenter, Open-label, Safety and Efficacy Study of Xermelo plus First-line Chemotherapy (Cis/Gem) in Patients with Locally Advanced, Unresectable, Recurrent, or Metastatic Biliary Tract Cancer (BTC)	Treatment	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com
QED (PROOF)	A Phase III, Multicenter, Open-Label, Randomized, Controlled Study of Oral Infigratinib versus Cis/Gem (Gemcitabine with Cisplatin) in Subjects with Advanced/Metastatic or Inoperable Cholangiocarcinoma with FGFR2 Gene Fusions/Translocations	Treatment	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com
EMERALD 2	A Phase III, Randomized, Double-blind, Placebo-controlled, Multicenter Study of Durvalumab, Monotherapy or in Combination, with Bevacizumab as Adjuvant Therapy in Patients with Hepatocellular Carcinoma who are at High Risk of Recurrence After Curative Hepatic Resection or	Treatment	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com
CP-MGAH22-06 (MAHOGANNY)	A Phase II-III Trial to Evaluate Margetuximab in Combination with INCMGA00012 and Chemotherapy or MGD013 and Chemotherapy in Patients with Metastatic or Locally Advanced, Treatment-naïve, HER2-Positive Gastric or Gastroesophageal Junction Cancer	Treatment	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com

NAPOLI-3	<p>An Open-label, Randomized, Multi-center, Phase III Study of Irinotecan Liposome Injection, Oxaliplatin, 5-Fluorouracil/Leucovorin vs Nab-paclitaxel + Gemcitabine in Subjects who have not Previously Received Chemotherapy for Metastatic Adenocarcinoma of the Pancreas</p>	Treatment	<p>CHCWM Physician Investigators</p>	<p>CHCWM (616) 954-9800 www.chcwm.com</p>
MK-7902	<p>MK-6482-011 : An Open-label, Randomized, Phase 3 Study of MK-6482 in Combination with Lenvatinib (MK-7902) vs Cabozantinib for Second-line Treatment in Participants with Advanced Renal Cell Carcinoma Who Have Progressed After 1 Prior Anti-PD-1/L1 Combination Regimen. Has experienced disease progression on or after first- or second-line systemic treatment with an anti-PD-1/L1 therapy for locally advanced or metastatic RCC. The anti-PD-1/L1 therapy may have been monotherapy or in combination with other agent(s) such as antiCTLA4 or VEGF-targeted-TKI. The immediately preceding line of treatment has to have been an anti-PD-1/L1 therapy</p>	Treatment	<p>CHCWM Physician Investigators</p>	<p>CHCWM (616) 954-9800 www.chcwm.com</p>
MK-3475	<p>MK-6482-012: An Open-label, Randomized Phase 3 Study to Evaluate Efficacy and Safety of Pembrolizumab (MK-3475) in Combination with MK-6482 and Lenvatinib (MK-7902), or MK-1308A in Combination with Lenvatinib, versus Pembrolizumab and Lenvatinib, as First-line Treatment in Participants with Advanced Clear Cell Renal Cell Carcinoma (ccRCC)</p>	Treatment	<p>CHCWM Physician Investigators</p>	<p>CHCWM (616) 954-9800 www.chcwm.com</p>

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A021703	Randomized Double-blind Phase III Trial of Vitamin D3 Supplementation in Patients with Previously Untreated Metastatic Colorectal Cancer (SOLARIS)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
WF-1806	Myopenia and Mechanisms of Chemotherapy Toxicity in Older Adults with Colorectal Cancer: The M&M Study	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EA2182	A Randomized Phase II Study of De-intensified Chemoradiation for Early Stage Anal Squamous Cell Carcinoma (DECREASE)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
S1613	A Randomized Phase II Study of Trastuzumab and Pertuzumab (TP) Compared to Cetuximab and Irinotecan (CETIRI) in Advanced/Metastatic Colorectal Cancer (mCRC) with HER-2 Amplification	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
URCC 16092	A Phase II Study of Exercise and Low-Dose Ibuprofen for Cognitive Impairment in Colorectal Cancer Patients Receiving Chemotherapy	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
NRG GI004	Colorectal Cancer Metastatic dMMR Immuno-Therapy (COMMIT) Study: A Randomized Phase III Study of mFOLFOX6/Bevacizumab Combination chemotherapy with or without Atezolizumab or Atezolizumab Monotherapy in the First-Line Treatment of Patients with Deficient DNA Mismatch Repair (dMMR) Metastatic Colorectal Cancer	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

S0820	A Double Blind Placebo-Controlled Trial of Eflornithine and Sulindac to Prevent Recurrence of High Risk Adenomas and Second Primary Colorectal Cancers in Patients with Stage 0-III Colon Cancer, Phase III- Preventing Adenomas of the Colon with Eflonithine and Sulindac (PACES)	Prevention	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
A021502	ATOMIC: Adjuvant Trial of Deficient Mismatch Repair in Colon Cancer: A Randomized Trial of Standard Chemotherapy Alone or Combined With Atezolizumab as Adjuvant Therapy for Patients With Stage III Colon Cancer and Deficient DNA Mismatch Repair	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EAQ162CD	Longitudinal Assessment of Financial Burden in Patients with Colon or Rectal Cancer Treated with Curative Intent	Cancer Care Delivery	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EA2165	A Randomized Phase II Study of Nivolumab After Combined Modality Therapy (CMT) in High Risk Anal Cancer	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
FC-11	A Phase II Study Evaluating the Combination of Neratinib Plus Trastuzumab or Neratinib Plus Cetuximab in Patients with "Quadruple Wild-Type" (KRAS/NRAS/BRAF/PIK3CA Wild-Type) Metastatic Colorectal Cancer Based on HER2 Status: Amplified, Non-Amplified (Wild-Type) or Mutated	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
NRG GI005	Phase II/III Study of Circulating tumOr DNA as a Predictive BiomaRker in Adjuvant Chemotherapy in Patients with Stage IIA Colon Cancer (COBRA)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

Merck-4280	MK-4280: A Phase I Trial of MK-4280 (LAG-3 Inhibitor) as Monotherapy and in Combination with Pembrolizumab in Subjects with Advanced Solid Tumors (Now in Expansion Cohort, All Patients get Pembrolizumab + LAG 3 Inhibitor (MK 4280))	Treatment	START Midwest Investigators	START Midwest (616) 954-5554
CRDF-01E	An Expanded Access Program of Onvansertib in Combination with FOLFIRI and BBEVAcizumab for the Second-Line Treatment of Patients with KRAS-Mutated Metastatic Colorectal Cancer (mCRC)	Treatment	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com

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CISTO	Comparison of Intravesical therapy and Surgery as Treatment Options for Bladder Cancer	Observational	SHORE Dr. Brian Lane	Zak Mulder zachary.mulder@spectrumhealth.org 616.391.2724
WF-1802	Influence of Primary Treatment for Prostate Cancer on Work Experience (PCW)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
NRG GU002	NRG GU002: Phase II-III Trial of Adjuvant Radiotherapy and Androgen Deprivation Following Radical Prostatectomy With or Without Adjuvant Docetaxel	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EA8153	Cabaz with Abiraterone versus Abiraterone alone Randomized Trial for Extensive Disease following Docetaxel: the CHARTED 2 Trial	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
A031102	GERM CELL TUMORS/Relapsed or Refractory; Male Only; Comparing Conventional-Dose Chemotherapy Using Paclitaxel, Ifosfamide, and Cisplatin (TIP) with High-Dose Chemotherapy Using Mobilizing Paclitaxel Plus Ifosfamide Followed by High-Dose Carboplatin and Etoposide (TI-CE) as First Salvage Treatment. Phase III Trial	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

A031501	Phase III Randomized Adjuvant Study of MK-3475 (Pembrolizumab) in Muscle Invasive and Locally Advanced Urothelial Carcinoma (Ambassador) Versus Observation	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
AREN03B2	COG-AREN03B2: Renal Tumors Classification, Biology, and Banking Study. Patients must be <30 Years Old at the Time of Diagnosis	Biobanking	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
S1802	Phase III Randomized Trial of Standard Systemic Therapy (SST) Versus Standard Systemic Therapy Plus Definitive Treatment (Surgery Or Radiation) of The Primary Tumor in Metastatic Prostate Cancer	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
A031803	Phase II Trial of Intravesical Gemcitabine and MK-3475 (pembrolizumab) in the Treatment Patients with BCG-Unresponsive Non-Muscle Invasive Bladder Cancer	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
S1806	Phase III Randomized Trial of Concurrent Chemoradiotherapy With or Without Atezolizumab In Localized Muscle Invasive Bladder Cancer (Study SWOG/NRG 1806)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

A031704	PD-Inhibitor (Nivolumab) and Ipilimumab Followed By Nivolumab vs. VEGF TKI Cabozantinib with Nivolumab: A Phase III Trial in Metastatic Untreated Renal Cell Cancer [PDIGREE]	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EA8171	Multiparametric MRI (mpMRI) for Preoperative Staging and Treatment Planning for Newly-Diagnosed Prostate Cancer	Treatment	CRCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com
AREN1921	Treatment of Newly Diagnosed Diffuse Anaplastic Wilms Tumors (DAWT) and Relapsed Favorable Histology Wilms Tumors (FHWT)	Treatment	CRCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com
NRG CC007CD	Increasing the Dose of Survivorship Care Planning In Prostate Cancer Survivors Who Receive Androgen Deprivation Therapy	Cancer Care Delivery	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
NRG GU005	A Phase III IGRT + SBRT vs IGRT + Hypofractionated IMRT for Localized Intermediate Risk Prostate Cancer	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EA8143	A Phase 3 RandOmized Study Comparing PERioperative Nivolumab vs. Observation in Patients with Renal Cell Carcinoma Undergoing Nephrectomy (PROSPER RCC)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

Merck 3475-641	A Phase III, Randomized, Double-blind, Trial of Pembrolizumab (MK-3475) Plus Enzalutamide versus Placebo Plus Enzalutamide in Patients with Metastatic Castration-Resistant Prostate Cancer (mCRPC)	Treatment	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com
Merck MK 3475-921	A Phase III, Randomized, Double-blind, Study of Pembrolizumab (MK-3475) Plus Docetaxel versus Placebo Plus Docetaxel in Patients with Metastatic Castration-Resistant Prostate Cancer (mCRPC)	Treatment	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com
AstraZeneca (PREVAIL)	A Prospective, Non-Interventional Study to Assess the Prevalence of PD-L1 Expression Using the Ventana SP263 Assay in the First-line Setting of Locally Advanced/Unresectable or Metastatic Urothelial Carcinoma	Other	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com
AstraZeneca (NILE)	A Phase III, Randomized, Open-Label, Controlled, Multi-Center, Global Study of First-Line Durvalumab in Combination with Standard of Care Chemotherapy and Durvalumab in Combination with Tremelimumab and Standard of Care Chemotherapy Versus Standard of Care Chemotherapy Alone in Patients with Unresectable Locally Advanced or Metastatic Urothelial Cancer	Biorepository	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com

<u>Short Name</u>	<u>Comments</u>	<u>Type</u>	<u>Investigator</u>	<u>Contact</u>
NRG GY018	A Phase III Randomized, Placebo-controlled Study of Pembrolizumab (MK-3475, NSC #776864) in Addition to Paclitaxel and Carboplatin for Measureable Stage III or IVA, Stage IVB or Recurrent Endometrial Cancer	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
NRG GY006	Newly Diagnosed Bulky Stage IB2, Stage II, IIIB, or IVA Cancer of the Uterine Cervix or Stage II-IVA Vaginal Cancer; Phase II; Radiation Therapy and Cisplatin Alone or in Combination with Intravenous <i>Triapine</i>	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
NRG GY005	PLATINUM-RESISTANT OR -REFRACTORY OVARIAN, FALLOPIAN TUBE, OR PRIMARY/Recurrent, Randomized Phase II/III Study of the Combination of <i>Cediranib</i> and <i>Olaparib</i> Compared to <i>Cediranib</i> or <i>Olaparib</i> alone, or Standard of Care Chemotherapy	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
CC008	A Non-Randomized Prospective Clinical Trial Comparing the Non-Inferiority of Salpingectomy to Salpingo-Oophorectomy to Reduce the Risk of Ovarian Cancer Among BRCA1 Carriers [SOROCK]	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
NRG GY009	A Randomized, Phase II/III Study of Pegylated Liposomal Doxorubicin and CTEP-Supplied Atezolizumab versus Pegylated Liposomal Doxorubicin/Bevacizumab and CTEP-Supplied Atezolizumab versus Pegylated Liposomal Doxorubicin/Bevacizumab in Platinum Resistant Ovarian Cancer	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

STMW 2020.01	Phase 1 Dose Escalation Study of the Agonist Redirected Checkpoint, SL-172154 (SIRP-αFc-CD40L) Administered Intravenously in Subjects with Ovarian Cancer	Treatment	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com
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<u>Short Name</u>	<u>Comments</u>	<u>Type</u>	<u>Investigator</u>	<u>Contact</u>
EA3132	Phase II Randomized Trial of Adjuvant Radiotherapy With or Without Cisplatin for p53 Mutated, Surgically Resected Squamous Cell Carcinoma of the Head and Neck	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
WF-97115	HEAD & NECK: A Phase III Prospective Randomized Trial of Acupuncture for Treatment of Radiation-Induced Xerostomia	Symptom Management	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
NRG HN005	A Randomized Phase II/III Trial of De-intensified Radiation Therapy for Patients with Early-stage, P16-Positive, Non-smoking Associated Oropharyngeal Cancer	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EA3161	A Phase II/III Randomized Study of Maintenance Nivolumab versus Observation in Patients with Locally Advanced, Intermediate Risk HPV Positive OPCA	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
RTOG 1216	Randomized Phase II/III Trial of Adjuvant Radiation Therapy with Cisplatin, Docetaxel-Cetuximab, or Cisplatin-Atezolizumab in Pathologic High-Risk Squamous Cell Cancer of the Head and Neck	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

<u>Short Name</u>	<u>Comments</u>	<u>Type</u>	<u>Investigator</u>	<u>Contact</u>
NHLBI-MDS	National Myelodysplastic Syndromes (MDS) Study	Registry	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
S1925	Randomized, Phase III Study of Early Intervention With Venetoclax And Obinutuzumab Versus Delayed Therapy With Venetoclax And Obinutuzumab In Newly Diagnosed Asymptomatic High-Risk Patients With Chronic Lymphocytic Leukemia/Small Lymphocytic Lymphoma (CLL/SLL): Evolve CLL/SLL Study	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
A041702	A Randomized Phase III Study of Ibrutinib Plus Obinutuzumab Versus Ibrutinib Plus Venetoclax and Obinutuzumab in Untreated Older Patients (> 70 YEARS OF AGE) with Chronic Lymphocytic Leukemia (CLL)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
AALL1521	A Phase 2 Study of the JAK1/JAK2 Inhibitor Ruxolitinib With Chemotherapy in Children With De Novo High-Risk CRLF2-Rearranged and/or JAK Pathway-Mutant Acute Lymphoblastic Leukemia	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
AALL15P1	A Groupwide Pilot Study to Test the Tolerability and Biologic Activity of the Addition of Azacitidine (IND# 133688, NSC# 102816) to Chemotherapy in Infants with Acute Lymphoblastic Leukemia (ALL) and KMT2A (MLL) Gene Rearrangement	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

AALL1631	International Phase 3 Trial in Philadelphia Chromosome-Positive Acute Lymphoblastic Leukemia (Ph+ALL) Testing Imatinib in Combination With Two Different Cytotoxic Chemotherapy Backbones	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
AALL1732	COG-AALL1732: A Phase 3 Randomized Trial of Inotuzumab Ozogamicin (IND#:133494, NSC#: 772518)for Newly Diagnosed High-Risk B-ALL; Risk-Adapted Post-Induction Therapy for High-Risk B-ALL,Mixed Phenotype Acute Leukemia, and Disseminated B-Lly	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
AAML1531	Risk-stratified Therapy for Acute Myeloid Leukemia in Down Syndrome	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
ACCL16N1CD	Documentation and Delivery of Guideline-Consistent Treatment in Adolescent and Young Adult AYA Acute Lymphoblastic Leukemia ALL	Cancer Care Delivery	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
ALLIANCE: A041501	A Phase III Trial to Evaluate the Efficacy of the Addition of Inotuzumab Ozogamicin (a Conjugated Anti-CD22 Monoclonal Antibody) to Frontline Therapy in Young Adults (Ages 18-39 Years) With Newly Diagnosed Precursor B-Cell ALL	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
A041701	A Randomized Phase II/III Study of Conventional Chemotherapy +/- Uproleselan (GMI-1271) in Older Adults With Acute Myeloid Leukemia Receiving Intensive Induction Chemotherapy	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

S1318	<p>A Phase II Study of Blinatumomab (NSC-765986) and POMP (Prednisone, Vincristine, Methotrexate, 6-Mercaptopurine) for Patients = 65 Years of Age with Newly Diagnosed Philadelphia-Chromosome Negative (Ph-) Acute Lymphoblastic Leukemia (ALL) and of Dasatinib (NSC-732517), Prednisone and Blinatumomab for Patients = 65 Years of Age with Newly Diagnosed Philadelphia-Chromosome Positive (Ph+) ALL, and Philadelphia-Chromosome-Like Signature (Ph-Like) ALL (Newly Diagnosed or Relapsed/Refractory) with Known or Presumed Activating Dasatinib-Sensitive Mutations or Kinase Fusions (DSMKF)</p>	Treatment	<p>CRCWM Physician Investigators</p>	<p>CRCWM (616) 391-1230 www.crcwm.org</p>
S1712	<p>A Randomized Phase II Study of Ruxolitinib (NSC-752295) in Combination with BCR-ABL Tyrosine Kinase Inhibitors in Chronic Myeloid Leukemia (CML) Patients with Molecular Evidence of Disease</p>	Treatment	<p>CRCWM Physician Investigators</p>	<p>CRCWM (616) 391-1230 www.crcwm.org</p>
Connect® MDS and AML	<p>The Myelodysplastic Syndromes (MDS) and Acute Myeloid Leukemia (AML) Disease Registry</p>	Registry	<p>CHCWM Physician Investigators</p>	<p>CHCWM (616) 954-9800 www.chcwm.com</p>
USC 9L-16-3	<p>A Phase II Multicenter Study Combining Guadecitabine, a DNA Methyltransferase Inhibitor, with Atezolizumab, An Immune Checkpoint Inhibitor, in Patients with Intermediate or High-risk Myelodysplastic Syndrome or Chronic Myelomonocytic Leukemia</p>	Treatment	<p>CHCWM Physician Investigators</p>	<p>CHCWM (616) 954-9800 www.chcwm.com</p>

<u>Short Name</u>	<u>Comments</u>	<u>Type</u>	<u>Investigator</u>	<u>Contact</u>
Wii Study	Managing Fatigue Using Virtual Reality for Post-Operative Lung Cancer Patients: Understanding the Post-Surgical Non-Small Cell Lung Cancer (NSCLC) Patient's Symptom Experience	Treatment	SHORE: Drs. Geoffrey Lam and Charles Wilekes	Cardiovascular Research Department (616) 391-3886
2019-278 SEER-Lung	A Prospective Blood Sample Collection Study to Develop and Validate a Panel of Protein-based Biomarkers in Patients with Pulmonary Nodules	Other	SHORE: Dr. Glenn VanOtteren	Tara Jager (616) 391-5038 tara.jager@spectrumhealth.org
S1900E	A Phase II Study of AMG 510 in Participants with Previously Treated Stage IV or Recurrent KRAS G12C Mutated Non-Squamous Non-Small Cell Lung Cancer (ECOG-ACRIN Lung-MAP Sub-Study)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
A151216	Non-Squamous, Large IB, II or IIIA, any EGFR/ALK, Resected. Adjuvant Lung Cancer Enrichment Marker Identification and Sequencing Trial (<u>ALCHEMIST</u>)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
E4512	Must be enrolled in A151216. NSCLC; ALK+; Surgically Resected Early Stage. <u>Crizotinib/Placebo</u>	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
A081105	Must be enrolled in A151216. NSCLC; EGFR Mutant, Resected Tumor. Treatment <u>Erlotinib/Placebo</u>	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
A221504	A Randomized, Double-blind, Placebo-controlled Pilot Study of an Oral, Selective Peripheral Opioid Receptor Antagonist in Advanced Non-small Cell Lung Cancer (Adenocarcinoma)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

NRG LU005	Limited Stage Small Cell Lung Cancer (LS-SCLC): A Phase II/III Randomized Study of Chemoradiation vs Chemoradiation + Atezolizumab	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
S1827	S1827 – MRI Brain Surveillance Alone Versus MRI Surveillance and Prophylactic Cranial Irradiation (PCI): A Randomized Phase III Trial in Small-Cell Lung Cancer (MAVERICK)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
S1900B	S1900B - A Phase II Study of LOXO-292 in Patients with RET Fusion-Positive Stage IV or Recurrent Non-Small Cell Lung Cancer (Lung-MAP Sub-Study)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EA5163: INSIGNA	A Randomized, Phase III Study of Firstline Immunotherapy Alone or in Combination with Chemotherapy in Induction/Maintenance or Postprogression in Advanced Nonsquamous Non-Small Cell Lung Cancer (NSCLC) with Immunobiomarker SIGNature-Driven Analysis	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
S1800A	A Phase II RANDOMIZED Study of Ramucirumab Plus MK3475 (Pembrolizumab) versus Standard of Care for Patients Previously Treated with Immunotherapy for Stage IV or Recurrent Non-Small Cell Lung Cancer (Lung-MAP Non-Matched Sub-Study)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
S1900A	A Phase II Study of Rucaparib in Patients with Genomic LOH High And/Or Deleterious Brca1/2 Mutation Stage IV or Recurrent Non-Small Cell Lung Cancer (Lung-Map Sub-Study)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

SWOG LungMAP	A Master Protocol to Evaluate Biomarker-Driven Therapies and Immunotherapies in Previously-Treated Non-Small Cell Lung Cancer (Lung-MAP Screening Study)	Screening	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EA5162	Phase II study of AZD9291 (Osimertinib) advanced NSCLC Patients with Exon 20 Insertion	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
NRG LU002	A Randomized Phase II/III Trial for Maintenance Systemic Therapy Versus Consolidative Stereotactic Body Radiation Therapy (SBRT) Plus Maintenance Systemic therapy for Limited Metastatic Non-Small Cell Lung Cancer (NSCLC)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
NRG CC003	Randomized Phase II/III Trial of Prophylactic Cranial Irradiation With or Without Hippocampal Avoidance for Small Cell Lung Cancer	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
S1900C	A Phase II Study of Talazoparib Plus Avelumab in Patients with Stage IV or Recurrent Non-Squamous Cell Lung Cancer Bearing Pathogenic STK11 Genomic Alterations (Lung-MAP Sub Study)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
S1929	Phase II Randomized Study of Maintenance Atezolizumab Versus Atezolizumab in Combination with Talazoparib in Patients with SLFN11 Positive Extensive Stage Small Cell Lung Cancer (ES-SCLC)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

CARMEN	Randomized, Open Label Phase 3 Study of SAR408701 versus Docetaxel in Previously Treated metastatic non-squamous non-small Cell Lung Cancer patients with CEACAM5 positive tumors	Treatment	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com
MS201923_0050 (DDRiver)	A Phase II, open-label, single-arm study of berzosertib (M6620) in combination with topotecan in participants with relapsed platinum-resistant small-cell lung cancer	Treatment	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com
ADRIATIC	AstraZeneca (ADRIATIC): A Phase III, Randomized, Double-blind, Placebo-controlled, Multi-center, International Study of Durvalumab or Durvalumab and Tremelimumab as Consolidation Treatment for Patients with State I-III Limited Disease Small-cell Lung Cancer Who Have Not Progressed Following Concurrent Chemoradiation Therapy	Treatment	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com
IPSEN MM-398-01-03-04	RESILIENT: A Randomized, Open Label Phase 3 Study of Irinotecan Liposome Injection (ONIVYDE) versus Topotecan in Patients with Small Cell Lung Cancer or after Platinum-based First-line Therapy	Treatment	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com
Helsinn (ANAM)	A Phase III Randomized, Double-Blind, Placebo-Controlled, Multicenter Study to Evaluate the Efficacy and Safety of Anamorelin HCl for the Treatment of Malignancy Associated Weight Loss and Anorexia in Adult Patients with Advanced Non-Small Cell Lung Cancer (NSCLC)	Treatment	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com

CodeBreak200: 20190009	A Phase III Multicenter, Randomized, Open-label, Active-controlled, Study of AMG 510 vs Docetaxel for the Treatment of Previously Treated Locally Advanced and Unresectable or Metastatic NSCLC Subjects with Mutated KRAS p.G12C	Treatment	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com
	An Open Label, Single Arm, Phase IB/II Study to Evaluate the Safety and Efficacy of Grapiprant (ARY-007) in Combination with Pembrolizumab in Patients with Advanced or Metastatic Post-PD-1/L1 Non-small Cell Lung Cancer (NSCLC) Adenocarcinoma	Treatment	START Midwest Investigators	START Midwest (616) 954-5554

<u>Short Name</u>	<u>Comments</u>	<u>Type</u>	<u>Investigator</u>	<u>Contact</u>
A051701	Randomized Phase II/III Study of Venetoclax (ABT 199) Plus Chemoimmunotherapy for MYC/BCL2 Double-hit and Double-expressing Lymphomas	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
S1826	A Phase III, Randomized Study of Nivolumab (Opdivo) Plus AVD or Brentuximab Vedotin (ADCETRIS) plus AVD in Patients (AGE>= 12 years) with Newly Diagnosed Advanced Stage Classical Hodgkin Lymphoma	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
A051301	A Randomized Double-Blind Phase III Study of Ibrutinib During and Following Autologous Stem Cell Transplantation Versus Placebo in Patients with Relapsed Refractory Diffuse Large B Cell Lymphoma of the Activated B-Cell Subtype	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
ANHL12P1	COG-ANHL12P1: A Randomized Phase II study of <u>Brentuximab</u> Vedotin and <u>Crizatinib</u> in Patients with Newly Diagnosed Anaplastic Large Cell Lymphoma (ALCL.) Patients Must be <22 Years of Age	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
S1608	Randomized Phase II Trial in Early Relapsing or Refractory Follicular Lymphoma	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EA4181	A Randomized 3 Arm Phase II Study Comparing 1.) Bendamustine + Rituximab + High Dose Cytarabine (BR/CR), 2.) Bendamustine + Rituximab + High Dose Cytarabine + Acalabrutinib (BR/CR-A) and 3.) Bendamustine + Rituximab + Acalabrutinib (BR-A) in Patients >70 years old with Untreated Mantle Cell Lymphoma	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

AALL1731	A Phase 3 Trial Investigating Blinatumomab (IND# 117467, NSC# 765986) in Combination with Chemotherapy in Patients with Newly Diagnosed Standard Risk or Down syndrome B-Lymphoblastic Leukemia (B-ALL) and the Treatment of Patients with Localized B-Lymphoblastic Lymphoma (B-LLy)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
CytomX	CTMX-M-2029-001: A Phase I/II, First-in-human Study of CX-2029 in Adults with Metastatic or Locally Advanced Unresectable Solid Tumors or Diffuse Large B-Cell Lymphomas	Treatment	START Midwest Investigators	START Midwest (616) 954-5554

<u>Short Name</u>	<u>Comments</u>	<u>Type</u>	<u>Investigator</u>	<u>Contact</u>
2019-362: Melanoma Surveillance	A Prospective Standardized Surveillance of Early-Stage Cutaneous Melanoma Based on Anatomic Stage and Molecular Profiling	Other	SHORE: Dr. G. Paul Wright	Zak Mulder zachary.mulder@spectrumhealth.org (616) 391-2724
EA6194	EA6194: Phase II Randomized Study of Neoadjuvant Pembrolizumab Alone or in Combination with CMP-001 in Patients with Operable Melanoma: Efficacy and Biomarker Study	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EA6141	Phase II/III Trial for STAGE III/IV/Unresectable Melanoma; <u>Nivolumab</u> + <u>Ipilimumab</u> + <u>Sargramostim</u> versus <u>Nivolumab</u> + <u>Ipilimumab</u>	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
A091802	Phase II Randomized Trial of Avelumab Plus Cetuximab Versus Avelumab Alone in Advanced Cutaneous Squamous Cell Carcinoma of the Skin	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EA6134	A Randomized Phase III trial of Dabrafenib + Trametinib followed by Ipilimumab + Nivolumab at Progression vs. Ipilimumab + Nivolumab followed by Dabrafenib + Trametinib at Progression in Patients With Advanced BRAFV600 Mutant Melanoma	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
S1801	A Phase II Randomized Study of Adjuvant Versus NeoAdjuvant MK-3475 (Pembrolizumab) for Clinically Detectable Stage III-IV High-Risk Melanoma	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

ARRAY-818-201	An Open-label, Randomized, Multicenter Trial of Encorafenib + Binimetinib Evaluating a Standard-dose and a High-dose Regimen in Patients with BRAFV600-mutant Melanoma Brain Metastasis	Treatment	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com
Regeneron	REGN3767-ONC-1613: A Phase I, Open-Label, Dose-Escalation and Cohort Expansion First-in-human Study of the Safety, Tolerability, Activity and Pharmacokinetics of REGN3767 (Lag 3 Inhibitor) Administered Alone or in Combination with REGN2810 (PD-1 Inhibitor) in Patients with Advanced Malignancies	Treatment	START Midwest Investigators	START Midwest (616) 954-5554

<u>Short Name</u>	<u>Comments</u>	<u>Type</u>	<u>Investigator</u>	<u>Contact</u>
S1803	Phase III Study of Daratumumab/rHuPH20 (NSC-810307) + Lenalidomide or Lenalidomide as Post-Autologous Stem Cell Transplant Maintenance Therapy in Patients with Multiple Myeloma (MM) Using Minimal Residual Disease to Direct Therapy Duration (DRAMMATIC STUDY)	Therapy	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
Janssen 54767414MM Y3021	AURIGA: A Randomized Study of Daratumumab Plus Lenalidomide Versus Lenalidomide Alone as Maintenance Treatment in Patients with Newly Diagnosed Multiple Myeloma who are Minimal Residual Disease Positive After Frontline Autologous Stem Cell Transplant	Treatment	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com

<u>Short Name</u>	<u>Comments</u>	<u>Type</u>	<u>Investigator</u>	<u>Contact</u>
2011-002: MSTA	Musculoskeletal Tumor Tissue Acquisition Protocol	Molecular Profiling	SHORE: Dr. Matthew Steensma	Zak Mulder zachary.mulder@spectrumhealth.org (616) 391-2724
2012-303: Genetic Outlier	The Identification of Novel Mutations in Outlier Patients	Biobanking	SHORE: Dr. Matthew Steensma	Zak Mulder zachary.mulder@spectrumhealth.org (616) 391-2724
2014-295: NF1 Tissue Harvest	A Clinical Translation Approach to Targeting the Mechanisms Underlying Cutaneous Neurofibroma Formation in NF1	Other	SHORE: Dr. Matthew Steensma	Zak Mulder zachary.mulder@spectrumhealth.org (616) 391-2724
ALTE16C1	Effects of Modern Chemotherapy Regimens on spermatogenesis and Steroidogenesis in Adolescent and Young Adult (AYA) Survivors of Osteosarcoma	Other	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
ARST1431	A Randomized Phase 3 Study of Vincristine, Dactinomycin, Cyclophosphamide (VAC) Alternating With Vincristine and Irinotecan (VI) Versus VAC/VI Plus Temsirolimus (TORI, Torisel, NSC# 683864) in Patients With Intermediate Risk (IR) Rhabdomyosarcoma (RMS)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

<u>Short Name</u>	<u>Comments</u>	<u>Type</u>	<u>Investigator</u>	<u>Contact</u>
SHUB	The Spectrum Health Universal Biorepository Provides a Robust, Diverse Offering of High Quality Research and Potential Diagnostic Samples Associated with Relevant De-identified Clinical Data to Investigators	Biobanking	SHORE: Dr. Sandra Cottingham	Spectrum Health Biorepository david.chesla@spectrumhealth.org (616) 267-2629
AGCT1531	A Phase 3 Study of Active Surveillance for Low Risk and a Randomized Trial of Carboplatin vs. Cisplatin for Standard Risk Pediatric and Adult Patients With Germ Cell Tumors	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
AGCT1532	Phase 3 Accelerated BEP: A Randomised Phase 3 Trial of Accelerated Versus Standard BEP Chemotherapy for Patients With Intermediate and Poor-risk Metastatic Germ Cell Tumours	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
S1609 DART	Dual Anti-CTLA-4 and Anti-PD-1 Blockade in Rare Tumors	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
A221701	Alliance A221701, Phase III placebo-controlled trial to evaluate dexamethasone use for everolimus-induced oral stomatitis: prevention versus early treatment approaches: MIST (My Individualized Stomatitis Treatment)	Symptom Management	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
A231601CD	Improving Surgical Care and Outcomes in Older Cancer Patients through Implementation of An Efficient Pre-Surgical Toolkit (OPTI-surg)	Cancer Care Delivery	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

A221602	Olanzapine with or without Fosaprepitant for the Prevention of Chemotherapy Induced Nausea and Vomiting (CINV) in Patients Receiving Highly Emetogenic Chemotherapy (HEC): A Phase III Randomized, Double-Blind, Placebo-Controlled Trial	Symptom Management	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
DCP-001	Use of a Clinical Trial Screening Tool to Address Cancer Health Disparities in the NCI Community Oncology Research Program (NCORP)	Screening	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
S1714	A Prospective Observational Cohort Study to Develop A Predictive Model of Taxane-Induced Peripheral Neuropathy in Cancer Patients	Quality of Life	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
URCC 18007	Randomized Placebo-Controlled Trial of Bupropion for Cancer-Related Fatigue	Symptom Management	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
WF-10217	Work Ability in Young Adult Survivors (WAYS)	Behavioral	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EAZ171	Prospective validation trial of taxane therapy (docetaxel or weekly paclitaxel) and risk of chemotherapy-induced peripheral neuropathy in African American Women	Supportive Care	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

A231602CD	Assessing Financial Difficulty in Patients with Blood Cancers	Economics of Care	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
EAY131-MATCH Screening Trial	Targeted Therapy Directed by Genetic Testing in Treating Patients with Advanced Refractory Solid Tumors, Lymphomas, or Multiple Myeloma	Molecular Profiling	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
ASCO TAPUR	Targeted Agent and Profiling Utilization Registry Study: All Drugs Provided	Molecular Profiling	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
NCI PDM 9846	Patient-derived Models Tissue Procurement Protocol for the National Cancer Institute	Biobanking	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
ACCRU 2018-01	Blood Sample Collection to Evaluate Biomarkers in Subjects with Untreated Solid Tumors	Other	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
APEC1621	Targeted Therapy Directed by Genetic Testing in Treating Pediatric Patients with Relapsed or Refractory Advanced Solid Tumors, Non-Hodgkin Lymphomas, or Histiocytic Disorders (The Pediatric MATCH Trial)	Treatment	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org
2020-285	NCI COVID-19 In Cancer Patients Study (N-CCaPS): A Longitudinal Natural History Study	Molecular Profiling	CRCWM Physician Investigators	CRCWM (616) 391-1230 www.crcwm.org

Exact Sciences	Blood Sample Collection to Evaluate Biomarkers in Subjects with Untreated Solid Tumors from Twelve Primary Cancer Diagnoses: Breast, Lung, Colorectal, Prostate, Bladder, Uterine, Kidney, Renal, Pelvis, Pancreatic, Liver, Stomach, Ovarian, and Esophageal Cancer	Biobanking	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com
STMW 2020.03	A Phase 1b, First-in-Human, Dose Escalation and Expansion Study of XMT-1536 In Patients with Solid Tumors Likely to Express NaPi2b	Treatment	CHCWM Physician Investigators	CHCWM (616) 954-9800 www.chcwm.com
ALX148 (Alexo)	<i>AT1488001</i>: Phase I, Dose Escalation Study of ALX148 in Patients with Advanced Solid Tumors or Lymphoma Examining Safety, Pharmacokinetics and Tolerability. Now in Combination Phase II with Pembrolizumab, Rituximab, and Trastuzumab	Treatment	START Midwest Investigators	START Midwest (616) 954-5554
AVID-100	<i>AVID100-01</i>: A Phase IA/IIA Cohort Dose Escalation Trial to Determine the Safety, Tolerance, Maximum Tolerated Dose, and Preliminary Antineoplastic Activity of AVID100, an Anti-Human Epidermal Growth Factor Receptor (EGFR) Monoclonal Antibody Linked to the Maytansinoid DM1 in Patients with Advanced or Metastatic Solid Tumors of Epithelial Origin	Treatment	START Midwest Investigators	START Midwest (616) 954-5554
APG-115	<i>APG-115</i>: A Phase I Study of the Safety, Pharmacokinetic and Pharmacodynamic Properties of Orally Administered APG-115 in Patients with Advanced Solid Tumors or Lymphomas	Treatment	START Midwest Investigators	START Midwest (616) 954-5554

Ascentage Pharma: 1252	<i>APG-1252-US-001: Phase I, Study of the Safety, Pharmacokinetic and Pharmacodynamic Properties of Intravenously Administered APG-1252 in Patients with Small Cell Lung Cancer (SCLC) or Other Solid Tumors</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
Ascentage Pharma: 1387	<i>APG-1387-US-001 : A Phase I Study of the Safety, Pharmacokinetic and Pharmacodynamic Properties of Intravenously Administered APG-1387 as a Single Agent or in Combination with Systemic Anti-cancer Agents in Patients with Advanced Solid Tumors or Hematologic Malignancies</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
BeiGene USA	<i>BGB-290-103: A Phase IB Study to Assess the Safety, Tolerability, and Clinical Activity of BGB-290 in Combination with Temozolomide (TMZ) in Subjects with Locally Advanced or Metastatic Solid Tumors</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
Bristol-Myers Squibb	<i>CA031002: A Phase I/II First-in-human Study of BMS-986258 (anti TIM3) Alone and in Combination with Nivolumab in Advanced Malignant Tumors</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
COM701- Compugen	<i>CPG-01-001: A Phase IA/IB Study of COM701 as Monotherapy and in Combination with an Anti-PD-1 Antibody in Subjects with Advanced Solid Tumors</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
COM902- Compugen	<i>CPG-02-101: A Phase I Study of the Safety and Tolerability of COM902 in Subjects with Advanced Malignancies</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556

Coordination Pharmaceutica ls: 100	<i>CPI-CL18-001: A Phase I, First in Human Study Evaluation the Safety, Tolerability, and Pharmacokinetics of CPI-100 via Intravenous Infusion in Patients with Advanced Solid Tumors</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
Coordination Pharmaceutica ls: 200	<i>CPI-200CL01: A Phase I, First in Human Study Evaluation the Safety, Tolerability, and Pharmacokinetics of CPI-200 via Intravenous Infusion in Patients with Advanced Solid Tumors</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
Exelixis	<i>XL092: A Dose-Escalation and Expansion Study of the Safety and Pharmacokinetics of XL092 in Subjects with Inoperable Locally Advanced or Metastatic Solid Tumors</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
Ikena	<i>IK-175-001: A Phase I, Open-Label, Dose-Escalation, and Expansion Study of IK-175, an Oral Aryl Hydrocarbon Receptor (AHR) Inhibitor in Patients with Locally Advanced or Metastatic Solid Tumors and Urothelial Carcinoma</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
InhibRx-105	<i>INBRX-105: An Open-Label, Multicenter, First-in-human, Dose-Escalation Phase I Study of INBRX-105 in Subjects with Locally Advanced or Metastatic Solid Tumors, Hodgkin or Non-Hodgkin Lymphoma</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
InhibRx-106	<i>INBRX-106: An Open-Label, Multicenter, First-in-human, Dose-Escalation, Phase I Study of INBRX-106 in Subjects with Locally Advanced or Metastatic Solid Tumors</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556

InhibRx-109	<i>INBRX-109: An Open-Label, Multicenter, First-in-human, Dose-Escalation, Phase I Study of INBRX-109 in Subjects with Locally Advanced or Metastatic Solid Tumors Including Sarcomas</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
Innovent	<i>IBI188: A Phase IA Study Evaluating the Safety, Tolerability, and Initial Efficacy of Recombinant Human Anti-Cluster Differentiation Antigen 47 (CD47) Monoclonal Antibody Injection (IBI188) in Patients with Advanced Malignant Tumors and Lymphomas</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
Klus	<i>KL264-01: A Phase I/II, First-in-human, Study of SKB264 in Patients with Locally Advanced Unresectable/Metastatic Solid Tumors who are Refractory to Available Standard Therapies</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
Expanded Access Loxo	<i>LOXO-RET-18037: A Multicenter Expanded Access Program (EAP) for the Treatment of Patients with Locally Advanced or Metastatic Solid Tumors with Rearranged During Transfection (RET) Activation (LIBRETTO-201)</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
MacroGenics-018	<i>MGC018-01: A Phase I/II, First-in-human, Open-Label, Dose-Escalation Study of MGC018 (Anti-B7-H3 Antibody Drug Conjugate) Alone and in Combination with MGA012 (Anti-PD-1 Antibody) in Patients with Advanced Solid Tumors</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
MacroGenics-019	<i>MGD019: A Phase I, First-in-human, Open-Label, Dose-Escalation and Cohort Expansion Study of MGA019, a Bispecific DART Protein Binding PD-1 and CTLA-4 in Patients with Unresectable or Metastatic Neoplasms</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556

Merck-5618	<i>MK-5618: A Phase IB Multicenter Clinical Study of Selumetinib (MK-5618) in Combination with Pembrolizumab (MK-3475) in Participants with Advanced/Metastatic Solid Tumors</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
Merck-8353	<i>MK-8353-013: A Phase IB Study to Evaluate the Safety and Tolerability of MK-8353 in Combination with Pembrolizumab in Patients with Advanced Malignancies</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
Odonate	<i>ODO-TE-S101: An Open-label, Study of the Effect of Tesetaxel on the QTc Interval and the Effect of Food, Itraconazole, and Rifampin on Tesetaxel Pharmacokinetics in Patients with Advanced Solid Tumors</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
Symphogen	<i>Sym021-01: A Phase I Open-Label, Multicenter Trial Investigating the Safety, Tolerability and Preliminary Antineoplastic Activity of Sym021 (Anti-PD-1) in Patients with Advanced Solid Tumor Malignancies or Lymphomas</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
Syros	<i>SY-5609-101: A Phase I Study of SY-5609, an Oral, Selective CDK7 Inhibitor, in Adult Patients with Selected Advanced Solid Tumors</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
SCI-CD47-001	<i>HU5F9-G4 : First-in-human Phase I Dose-escalation Trial of HU5F9-G4 in Patients with Advanced Solid Malignancies</i>	Treatment	START Midwest Investigators	START Midwest (616) 954-5556

R3767-ONC-1613	REGN3767 : A Phase I, Open-label, Dose-escalation and Cohort Expansion First-in-human, Study of the Safety, Tolerability, Activity, and Pharmacokinetics of REGN3767 (Anti-LAG-3 mAb) Administered Alone or in Combination with REGN2810 (Anti-PD-1-mAb) in Patients with Advanced Malignancies	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
CP-MGA012-01	MGA012 : A Phase I Study of the Safety, Tolerability, and Pharmacokinetics of MGA012 in Patients with Advanced Solid Tumors	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
ASN003-101	ASN003 : A Phase I, Open-label, Dose-finding, and Cohort Expansion Study of ASN003 in Subjects with Advanced Solid Tumors	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
5F9004	Hu5F9-G4-Cetuximab : A Phase Ib/II Trial of Hu5F9-G4 in Combination with Cetuximab in Patients with Solid Tumors and Advanced Colorectal Cancer	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
APX-CLN-0011	A Phase I Study of APX3330 in Patients with Advanced Solid Tumors	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
CP-MGA271-03	Macrogenics Protocol CP-MGA271-03: A Phase 1, Open-Label, Dose Escalation Study of MGA271 in Combination with Pembrolizumab in Patients with B7-H3-Expressing Melanoma, Squamous Cell Cancer of the Head and Neck, Non-Small Cell Lung Cancer, Urothelial Cancer and Other B7-H3-Expressing Cancers	Treatment	START Midwest Investigators	START Midwest (616) 954-5556

CPI-0209	A Phase I/II Study of CPI-0209 Monotherapy in Combination with Other Therapy in Patients with Advanced Solid Tumors	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
Sym 021	Sym 021: A Phase 1, Open-Label, Multicenter Trial investigating the Safety, Tolerability, and Preliminary Antineoplastic Activity of Sym021 (Anti-PD-1) as Monotherapy and in Combination with either Sym022 (Anti-LAG-3) or SYM023 (Anti-TIM-3) in Patients with Advanced Solid Tumor Malignancies or Lymphomas	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
eFT226	A Phase I-II Dose-escalation and Cohort-expansion of Study of Intravenous Zotatfin (eFT226) in Subjects with Selected Advanced Solid Tumor Malignancies	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
	Phase 1/2A Dose Escalation, Finding and Expansion Study Evaluating Safety, Tolerability, Pharmacokinetics, Pharmacodynamics and Anti Tumor Activity of PF-07104091 as a Single Agent and In Combination Therapy	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
EPZ-6438	A Phase I, Open-label Multi-dose Two-Part Study to Characterize the Effects of a Strong CYP3A4 Inhibitor on the Steady-State Pharmacokinetics of Tazemetostat (EPZ-6438), and the Effects of a Strong CYP3A4 Inducer on the Steady-State Pharmacokinetics of Tazemetostat in Subjects with Advanced Malignancies	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
NEON-1	An Open-Label Study of ALPN-202 in Subjects with Advanced Malignancies (NEON-1)	Treatment	START Midwest Investigators	START Midwest (616) 954-5556

	A Phase 1 Multicenter Global First in Human Study of the CD73 Inhibitor LY3475070 as Monotherapy or in Combination with Pembrolizumab in Patients with Advanced Solid Malignancies	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
COM902	A Phase 1 Study of the Safety and Tolerability of COM902 in Subjects with Advanced Malignancies	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
eFT226	A Phase 1-2 Dose-Escalation and Cohort-Expansion Study of Intravenous eFT226 in Subjects with Selected Advanced Solid Tumor Malignancies	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
JTX-8064	Phase 1 First-in-Human (FIH) Study of Leukocyte Immunoglobulin- Like Receptor B2 (LILRB2) Inhibitor Monoclonal Antibody (mAb) JTX-8064, as Monotherapy and in Combination with a Programmed Cell Death Receptor-1 (PD-1) Inhibitor, in Adult Subjects with Advanced Refractory Solid Tumor Malignancies	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
LYT-200	A Phase 1/2 Open-label, Multi-center Study of the Safety, Pharmacokinetics, and Anti-tumor Activity of LYT-200 Alone and in Combination with Chemotherapy or Anti-PD-1 in Patients with Metastatic Solid Tumors	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
ARRY-558	A Phase 1, Open-Label, Multi-Center, Dose Escalation and Dose Expansion Study to Evaluate the Safety, Tolerability, Pharmacokinetics, and Preliminary Evidence of Anti-Tumor Activity of PF-07284892 (ARRY-558) as a Single Agent and in Combination Therapy in Participants with Advanced Solid Tumors	Treatment	START Midwest Investigators	START Midwest (616) 954-5556

GEN 1046	First-in-human, open-label, dose-escalation trial with expansion cohorts to evaluate safety of GEN1046 in subjects with malignant solid tumors	Treatment	START Midwest Investigators	START Midwest (616) 954-5556
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