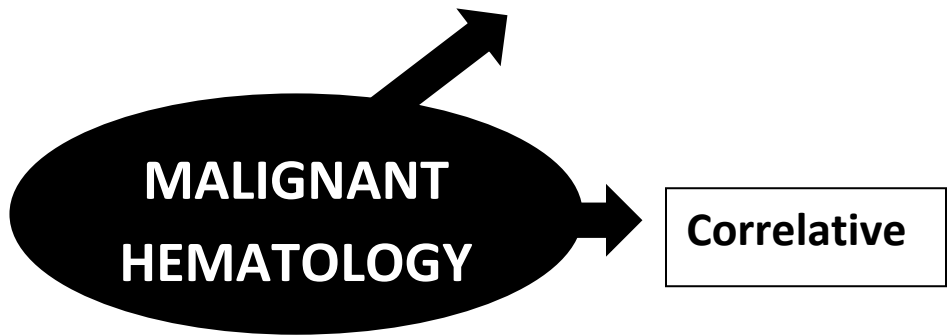


Epidemiologic

UCI-13-02 Assessing Fitness and Physical Activity in Acute Lymphoblastic Leukemia (ALL) Patients in Remission (Aizik)



UCI-14-03 Role of inflammation in the pathogenesis of myeloproliferative neoplasm (Fleischman)

UCI-16-01 Pathway Analyses for Individualized Network Therapeutics for Cancer (PAINT Cancer) (Nelson)

UCI 15-65**
Effect of candidate blood cancer therapies on normal human lymphocytes (Fruman)

CLL

UCI 15-88

A Randomized, Multicenter, Open-Label, Non-Inferiority, Phase 3 Study of ACP-196 Versus Ibrutinib in Previously Treated Subjects with High Risk Chronic Lymphocytic Leukemia (O'Brien)

UCI 17-03**

Phase 2 Study of the Combination of Ibrutinib Plus Venetoclax in Subjects with Treatment Naïve Chronic Lymphocytic Leukemia (CLL)/Small Lymphocytic Lymphoma (SLL) (O'Brien)

UCI 16-95

A Phase 1b/2 Dose-Escalation and Cohort-Expansion Study of a Non-Covalent, Reversible Bruton's Tyrosine Kinase Inhibitor, SNS 062, in Patients With B-Lymphoid Malignancies (O'Brien)

UCI 15-18

An Open-Label, Multicenter Phase 1 Study to Investigate the Safety and Tolerability of REGN1979, an Anti-CD20 x Anti-CD3 Bispecific Monoclonal Antibody, in Patients with CD20+ B Cell Malignancies Previously Treated with CD20-Directed Antibody Therapy (O'Brien)

UCI 16-65

A Phase 1 clinical trial to evaluate obinutuzumab with high-dose Ibrutinib for the treatment of patients with chronic lymphocytic leukemia with progressive disease on single agent Ibrutinib (Jeyakumar)

UCI 17-19 (CLL/SLL/Richters)**

A Phase 1, Open-Label, Multicentre Study to Assess the Safety, Tolerability, Pharmacokinetics and Preliminary Antitumor Activity of Ascending Doses of AZD5991 in Subjects with Relapsed or Refractory Haematologic Malignancies (O'Brien)

ALL

ECOG E1910

A phase II Randomized Trial of Blinatumomab for Newly Diagnosed BCR-ABL-negative B lineage Acute Lymphoblastic Leukemia (ALL) (Jeyakumar)

UCI 15-58*On Hold*

A Phase I/II Study Evaluating KTE-C19 in Subjects with Relapsed/Refractory ALL (O'Brien)

UCI 16-13**

A Phase I Dose Escalation with Two Disease Specific Expansions, Multicenter, Open-label, Safety, Pharmacokinetic and Pharmacodynamic Study of APTO-253 in Patients with Relapsed or Refractory Hematologic Malignancies (Jeyakumar)

SWOG S1318 (Ph- Only)

A Phase II Study of Blinatumomab (NSC-765986) and POMP (Prednisone, Vincristine, Methotrexate, 6-Mercaptopurine) for Patients \geq 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 \geq 18 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 (Jeyakumar)

UCI 16-10

A Phase 1b, Open-Label, Dose Escalation and Expansion Study Evaluating the Safety and Efficacy of Entospletinib (GS-9973) with Vincristine and Dexamethasone in Adult Subjects with Relapsed or Refractory Acute Lymphoid Leukemia (ALL) (O'Brien)

UCI 16-95

A Phase 1b/2 Dose-Escalation and Cohort-Expansion Study of a Non-Covalent, Reversible Bruton's Tyrosine Kinase Inhibitor, SNS 062, in Patients With B-Lymphoid Malignancies (O'Brien)

T-Cell

UCI 16-45

A Phase I/Ib, Dose Escalation Study to Evaluate Safety and Efficacy of RP6530, a dual PI3K δ/γ inhibitor, in Patients with Relapsed or Refractory T-cell Lymphoma. (Pinter-Brown)

UCI 16-46

A Phase 1 dose-ranging study to investigate the safety, tolerability, and pharmacokinetics of MRG-106 following local intratumoral and subcutaneous injection in patients with cutaneous T-cell lymphoma (CTCL), mycosis fungoides (MF) sub-type. (Pinter-Brown)

UCI 17-19 (T-Cell/NHL)**

A Phase 1, Open-Label, Multicentre Study to Assess the Safety, Tolerability, Pharmacokinetics and Preliminary Antitumor Activity of Ascending Doses of AZD5991 in Subjects with Relapsed or Refractory Haematologic Malignancies (O'Brien)

Lymphoma

ECOG-EAY131

Molecular Analysis for Therapy Choice (MATCH). (Seery)

UCI 16-13**

A Phase I Dose Escalation with Two Disease Specific Expansions, Multicenter, Open-label, Safety, Pharmacokinetic and Pharmacodynamic Study of APTO-253 in Patients with Relapsed or Refractory Hematologic Malignancies (Jeyakumar)

B-Cell

UCI 14-84

Phase 3 Randomized, Double-Blind, Placebo Controlled, Multicenter Study to Compare the Efficacy and Safety of Lenalidomide Plus R-CHOP versus Placebo Plus R-CHOP Chemotherapy in Subjects with previously Untreated Activated B-Cell Type Diffuse Large B-Cell Lymphoma (Jeyakumar)

UCI 16-95

A Phase 1b/2 Dose-Escalation and Cohort-Expansion Study of a Non-Covalent, Reversible Bruton's Tyrosine Kinase Inhibitor, SNS 062, in Patients With B-Lymphoid Malignancies (O'Brien)

UCI 15-18

An Open-Label, Multicenter Phase 1 Study to Investigate the Safety and Tolerability of REGN1979, an Anti-CD20 x Anti-CD3 Bispecific Monoclonal Antibody, in Patients with CD20+ B Cell Malignancies Previously Treated with CD20-Directed Antibody Therapy (O'Brien)

AML

UCI 15-29

A Phase III, randomized, double-blind, placebo-controlled study to compare efficacy and safety of oral Azacitidine plus best supportive care versus best supportive care as maintenance therapy in subjects with AML in complete remission.
(Jeyakumar)

UCI 16-100**

A Phase 2/3 Multicenter, Open-label, 3-arm, 2-Stage Randomized Study of ASP2215 (Gilteritinib), Combination of ASP2215 Plus Azacitidine and Azacitidine Alone in the Treatment of Newly Diagnosed Acute Myeloid Leukemia With FLT3 Mutation in Patients Not Eligible for Intensive Induction Chemotherapy (Jeyakumar)

UCI 16-11**

A Randomized Phase 2/3 Study of DACOGEN® (Decitabine) Plus JNJ-56022473 (Anti-CD123) Versus DACOGEN (Decitabine) Alone in Patients with AML who are not Candidates for Intensive Chemotherapy (Jeyakumar)

UCI 15-69

Phase 3 study of ASP2215 vs. Salvage Chemo in Pts w/ Relapsed or Refractory AML w/ FLT3 Mutation (Jeyakumar)

UCI 16-13**

A Phase I Dose Escalation with Two Disease Specific Expansions, Multicenter, Open-label, Safety, Pharmacokinetic and Pharmacodynamic Study of APTO-253 in Patients with Relapsed or Refractory Hematologic Malignancies (Jeyakumar)

MPN

UCI 17-31**

An Open-Label Phase 2 Study of Itacitinib (INCB039110) in Combination With Low Dose Ruxolitinib or Itacitinib Alone Following Ruxolitinib in Subjects With Myelofibrosis (Fleischman)

MDS

UCI 14-93

Phase 1b Study of Ibrutinib and Azacitidine for the treatment of patients with higher risk MDS (Jeyakumar)

UCI 16-13**

A Phase I Dose Escalation with Two Disease Specific Expansions, Multicenter, Open-label, Safety, Pharmacokinetic and Pharmacodynamic Study of APTO-253 in Patients with Relapsed or Refractory Hematologic Malignancies (Jeyakumar)

MALIGNANT HEMATOLOGY



ECOG-EAY131

Molecular Analysis for Therapy Choice (MATCH) (Seery)

Multiple Myeloma

ECOG E1A11

Randomized Phase III Trial of Bortezomib, LENalidomide and Dexamethasone (VRd) Versus Carfilzomib, Lenalidomide, Dexamethasone (CRd) Followed by Limited or Indefinite DURATION Lenalidomide Maintenance in Patients with Newly Diagnosed Symptomatic Multiple Myeloma (ENDURANCE) (Brem)

UCI 17-19 (Myeloma)**

A Phase 1, Open-Label, Multicentre Study to Assess the Safety, Tolerability, Pharmacokinetics and Preliminary Antitumor Activity of Ascending Doses of AZD5991 in Subjects with Relapsed or Refractory Haematologic Malignancies (O'Brien)

UCI 14-96

A Phase I/II Study of PiC-D (Ixazomib in Combination with Pomalidomide, Clarithromycin and Dexamethasone) in Patients with Double Refractory Multiple Myeloma (Brem)